Metagenomic analysis of microbial communities in rapid sand filter treating groundwater. Community diversity and metabolic potential

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Metal specific modulation of community permissiveness towards broad host range plasmids through stress

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Giant viruses

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More than a century ago, the notion of "virus" was introduced to designate infectious agents invisible to the light microscope and capable of passing through "sterilizing" filters. In addition to their extremely small size, most viruses studied over the years also exhibited minimal genomes and gene contents, almost entirely relying on cell-encoded functions to multiply, as expected from absolute intracellular parasites. Unexpectedly, the last ten years have seen the discovery of 4 different families of eukaryotic "giant viruses" exhibiting particles of cellular dimensions as well as gene contents overlapping in size with that of bacteria and some parasitic eukaryotes. Although all presently known giant viruses have been isolated using Acanthamoeba as laboratory host, related members are now starting to be found in other protozoa, such as marine picoplankton species the population of which they regulate. Representatives of two families of giant viruses have been revived from a layer of Siberian permafrost radiocarbon dated from 30,000 years ago, raising concern that pathogenic viruses from long past epidemics might also remain infectious and resurface in circumpolar regions as a consequence of global warming and industrial exploitations.

The unexpected abundance, ubiquity and diversity of giant viruses, as well as the alien nature of their gene contents deeply challenge conventional conceptions about the origin and evolution of all DNA viruses and raise the question of their evolutionary relationship with the 3 cellular domains forming today's Tree of Life, and possibly other long extinct cellular lineages.

I will conclude by discussing the nature of the epistemological barrier that led to the delayed discovery of these highly visible viruses, most of which were spotted 10 years before their viral nature was eventually recognized.
(human) microbiome

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The performance of the human microbiome is perturbed in unknown ways by a variety of factors including disease and diet. Here we aimed to determine the impact of inflammatory bowel disease (IBD) and a resistant starch diet on the gut microbiome community composition and functional potential by application of an array of omics technologies: 16S rRNA sequencing, metagenomics, metaproteomics and metabolomics. The first studies focused on a Swedish twin cohort and a longitudinal cohort for IBD. Specific bacterial species, proteins and metabolites correlated with different sub-phenotypes of the IBD; including a reduced amount of normal beneficial microbes as well as proteins involved in butyrate metabolism, suggesting potential biomarkers of IBD. For the dietary study, subjects transitioned from a baseline diet to one with low or high amounts of resistant starch. Specific bacteria, proteins and metabolites were shown to significantly vary in abundance when comparing baseline to the high resistant starch diet. Together these studies exemplify the use of omics to provide potential bioindicators of specific physiological states of importance to human health.
Fungi are notorious plant pathogens that cause serious diseases on natural vegetations and crop plants. The majority of plants, however, is not infected by fungal pathogens as they recognize pathogen-associated molecular patterns (PAMPs) like chitin or glucans by pattern recognition receptors that mediate PAMP-triggered immunity (PTI), a basal defense response effective against potential fungal pathogens. Successful fungal plant pathogens secrete effectors to suppress PTI and alter host plant physiology enabling them to infect plants. In turn, plants have evolved immune receptors that recognize effectors resulting in effector-triggered immunity (ETI) including the hypersensitive response, effective against biotrophic fungal plant pathogens that require living cells to feed on. Other fungal pathogens are hemi-biotrophic, which start infection as a biotroph, but after having colonized the host tissue can also feed on death tissue. Necrotrophic fungal pathogens kill host tissue before they start to feed on it. Co-evolution between fungal pathogens and their hosts had led to the development of numerous effectors in fungal plant pathogens and corresponding resistance proteins in host plants, which has generated an arms race genetically described by the gene-for-gene concept. Resistance genes encoding resistance proteins have now been cloned and are successfully transferred to crop plants by classical breeding or as transgenes stapled into one plant cultivar.
Ranking risks in resistomes. The intrinsic resistome

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Before the antibiotic era, bacterial pathogens were by definition susceptible to the antibiotics that pharmaceutical companies have developed to fight infections. Nevertheless, soon after the introduction of antibiotics for therapy, resistant isolates emerge and spread among the population. Resistance can be acquired either because of mutations or by acquiring resistance genes from a donor strain. Since pathogens were susceptible, resistance genes should come from non pathogens and, most likely from environmental microorganisms. Indeed, the analysis of metagenomes from very different ecosystems has shown that antibiotic resistance genes are ubiquitously find in any place in Earth. These genes (the resistome) can be transferred to human pathogens, then constituting a risk for human health. However, if all ecosystems contain resistance genes, all of them will be risky, which is the same as saying that no one has a relevant differential risk. In the presentation the possibility of ranking risk of resistomes will be discussed. The ranking is based in the likehood that the finding of a specific gene in a given ecosystem constitutes a real risk for human health. One of the relevant aspects to take into consideration is whether those genes belong to the transferrable resistome (the mobilome) or are just a part to the set of elements that contribute to the characteristic phenotype of susceptibility to antimicrobials of a given microorganisms (its intrinsic resistome). Current information on the elements forming part of the intrinsic resistome will be discussed in the presentation.

Further reading:
http://www.nature.com/nrmicro/journal/vaop/ncurrent/full/nrmicro3399.html
Antibiotic resistance is a growing public health concern that negatively impacts treatment outcomes of most infections. The issue is projected to grow substantially in the coming decades and new approaches are needed to curb the spread of resistance. Notably, resistance evolves through a complex interplay between residents of most microbial communities and organisms causing disease. In spite of significant efforts using a variety of experimental and computational approaches our understanding of the dissemination of resistance genes is still limited. To institute preventative regulations on key environments harboring microbial communities that influence resistance gene dissemination general and unbiased approaches are necessary. Yet, to date our ability to assess the impact of specific microbial communities on the dissemination of antibiotic resistance is inadequate. Recent improvements in experimental and computational approaches aimed at addressing this overall goal are discussed with a focus on key environments believed to influence resistance gene dissemination.
The introduction of antibiotics revolutionized human medicine and agricultural industry, but their success has created a public health threat caused by antibiotic resistance (ABR). Indisputably, human use of antibiotic has contributed to the expansions of ABR. At the same time it is becoming increasingly clear that antibiotic use in animal husbandry has also created a large reservoir of ABR with increasing risk of spill over of ABR from animals to humans. Whole genome sequencing (WGS) has revolutionizing research into transmission of ABR by providing the optimal resolution to infer epidemiological linkage of multidrug resistant isolates and their resistance genes and plasmids. However, frequently used WGS technologies rely on the generation of sequences with short-read length resulting in many contigs of assembled reads without easy assorting contigs to plasmids and the main chromosome. We used a novel algorithm called PLACNET that identifies plasmid contigs in WGS projects to study genetic relatedness of ESBL-genes containing plasmids from animal and human origin. Plasmid reconstructions, validated by long-read DNA sequencing, indicated that genetically unrelated human isolates shared virtually identical plasmids with animal isolates.

Another potential drawback of WGS for studying transmission of ABR is the lack of a standardized and expandable classification scheme since most WGS-based analyses rely on SNP detection from reads mapped to reference genomes that are often distinct for different epidemiological studies. This limitation may be overcome by using a gene-by-gene comparison approach just like classical MLST but with an important extension of the numbers of analysed genes from seven to several hundreds/thousands. We developed an extended, core genome MLST scheme and showed its value in determining the clonal relatedness of multi-resistant Enterococcus faecium from hospital outbreaks in three different countries.
Resistome

Networks of Exchanging Antibiotic Resistance in Human and Environmental Microbiota

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While the most acute effects of increasing antibiotic resistance in pathogens are observed in clinical settings, it is becoming increasingly clear that the evolution and transmission dynamics of resistance gene dissemination is an ecological problem. Indeed, steady use and abuse of antibiotics over the past century in food animals, humans, and the environment has provided substantial selective pressure for enrichment of resistance genotypes in each of their associated microbiomes. An over-reliance on culture-based methods, the standard in the study of clinical resistance, has vastly underestimated these reservoirs of resistance genes (or ‘resistomes’). To address this issue, we have recently developed high-throughput metagenomic functional selections, aided by next-generation sequencing, to characterize resistomes encoded by the microbiota of healthy human adults and children as well as diverse soils. Hundreds of resistance genes we identify from specific taxa in these different microbial communities are identical to resistance genes found in major human pathogens, indicating recent genetic exchange between these microbes. We also find thousands of functionally validated resistance genes which are genetically novel, but flanked by genes involved in horizontal gene transfer, including transposases and integrases. Together, these findings highlight the substantial antibiotic resistome encoded by microbes from diverse environments, which is available for exchange with pathogens, with the potential to severely exacerbate the problems with clinical resistance.
Neisseria meningitidis is a strictly human pathogen that colonizes the nasopharynx as a commensal but occasionally crosses the epithelial barriers to cause sepsis and/or meningitis. Proteins secreted by this Gram-negative diplococcus are considered to be important for colonization and/or infection. Of six protein secretion systems widely disseminated among Gram-negative bacteria, only two are present in N. meningitidis, i.e. type I and type V, the latter being subdivided in autotransporters and two-partner secretion (TPS) systems.

We are investigating the functions of type V secreted proteins and found that many of them are involved in interbacterial interactions. Neisserial IgA protease was the first autotransporter ever described. Besides its protease domain, also a ~40-kDa polypeptide of unknown function, the α-peptide, is transported to the bacterial cell surface. Our results demonstrate that this positively charged α-peptide and a cell-surface-exposed lipoprotein, the neisserial heparin-binding antigen (NHBA), are involved in biofilm formation by binding extracellular DNA, an important matrix component of meningococcal biofilms. Another autotransporter, the protease NalP, inhibits biofilm formation by cleaving the α-peptide of IgA protease and NHBA from the cell surface. Yet another autotransporter, designated AutA, is only expressed in a limited number of meningococcal isolates. Its expression causes bacterial autoaggregation and influences the architecture of the biofilms formed.

The substrates of the meningococcal TPS systems are large proteins. These proteins were found to contain a small toxic domain at their C-terminus that inhibits the growth of related bacteria. Similar toxic domains were found at the C-terminus of another family of proteins, MafB. MafB proteins are transported to the cell surface via a protein, MafA, representing a novel protein secretion system. Like the TPS system, the MAF system is involved in interbacterial competition.
Background
The cell envelope protects bacteria from their surroundings. Breaches in its integrity or defects in its assembly are sensed by signal transduction systems, allowing cells to rapidly adjust. The Rcs phosphorelay responds to outer membrane (OM)- and peptidoglycan-related stress in enterobacteria. An OM lipoprotein, RcsF, is required for sensing and transducing the stress signals to downstream components in yet unidentified manners.

Objectives
We set out to discover how RcsF detects stress and activates the Rcs signalling cascade.

Methods
We used a combination of methods, including site-specific photo-crosslinking, co-immunoprecipitation and fluorescence microscopy.

Conclusions
We found that RcsF interacts with BamA, the major component of the OM beta-barrel assembly machinery. BamA funnels RcsF to the beta-barrel OmpA, allowing RcsF to reach the cell surface in an unresponsive-to-stress conformation. This spatially separates RcsF from the inner membrane-located IgaA, which we show is the next component of the cascade. Stress conditions prevent RcsF from interacting with BamA, allowing it to reach IgaA. Thereby the BamA-RcsF interaction acts as a checkpoint for lipoprotein transport to the cell surface, which we propose is mediated by BamA itself.
Staphylococcus aureus colonizes the moist squamous epithelium of the nasal cavity of about 20% of the population where it resides as a harmless commensal. It can cause infections ranging from superficial skin and soft tissue infections to those that are invasive and seriously life threatening. The surface of S. aureus is decorated with up to 24 cell wall anchored (CWA) proteins that are covalently linked to peptidoglycan by sortase. CWA proteins perform a variety of functions viz adhesion to host cells and tissue, evasion of innate and adaptive immune responses, invasion of host cells, acquisition of iron from haemoglobin, biofilm formation and stimulating inflammation. They can be classified into six groups based on structural and functional characteristics. The largest class are called microbial surface components recognising adhesive matrix molecules (MSCRAMMS) which have at their N terminus an A domain containing two adjacent subdomains each comprising IgG-like folds. MSCRAMMS bind to short peptide sequences within host proteins by the dock latch lock mechanism. Clumping factor A is the archetypal MSCRAMM which binds to the C-terminus of the gamma-chain of fibrinogen. The N-terminal A domains of MSCRAMMS are displayed on the cell surface aided by unfolded flexible stalks. In the case of ClfA and many other MSCRAMMS the stalk comprises serine-aspartate dipeptide repeats. In fibronectin binding proteins the linker for the A domain comprises fibronectin binding repeats. Several CWA proteins are being tested as components of vaccines aimed at reducing S. aureus infections
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Pathogenicity and cell surface

Role of the ESX-5 protein secretion system in functioning of the mycobacterial cell envelope and in virulence

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Mycobacteria use type VII secretion (T7S) systems to secrete proteins across their complex cell envelope. Pathogenic mycobacteria, such as the notorious pathogen Mycobacterium tuberculosis, have up to five of these secretion systems, named ESX-1 to ESX-5. At least three of these secretion systems (ESX-1, ESX-3 and ESX-5) are essential for mycobacterial virulence and/or viability. Elucidating T7S is therefore essential to understand the success of M. tuberculosis and other pathogenic mycobacteria as pathogens, and could be instrumental to identify novel targets for drug- and vaccine-development. Recently, significant progress has been achieved in the identification of T7S substrates. We have identified a general motif (YxxxD/E) that is required for secretion. Surprisingly, this motif does not determine system specificity. Secretion of substrates is also dependent on the function of dedicated chaperones. We are currently determining whether these chaperones are involved in system specificity. In addition, a start has been made with unraveling the mechanism of secretion and the structural analysis of the different subunits. We have identified the membrane translocation complex of both the ESX-1 and ESX-5 system. Finally, one of the T7S systems, ESX-5, is only present in slow-growing mycobacterial species and is responsible for the secretion of dozens of substrates. We have recently shown that ESX-5 substrates play a role both in virulence and in nutrient uptake. The latter characteristics could be linked to its exclusive presence in slow-growing mycobacteria.
Saccharomyces cerevisiae is a robust industrial micro-organism and a popular platform for metabolic engineering. Breath-taking advances in synthetic biology have greatly increased the pace and ease with which biochemical pathways towards non-native compounds can be expressed in this yeast. For environmentally and economically sustainable production of such compounds, it is essential that product yields on (sugar) substrate are as close to their thermodynamic maxima as possible. This not only requires ‘tuning’ of the non-native product pathways, but also optimization of their interaction with central metabolism, for example for the generation of product precursors and for redox balancing. While central metabolism of S. cerevisiae and its regulation have been shaped in aeons of natural evolution to improve competitiveness in natural environments, several key aspects of its redox and energy metabolism are clearly suboptimal for industrial application. In this presentation, I will discuss novel metabolic engineering strategies to address these limitations and, thereby, enable improved yields of existing and novel products of engineered S. cerevisiae strains.
Coordination of yeast metabolism through phosphorylation

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While our knowledge on the topology of regulation network steadily increases, we understand much less about the functionality of individual regulation events and their quantitative relevance for controlling a given biological function and eventually a phenotype. Focussing on metabolism, we ask which mechanisms *Saccharomyces cerevisiae* employs to make decisions on the use of different pathways. Although obviously important, transcriptional regulation does not appear to be the major mechanism to control the activity of metabolic pathways in bacteria and yeasts (1). By combining quantitative proteomics, P-proteomics and flux measurements, we instead demonstrated in a pilot study that enzyme phosphorylation, in contrast to transcriptional regulation, frequently contributes to the in vivo control of pathway fluxes in bakers yeast (2). To more broadly investigate the role of phosphorylation in controlling enzyme activity, and thus pathway fluxes, we performed large-scale metabolomics on 118 kinase and phosphate deletion mutants. Combining the metabolome with P-proteomic data, we predicted functional phosphorylation for about 50 metabolic enzymes (3). Different from bacteria, the yeast kinase network is rather interconnected with often multiple kinases acting on a given enzyme, a redundancy that in many cases might obscure functionality assessment. Here I will discuss how we tackle the problem of functionality by two approaches: i) integration of dynamic metabolomics and P-proteomics data, focussing on TOR signalling, and ii) large-scale point mutations in P-enzymes that either mimic permanent phosphorylation (serine to glutamate) or prevent phosphorylation at a given site entirely (serine to alanine).

References:


Many yeasts used in industrial fermentations are not pure species but hybrids between 2 or more species. Their advantage is that they can combine useful properties from each of the parental species. The disadvantage for strain improvement is that these hybrids are sterile and therefore not amenable to genetic analysis. Complex trait analysis in these is virtually impossible due to this sterility. The most famous yeast hybrid is the lager yeast, *S. carlsbergensus*, also known as *S. pastorianus*, which actually arose from at least two independent hybridization events between *S. cerevisiae* and another *Saccharomyces* species. Genetic studies and strain improvement has generally been limited to single gene mutations and analysis of rare spore progeny. The recent discovery of the second parental species, *S. eubayanus*, has provided the opportunity to create new hybrids using diverse parental strains in order to enhance genetic analysis. These new hybrids still suffer from the problem of sterility however. There are other hybrids used in industry such as the *S. cerevisiae* by *S. kudriavzevii* hybrids used in wine fermentation. These are less well characterized than the lager strains but share many of the features, such as colder fermentation temperatures, and issues, such as sterility and genome stability. There are also many other hybrids isolated from nature that may or may not have industrial uses. We have recently utilized a tetraploid intermediate strategy to overcome the sterility of hybrids, which allows us to both genetically study exiting hybrids as well as create new hybrids such that quantitative genetic analysis can be done. We have successfully restored fertility in a wine hybrid and are exploring the industrial potential of new hybrids, incorporating all the extant species.
Lignocellulosic bioethanol – from laboratory proof-of-concept to full scale plant

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Lignocellulosic feedstocks are considered to be of great economic and environmental significance for sustainable production of valuable fermentation products. For cost-effective and efficient industrial processes, complete and fast fermentation of all biomass sugars is a prerequisite. In this respect, the main challenge emerging from the use of lignocellulosic feedstocks for ethanol production by *Saccharomyces cerevisiae* is the efficient fermentation of the pentose sugars xylose and arabinose, as these sugars cannot be utilized by natural *S. cerevisiae* strains. Another significant challenge is the inevitable presence of inhibitors, such as furfural and acetic acid, which are formed during pretreatment and hydrolysis of the feedstocks and severely inhibit yeast growth and product formation.

DSM has developed advanced yeast strains by introducing heterologous pathways into robust *S. cerevisiae* hosts which resulted in the ability to ferment xylose and arabinose in lignocellulosic hydrolysates. Subsequently, by the application of evolutionary engineering the total time required to ferment hexoses and pentoses in lignocellulosic sugar mixtures was significantly reduced.

In order to take these developments to the next level, a toolbox for the new generation of advanced yeast strains is in development, in which the main challenges are further addressed by accelerating pentose fermentation through engineered pentose transport, as well as converting lignocellulosic inhibitors and biorefinery waste streams into additional ethanol. A roadmap to implementation of commercial strains in a full scale plant will be presented.
The phyllosphere microbiota: responses to and impacts on plants

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The aerial parts of the plants, which are dominated by leaves, represent one of the largest terrestrial habitats for microorganisms. This habitat, called the phyllosphere, is occupied by a diverse community of microorganisms, which is important for plant health and growth. Most of the phyllosphere inhabitants are not well investigated; however, there is a growing interest to study commensal bacteria to elucidate their interactions with the plants, among each other and to learn how they withstand the hostile conditions of their habitat. A predominance of Proteobacteria, Actinobacteria and Bacteroidetes living in the phyllosphere of numerous plants has been revealed, while metagenomics and metaproteomics approaches gave insights into the general bacterial adaptation strategies to the phyllosphere. Complementary to these cultivation-independent approaches we established a comprehensive strain collection which covers a broad diversity of strains colonizing the model plant Arabidopsis thaliana. Targeted studies with model strains allowed us to identify metabolic traits important for plant colonization and to uncover a novel bacterial regulatory system essential for plant colonization which is responsible for the general stress response in Alphaproteobacteria. The establishment of a gnotobiotic system led to the identification of plant probiotic effects of commensal bacteria and candidate genes for plant protection against bacterial pathogens. Moreover, the experimental system paired with synthetic bacterial communities helped identifying plant genes involved in shaping the bacterial community structure.
Metalliferous plants including (hyper)accumulators with tissues containing extremely high trace elements (TE) concentrations are associated with complex microbiomes, which are likely to be important for plant growth under such adverse conditions. Our studies addressed the ecology and microbial community behavior under different conditions as well as the role of plant-associated bacteria for the TE mobilization / availability and plant uptake.

We analyzed microbiomes associated with Zn-accumulating plant Salix species, which are applied for phytoextraction applications. Plants were associated with complex rhizosphere microbiomes, which correlated with the plant genotype and which were particularly rich in Actinobacteria. Bacteria isolated from Zn accumulating S. caprea showed highly different effects on TE mobilization. Several strains decreased bioavailability, whereas others, particularly Actinobacteria such as Microbacterium spp. strains, increased bioavailability. The mechanism of TE mobilization is poorly understood, however, we hypothesize that microbial secondary metabolites (such as siderophores) are involved in the mobilization process. We therefore have analyzed the genomes and metabolomes of Zn/Cd mobilizing actinobacterial strains to further elucidate the mechanisms involved in TE mobilization and plant uptake.
Back to the roots: microbiology and chemistry at the root-soil interface

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Back to the Roots: microbiology and chemistry at the root-soil interface

Plant roots are colonized by an astounding number of (micro)organisms that can reach cell densities much greater than the number of plant cells. Many studies have shown that members of the rhizosphere microbiome can have profound effects on seed germination, seedling vigour, plant growth and development, nutrition and tolerance to pests, diseases and abiotic stress. For the vast majority of rhizosphere microorganisms, however, there is limited knowledge on the mechanisms involved in modulation of plant growth and plant health. Here, new results are presented on how rhizosphere bacteria and fungi impact on root architecture, root growth and plant tolerance to infections by soil-borne pathogens. For the rhizosphere bacteria, results showed that representatives of the γ-Proteobacteria protect plants from pathogen infection by the production of chlorinated peptides and alter root architecture and plant growth via modulation of the plants’ sulfur metabolism. For the fungi-plant interactions, our results showed that specific soil-borne pathogenic fungi induce changes in root architecture and plant growth via volatile organic compounds. An overview will be given on the wealth of yet unknown functions and metabolic potential of the rhizosphere microbiome.
Fungi from the Phylum Glomeromycota establish one of the most ancient associations with plant roots helping plants to thrive and survive under conditions of mineral deprivation. These fungi are particularly good at improving plant phosphate nutrition while in return, plants provide fungi with carbohydrates. This mutualistic symbiosis is characterized by the formation of fungal specialized structures in the cortex, arbuscules, key to the nutrient exchange and giving name to the association: arbuscular mycorrhizal (AM) symbiosis. AM fungi, like all biotrophic fungi, are committed to keep plant defenses at bay and prevent the mutualistic association to become parasitic. It is only recently that the mechanisms of how AM fungi overcome the surveillance system of the plant and manipulate their metabolism are starting to be understood. In general AM fungi use at least three mechanisms to achieve that: i) avoiding being recognized; ii) suppressing defense responses elicited after recognition iii) using a specific dialogue that allows manipulation of the plant cell program. We were first to show that similar to pathogenic microorganisms, AM fungi do have and are able to deliver effector proteins to the plant to counteract MAMP-triggered immune responses (Kloppholz et al., 2011). With the recent outcome of the first genome of an AM fungus, the effector repertoire is starting to be envisaged, and the presence of LysM containing proteins, Crinkler-like effectors and secreted proteins with NLS and DNA modification motifs points out that the similarities with pathogenic fungi are likely to be more than previously anticipated. The aim of our work is to investigate how plant roots and arbuscular mycorrhizal fungi (AMF) perceive signals from each other and establish the molecular dialogue that permits a long-lived and harmonious symbiosis.
We have reported direct connections between the SOS response activation and HGT mechanisms such as conjugation and natural transformation, but also with gene capture in integrons. We also found that most antibiotics at sub-inhibitory concentrations induce the SOS response in *Vibrio cholerae*, and in other Gram-negative pathogens such as *Klebsiella pneumoniae*. This coupling enhances the potential for cassette swapping and capture in cells undergoing stress, while freezing the cassette arrangement in steady environments, but it also favors any mechanisms leading to resistance development through mutation. We have now undertaken the characterization of the different partners and pathways that are involved in these connections, in order to better understand these physiological responses and propose innovative strategies against the rise of resistance.
Gram-negative bacterial cells show copious evidence of evolution through the horizontal acquisition of novel genetic sequences with a high A+T content. In *Salmonella enterica* serovar Typhimurium, the majority of the main virulence genes fit this profile. These genes allow the bacterium to gain access to privileged sites in the host, such as the interior of epithelial cells, or to avoid being killed by components of the host defences such as macrophage. Understanding how such horizontally acquired genes are integrated into the regulatory regime of the bacterium presents a major challenge. We will provide evidence that in the case of *S. Typhimurium* regulatory integration has been achieved through combining horizontally acquired regulatory genes with regulators encoded by the ancestral genome with both operating on the virulence genes through a mechanism that exploits the special structural characteristics of A+T-rich B-DNA.
Viruses that infect bacteria – termed phages – are recognized vectors of lateral gene transfer between microbial cells. Bacteria and their parasitic phages are co-evolving in a constant arms race, yet their interaction may include also mutualistic aspects. The beneficial contribution of phage-mediated gene transfer to the host fitness has been documented in diverse species and environments. Here we study the contribution of phage-mediated gene transfer to the lateral component of microbial evolution using phylogenomic transduction networks. The networks are composed of donors, phages, and recipients that are connected by 23,014 lateral gene transfer events reconstructed from genomic data. Large-scale structural differences among the stable and transient sub-networks reveal the differential contribution of temperate and lytic phage-bacteria interactions to transduction dynamics in Nature. Our evolutionary reconstruction reveals high phage-host specificity in stable interactions, while transient interactions may occur with multiple hosts. Edge weight distribution in the network suggests that gene uptake from transient hosts into the phage genome typically includes a single gene, while gene transfer into stable hosts usually comprises several genes. Yet, transient interactions constitute an important contribution to the global network structure by connecting among clusters of stable hosts. Self-donor recipients constitute a significant portion of the network (21%), implicating transduction as an important gene duplication mechanism during microbial evolution. Furthermore, donors and recipients that have a similar genome sequence and similar codon-usage are more highly connected by transduction events than the expected by chance. This indicates that donor-recipient whole genome similarity is an important factor that shapes the transduction network connectivity pattern. Our results demonstrate the implication of phage-bacteria co-evolution to phage-mediated gene transfer during microbial evolution.
Horizontal gene transfer, evolution

Co-transfer of hypermutagenesis and symbiotic genes drives the evolution of legume endosymbionts

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Horizontal gene transfer (HGT) is an important mode of adaptation and diversification of prokaryotes and eukaryotes and a major event underlying the emergence of bacterial pathogens and mutualists. Yet it remains unclear how complex phenotypic traits such as the ability to fix nitrogen with legumes have successfully spread over large phylogenetic distances.

To get an insight into the evolutionary mechanisms that facilitated rhizobium diversification, we experimentally replayed the evolution of rhizobia. Following introduction of the symbiotic plasmid of Cupriavidus taiwanensis, the Mimosa symbiont, into pathogenic Ralstonia solanacearum we challenged transconjugants to become Mimosa symbionts through serial plant-bacteria co-cultures.

Using whole genome sequencing we showed that co-transfer of imuABC error-prone DNA polymerase genes with key symbiotic genes accelerates the evolution of a soil bacterium into a legume symbiont. We demonstrate that a mutagenesis imuABC cassette encoded on the C. taiwanensis symbiotic plasmid triggered a transient hypermutability stage in R. solanacearum transconjugants that occurred before the cells entered the plant. The generated burst in genetic diversity accelerated symbiotic adaptation of the recipient genome under plant selection pressure, presumably by improving the exploration of the fitness landscape. Finally, we show that plasmid imuABC cassettes are over-represented in rhizobial lineages harboring symbiotic plasmids. Our findings shed light on a mechanism that may have facilitated the dissemination of symbiotic competency among a- and b-proteobacteria in natura and provide evidence for the positive role of environment-induced mutagenesis in the acquisition of a complex lifestyle trait.
We employ nL-sized gel-carriers (nanoliter reactors or NLRs) kept in suspension arrays as growth and reaction compartments. Single library cells are embedded into NLRs and proliferated to microcolonies of a few 1'000 cells required for increase of the signal-to-noise ratio. The phenotype (e.g. biocatalyst productivity) is indicated by genetically encoded or chemical fluorescence reporters. Putative positive strains are
detected on the basis of the reporter signals and isolated by an appropriate technique such as large particle flow-cytometers.

We will provide a detailed overview of the technical elements required for nLR-screenings and present two application examples: First, screening for antibacterial peptides secreted by Lactobacillus libraries at rates of 1E5 / day. Second, identification of Bacillus subtilis strains catalyzing the conversion of cellobiose to the industrial product vitamin B2 at an elevated yield.

Both protocols are used on a regular basis. While the peptide libraries screens are performed by us, the B2 protocol has successfully been transferred to a professional environment thereby further accentuating the robustness of NLR technology.
The chemical make-up of microbiology is incredibly complex. It is already difficult to analyze the molecular make up of one organism, let alone 100s or 1000s of different organisms found in microbial communities? How can we look at the molecular make-up of all organisms at once? How many molecules do they have in common? How many are different? How do we retain the collective mass spectrometry knowledge of the community? As mass spectrometers are becoming faster and more sensitive we can detect a lot of molecular information. There are now mass spectrometers that can analyze 10,000 samples a day. However, there is no infrastructure to analyze this amount of molecular information or to correlate this information to other Big Data generating approaches such as sequencing. On average, however, only 1-5% of all the molecular information that is collected by mass spectrometry can be annotated. It is simply too much information for one person or lab to analyze this information with the existing tools that are available. In this lecture we will explore the strategies for organizing and visualizing the massive amount of information. Topics such as molecular networking, crowd source molecular analysis, 3D topographical mapping, MS/MS rarefaction, large scale mass spectrometry based genome mining, cross correlative analysis with molecular networking and pattern based genome mining will be covered.
Fungi possess inducible pathways that are activated in responses to stress that can produce highly bioactive defense compounds upon elicitation that need to be identified and investigated further for an evaluation of their biological properties and for a fundamental understanding of the chemical events that trigger various interactions between microorganisms. Metabolomics represents an ideal approach to highlight biomarker induction in such complex extracts. In this context the confrontation of various strains of human and plant pathogen fungi have been studied by an MS-based metabolomics approach that take advantages of UHPLC-TOF-MS fingerprinting and high resolution metabolite profiling for an efficient localization of stress-induced biomarkers. New chemometric algorithms that serve at the detection of MS features in complex mixture of fungal metabolomes have been devised. The de novo structure identification of novel stress-induced biomarkers was assured by subsequent LC-MS targeted microisolation and microflow NMR analyses. Various sampling, MS-based and data mining strategies were compared to highlight at best de novo metabolite stress-induction at the confrontation zone between microorganisms. Comparison of replicates obtained from miniaturized solid media coculture produced in the 12-well plate format were found to be the best compromise for generating reproducible series of samples for further metabolite profiling. In some of the co-culture studied novel sulfated polyketides were evidenced in others quinonic pigments were highlighted to be strongly induced. No clear correlation between confrontation morphological patterns and metabolite induction could be made. In all cases however significant induction mechanisms were evidenced at the molecular level thanks to the advanced data mining approaches applied. De novo structure identification of the targeted metabolite was be performed by HRMS and micro NMR after careful sample enrichment and chromatographic gradient transfer and MS-directed purification.
**FEMS-1960**  
*Cell polarity and virulence*

**Brucella abortus cell cycle in culture and inside host cells**  
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*Brucellae* are responsible for a worldwide zoonosis called brucellosis. These bacteria are able to invade a large range of host cell types. The intracellular trafficking of these bacteria is characterized by two successive steps: a first step in the endosomal compartments, where the bacteria do not grow, and a second step during which the bacteria proliferate massively, usually in the endoplasmic reticulum. Using fluorescent reporters for the monitoring of growth and chromosomal segregation at the single cell level, we showed that the cell cycle of *Brucella abortus* is blocked at the G1 stage during infection of HeLa cells and RAW 264.7 macrophages. This blockage occurs during the first stage of the infection. The cell cycle is resumed at the end of this first trafficking step, when bacteria are still in the endosomal compartments. These data show that cell cycle progression is coordinated with the intracellular trafficking of the bacterium. We thus investigated the role of a conserved transcription factor and regulator of cell cycle progression called CtrA. The predicted and experimental (ChIP-seq) CtrA regulons are consistent with a major role of CtrA for the control of cell growth, cell division and DNA methylation. Interestingly, CtrA is also proposed to control the segregation of chromosome II. Indeed, the *B. abortus* genome is split in two chromosomes, the segregation of newly replicated chromosome I origins occurring before the segregation of chromosome II origins. We propose that *B. abortus*, similarly to the model bacterium *Caulobacter crescentus*, is characterized by ordered cell cycle processes, such as pole maturation and successive initiations of chromosomal replication.
Agrobacterium tumefaciens is best known for its ability to genetically modify plants through transfer and integration of a segment of DNA (T-DNA) into the plant genome, causing the disease called crown gall. Despite extensive studies of T-DNA transfer, little is known regarding the initial processes at the plant surface. A. tumefaciens forms dense biofilms on both abiotic and biotic surfaces which may increase the overall efficiency of transformation while affording protection against host defense responses. One of the primary factors mediating surface attachment in A. tumefaciens is called the unipolar polysaccharide (UPP), an adhesion produced at a single pole of the cell following surface contact. The UPP is one of several polarly localized cellular structures reported in A. tumefaciens including a tuft of unipolar flagella, and components of the Type IV secretion system (T4SS) that exports T-DNA. The cellular asymmetry is even more profoundly reflected in a polar mode of cell division for A. tumefaciens, in which cells multiply by budding and utilize a zonal mechanism of cell wall growth. Surface-attached A. tumefaciens cells generate daughter cells from their unattached pole, from which the flagella are also elaborated. The cellular asymmetry is under the control of a complex regulatory cascade designated the Control of Division and Development (CDD) pathway. Several related sensor kinases (PleC, DivJ, PdhS1 and PdhS2) and their cognate response regulators (DivK and PleD) converge on control of the CckA-ChpT-CpdR/CtrA CDD central circuit, and provide cell pole-specific modulation of division and polar development. UPP targeting to the old pole of the target cell is dependent on the PodJ protein, thought to function as a localization factor, whereas T4SS components concentrate at the pole independently of PodJ.
Mycobacterium tuberculosis infects billions of people worldwide and kills more than 1.5 million per year. TB remains extremely difficult to treat with antibiotics, requiring months to years of therapy for cure. The variable course of disease and treatment response suggests that functionally heterogeneous populations of mycobacteria respond differently to stress. Using a quantitative single-cell approach, we show that mycobacteria deterministically generate diversity in their growth characteristics through an asymmetric growth pattern. Coupled with a cell cycle regulated by time and not size, this asymmetry creates subpopulations of cells with distinct growth rates and cell sizes that are differentially susceptible to clinically relevant classes of antibiotics. Thus, the growth pattern intrinsic to mycobacteria deterministically creates a diverse population structure that may underlie phenotypes previously thought to be controlled by external stressors. We have also observed variation among microcolonies in antibiotic susceptibility that cannot be explained by growth pole age. Armed with new reporters of cell state and mathematical models, we seek to quantitatively characterize mycobacterial subpopulations with distinct behaviors arising from asymmetric growth and division.
It takes two to tango - the interplay between two nanomachines drives horizontal gene transfer in Vibrio cholerae

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Vibrio cholerae is considered an important model organism for elucidating virulence regulation of facultative pathogens. In this context, the involvement of quorum sensing (QS) and of the type VI secretion system (T6SS) have been extensively studied even though the T6SS is been considered to be "silent" under laboratory conditions in pandemic V. cholerae O1 El Tor strains. Much less is known about the bacterium’s lifestyle in its natural environment, where it often associates with small crustaceans and their chitinous surfaces.

Here, we show that the chitin- and QS-dependent competence regulon of diverse pandemic and epidemic V. cholerae strains includes the T6SS-encoding gene clusters and that the T6SS contributes to enhanced horizontal gene transfer by means of natural transformation. We used live cell imaging to visualize components of the two competence-induced nanomachines of V. cholerae, namely the DNA uptake complex and the T6SS. Notably, such live-cell time-lapse imaging revealed that competence-mediated induction of the T6SS enhances horizontal gene transfer by deliberately killing neighboring non-immune cells and taking up their DNA via the competence-induced DNA uptake machinery.
Experimental mixed species biofilms: A novel approach to evaluating interactions at a community level

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FEMS Conference presentation

Speaker: Prof. Staffan Kjelleberg

Abstract title: Experimental mixed species biofilms: A novel approach to evaluating interactions at a community level

Our understanding of the mechanisms of biofilm development has largely been derived from studies on biofilm populations, yet biofilms predominantly exist as communities. This lack of investigation into multispecies consortia reflects the relative ease by which the biofilm life cycle can be experimentally interrogated for single species systems and the difficulty to maintain experimentally tractable biofilm community systems. We have employed experimental mixed species biofilm communities to bridge this gap, also allowing us to explore mechanisms in more ecologically relevant systems and test theories from eukaryote ecology. The biofilm life cycle of two biofilm community systems – a defined three species biofilm community and a phylogenetically highly diverse biofilm granulation system – will be addressed. For the former, the establishment of a mixed defined community of *Pseudomonas aeruginosa*, *Pseudomonas protegens*, and *Klebsiella pneumonia*, has been employed to demonstrate community specific contributions to biofilm development, resistance to chemicals and predation, and variant formation by the individual member species. These findings report on mechanisms that regulate cross species interactions but which are not available for monospecies biofilms formed by either of the three members. The second experimental biofilm community system, exploring the life cycle of high species diversity granular biofilms reveals that quorum sensing signaling is a distinct community regulatory process. Here, signal producers and degraders are phylogenetically diverse and dissimilar, while a minority of the community members are capable of entertaining both signaling synthesis and quenching, as is the case for our understanding of QS signaling biology in population based model biofilms. Crucially, the granular biofilm life cycle is driven by differentiated community signaling that reflects community shifts in phylogeny, signaling synthesis and quenching.
Clinical Implications of Biofilm-Associated Polymicrobial Infections

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The clinical significance of microbial interactions in a host particularly those between opportunistic pathogens such as Candida albicans and Staphylococcus aureus remains largely understudied mainly due to lack of suitable animal models. To that end, we developed mouse models of co-infection to investigate the clinical implications of mixed species biofilms on host tissue and implanted catheters. In the first model, animals were orally infected with C. albicans and following development of oral candidiasis animals were exposed to S. aureus through drinking water. Findings from this model demonstrated that the extensive tissue damage caused by invading C. albicans hyphae provided an inlet for the typically non-invasive S. aureus to disseminate to the kidneys causing high morbidity and mortality. Additionally, a mouse subcutaneous catheter model was used to study in vivo-grown biofilms where co-infected catheter fragments were implanted subcutaneously in the animals. Following 6 days, analysis of explanted catheters revealed the formation of a thick biofilm within the catheter lumen consisting of both species embedded in a matrix. Further analysis indicated increase in S. aureus produced eDNA in mixed biofilms and enhanced tolerance of S. aureus to vancomycin. Combined the findings from these studies demonstrated that the onset of oral candidiasis may constitute a risk factor for disseminated bacterial disease. Further the co-adherence of C. albicans and S. aureus to abiotic surfaces results in formation of resistant biofilms on indwelling medical devices. These novel findings demonstrate the gravity of C. albicans-S. aureus interactions in the host, warranting awareness in terms of prophylactic therapeutic measures.
Candida albicans-Staphylococcus aureus intra-abdominal infections: at the crossroads of inter-Kingdom virulence and host response

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The polymorphic fungus Candida albicans and the ubiquitous bacterial pathogen Staphylococcus aureus are common co-infecting microbes during intra-abdominal infections (IA), often resulting in high morbidity and mortality. Using a murine model of IAI, our previous work demonstrated that synergistic host inflammation (specifically, prostaglandin E2) is associated with mortality. However, the microbial virulence factors contributing to infectious synergism remain undefined.

Previous studies suggested that staphylococcal toxins were associated with poor outcome during murine IAI. Thus, we hypothesized that C. albicans may augment the expression of S. aureus toxins during co-infection. To test this hypothesis, cultures of S. aureus, C. albicans, or both organisms together were grown under planktonic or biofilm growth conditions. Cell-free supernatants from polymicrobial cultures showed increased hemolysis in vitro. Use of an isogenic S. aureus ΔagrA mutant confirmed that enhanced hemolysin production was agrA-dependent. Use of a GFP-reporter plasmid demonstrated enhanced agr activity in co-culture as compared to monomicrobial growth, and qPCR and Western blot analysis of agr-regulated genes further supported these results. Lastly, mice were intraperitoneally inoculated with sub-lethal doses of C. albicans+S. aureus, C. albicans+S. aureus (ΔagrA), or S. aureus alone and monitored for morbidity and mortality. Alarmingly, 80% of the mice co-inoculated with C. albicans+S. aureus (WT) succumbed by d 1 post-inoculation, while only 30% of the mice co-inoculated with C. albicans+ΔagrA succumbed to the infection (d 8 post-inoculation). All mice inoculated with JE2 alone survived throughout the infection time course. Furthermore, assessment of α-toxin in the lavage fluid by immunoblot and ELISA confirmed increased toxin during co-infection, despite similar bacterial burden as monomicrobial infection.

In summary, C. albicans can augment S. aureus toxin production, potentially explaining the devastating pathology of fungal-bacterial IAI.
Contribution of bacterial volatile molecules to the dynamics of polymicrobial bacterial biofilm communities

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Bacteria produce and sense a high diversity of signals and cues to adapt their behaviors to changing environments. Among them, volatile compounds produced by fungi and plants were shown to modify bacterial phenotypes and activities. However, besides being perceived as odors, attractive scents or pollutants, the influence of bacterial volatile molecules on bacterial biology itself remained poorly explored. I will discuss how some bacterial volatile compounds diffusing in heterogeneous environments, such as soil, organic tissues and microbial mats can mediate airborne chemical interferences and contribute to shape polymicrobial bacterial communities.
An evolutionary view of the Holobiont theory and its implications for host development

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The mammalian microbiome is formed out of a maternal hand-over of microbes, diet and lifestyle that will influence host physiology throughout life. Moreover, microbes and its host also share, in part, their biological and biochemical needs. This assumption forms the basis of the Holobiont theory. This theory of evolution proposes that the object of natural adaption and selection is not the host (its genes and organs), but the host together with its microbiome (all genes and its metabolites) - The Holobiont. When the host – The Holobiont - is challenged by dramatic changes, such as alteration in dietary intake, malnutrition, reduced physical activity or unwanted severe stress, a Holobiont will employ strategies and adaptation mechanisms to handle these changes including signals transmitted to the microbiome community. The microbiome in turn, will reciprocate and transmit signals to its host. The last decades of reductionist research aiming to decipher the underlying mechanisms responsible for the epidemic development of lifestyle related diseases, by use of Genome Wide Association Studies has not considered the Holobiont perspective and consequently, the adaptation strategies developed by the microbiome and its signals transmitted to the host has not be included in the analysis. In my presentation I will discuss the Holobiont concept and present data which support how a host respond to this bilateral microbiome-host communication related to development and metabolic homeostasis.
The human gut is inhabited with trillions of bacteria, gut microbiota, that have co-evolved with us and affect our physiology within and outside the gut. The gut microbiota has recently been suggested as a novel contributor to obesity and related comorbidities, such as type 2 diabetes (T2D) and cardiovascular diseases (CVD). We recently found that the gut microbiota is altered in patients with CVD and T2D and that we can classify patients and T2D patients based on the microbiota. Using germ-free mice we have causally linked the gut microbiota to obesity and insulin resistance and have recently found that the gut microbiota modulates adipose inflammation, bile acid signaling, and enteroendocrine cell function. However, the underlying mechanisms(s) by which the gut microbiota induces signalling is yet to be dissected. By combining defined microbial communities, multi-omics approaches with genetically modified mice we are beginning to clarify host microbial interactions and their metabolic responses.
Genome sequences as nomenclatural type material for novel taxa

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Major goals of the Bacteriological Code are to provide all prokaryotic species with a single name and insure that each name refers to a single species. Currently, these goals are achieved in part through designation of type strains, which are viable cultures deposited in two public culture collections. These cultures then become the reference material for comparison to the type strains of other validly described species.

The genome sequence provides sufficient information to satisfy the goals of the Bacteriological Code for type material. Based upon genome sequence similarities, it is possible to determine if two strains are members of the same species. The genome sequence also provides sufficient information about the growth properties, physiology, metabolism, and lifestyle to support polyphasic classifications. Thus, the Bacteriological Code should be modified so that genome sequences from clonal populations or single cells are acceptable type materials upon deposition of DNA to insure authenticity.

This change will allow naming of taxa not readily cultivated by service culture collections and uncultured taxa. Moreover, the current system was developed when the number of type strains was in the thousands. The number of type strains is now greater than 12,000 and continues to increase. Culture collections are unfairly burdened with the task of maintaining and distributing this large number of strains. Allowing genome sequences to serve as type material will allow culture collections to reallocate resources to the most important and valuable cultures.
Applying an evolution-based species concept for the delineation of bacterial species: The Pantoea example

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Bacterial species are commonly defined by applying a set of predetermined criteria, such as DNA:DNA hybridization values, 16S rRNA sequence similarity and phenotypic and chemotaxonomic comparisons. Alternative genome-based similarity criteria have also been proposed to define bacterial taxa. We have previously shown that the metrics based on genome data support all of the validly described Pantoea species. However, the question remained whether the same species would have been recognised when taxa were delineated using an evolution-based species concept instead of fixed or quantitative criteria. Here we utilized Pantoea and the available genome data to investigate whether an evolution-based concept, in which species represent unique and genealogically exclusive groups of isolates on phylogenetic trees, would allow for delineation of the known species.

The core genomes (genes present amongst all member of the group) for Pantoea species were determined. These genes were subjected to phylogenetic analyses as they collectively represent the dominant evolutionary signal that depicts vertical descent. The core gene content was also characterized in terms of general cellular functions.

The Pantoea core genome consisted of approximately 2500 genes and differed markedly from the genomes of other closely related genera. Similar phylogenies were inferred for the various functional sets of genes. Although slight changes in sister relationships were observed for some of the species, the examined Pantoea species all represented exclusive genealogical groups. Our results thus indicated that a genome-based comparative and evolutionary approach could be a preferred alternative to fixed criteria when delineating coherent bacterial taxa.
Background:

Molecular tools applied to understand microbial diversity have evidenced the very large extent of undescribed taxa. The estimated richness exceeds in orders of magnitude the current ~12,000 classified species. However, there is a need to find faster ways to catalogue the biosphere’s diversity, and try to overcome the major problem in classification that is bringing the type material into pure culture.

Objective:

For this study we have measured the extent of hitherto discovered putative taxa in 16S rRNA gene databases using taxonomic thresholds that have been previously calculated with all cultured Bacteria and Archaea. In addition, we have evaluated the current species definition and its application to recognize uncultured microbial populations thriving in natural environments and detected through metagenomic approaches.

Conclusions:

The current species definition for prokaryotes is strongly based on its circumscription by genome sequence comparisons and genetic distances of housekeeping genes, together with a discriminative phenotypic property. The set of environmental 16S rRNA gene sequences in public repositories matched about 250,000 species, 80,000 genera and 1,300 phyla, and the predicted total number of species in the biosphere ranges from $10^6$ – $10^7$. In addition, the new sequencing technologies allow retrieving individual population’s (meta)genomes of uncultured organisms that can be used to understand their phylogeny and genomic uniqueness, and predict their metabolism as for cultured organisms. This will allow implementing metagenomics into the Candidatus proposals of organisms not showing conspicuous prominent characters, and speed up the description of uncultured organisms in a similar way to those in pure culture.
Type species in the genomic era of Mollicutes

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Since its inception, the International Code of Nomenclature of Bacteria has emphasized the importance of type material as a reference to be used when considering the identity of specimens. Nomenclatural types also permanently bear the name of the taxon. Investigators expect that types should exemplify the defining characteristics of the taxon as it exists within a hierarchy of superior taxa. The Code is sufficiently flexible to allow that types later realized to be unsuitable can be replaced with neotypes as knowledge advances.

The type of genus Mycoplasma, hence the Mycoplasmataceae, Mycoplasmatales and class Mollicutes is Mycoplasma mycoides subsp. mycoides strain PG-1T. Modern phylogenetic analyses situate M. mycoides in the distant order Entomoplasmatales, so if discovered today PG-1 would be doubtfully referred to Mycoplasma. Our objective was to explore genomic methods that might be used to identify candidates for neotype of the genus.

A well-chosen neotype might be expected to exhibit genomic consistency with others in its genus. JSpecies was used to calculate ANI values by pairwise comparisons of the complete genome sequences of 20 species representing six families within class Mollicutes. ANIm was calculated by the NUCmer algorithm of MUMmer and ANIb was calculated by reciprocal BLASTN. The alignment-free tetranucleotide frequency correlation coefficient was also calculated. The matrices were visualized as unrooted trees using PHYLIP, then reviewed by the ICSP Subcommittee on the taxonomy of Mollicutes. The Subcommittee unanimously agreed that the genomic trees were not congruent with the shared view of the taxonomic structure of the class. Although evidently not suitable for determination of supraspecific relationships, genomic analyses remain valuable for species-level circumscription of mollicutes.

Antibiotic resistance in the urban water cycle: origins, fate and risks

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The urban water cycle refers to the journey of water from the collection, disinfection and purification for human consumption to its return to nature as treated wastewater. Over this cycle, water is inhabited by different bacterial populations, some strictly environmental, others of human or animal origin, most of which considered contaminants. Multidrug resistant bacteria, sometimes harboring antibiotic resistance (AR) genes of clinical relevance, are commonly found in each of these aquatic environments.

It is estimated that AR bacteria are responsible annually for > 20000 deaths, in both the US and the EU. Although these values are registered in the clinical settings, it is recognized that the environment, in particular water, plays an important role on the dissemination of AR from and to humans.

Urban wastewater, either raw or treated and ready to discharge in the environment, presents a high prevalence of AR to “old” antibiotics, suggesting the cumulative effect of this form of contamination. Moreover, almost all AR genes of clinical concern can be detected in wastewater, suggesting their rapid dissemination.

In contrast, drinking water, in principle free of pathogens, is inhabited by an impressive diversity of environmental bacteria, some of which exhibit multidrug resistance phenotypes, probably intrinsic in those species. Although not posing an acute problem of AR dissemination, intrinsically AR bacteria may have also adverse human health effects.

Nowadays, it is difficult to assess the risks of the transmission, directly or indirectly, of AR bacteria from the environment to humans. A major limitation is related with the absence of surveillance and data sharing systems comprising both environmental and clinical data, allowing a global overview of the emergence and evolution of AR and how it can affect human health.

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The effluents of urban wastewater treatment plants (UWWTPs) are among the main point sources of antibiotic resistance pollution of surface waters. In several countries, disinfection process is the last step of UWWTPs to minimize the release of antibiotic resistant bacteria and genes (ARB&Gs) in the effluent. But in some country UWWTP effluent is disposed in the receiving water body without any disinfection treatment. Typical disinfection processes include chlorination and UV radiation but other disinfectants/oxidants such as ozone, chlorine dioxide and hydrogen peroxide are also used. Unfortunately, ARB&Gs have been detected in the effluents of UWWTPs raising the suspect that conventional disinfection processes (particularly chlorination and UV radiation) not effective in controlling antibiotic resistance spread into the environment. Therefore, in the last years non conventional disinfection technology such as that based on Advanced Oxidation Processes (AOPs) has been investigated as possible alternative to conventional disinfection processes for controlling antibiotic resistance spread.

In this work the results of disinfection experiments by chlorination, UV radiation and AOPs are summarized. The effect of disinfection processes on antibiotic resistant *E. Coli* (from the effluent of secondary treatment of an UWWTP) in terms of inactivation, bacterial regrowth and antibiotic resistance was evaluated. Moreover, the effect of an AOP (namely TiO2 photocatalysis) on the mechanism of antibiotic resistance transfer was also investigated.

AOPs can be more effective than conventional disinfection processes in the inactivation of ARB but further studies on the effect of AOPs on antibiotic resistance transfer are needed to better understand the potential of residual ARB and mobile genetic elements of transferring antibiotic resistance after disinfection treatment.
DETERMINING THE MINIMAL SELECTIVE CONCENTRATIONS OF ANTIBACTERIAL AGENTS IN COMPLEX AQUATIC BACTERIAL COMMUNITIES

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Background

A mixture of antibiotics and co-selective agents, such as biocides and metals, reach the environment through sewage effluents. It is unknown to what extent antibacterial agents select and co-select for resistance in aquatic ecosystems e.g. wastewater treatment plants and receiving waters.

Objectives

The aim was to determine the minimal selective concentration (MSC) of an antibiotic (tetracycline) and the minimal co-selective concentrations (MCCs) of a common biocide (cetyl trimethyl ammonium chloride; CTAC) in complex aquatic bacterial communities.

Methods

Treated sewage effluent (Gothenburg, Sweden) was diluted and used as inoculum to establish biofilms in flow-through systems containing different concentrations of antibacterial agents. Biofilms were screened for their resistance/tolerance using selective plating, characterization of isolates, metagenomics analyses (resistance genes and taxonomic composition), and pollution induced community tolerance (PICT).

Conclusions

Concentration-response curves for tetracycline were obtained for selective plating, tet-genes and PICT. Based on selective plating the MSC of tetracycline was 10ng/mL (p=0.009), which correlates well with an increased MIC50 value (concentration inhibiting 50% of the isolates). Increases of other resistance genes, ISCR2 and intI1 were observed with increasing tetracycline concentrations. Effects of tetracycline on taxonomic composition are under evaluation.
CTAC selects for self-tolerance and co-tolerance to chloramphenicol, tetracycline and erythromycin. The MCCs of CTAC as well as effects on resistance gene frequencies and taxonomy remain to be evaluated.

Knowledge of the selective and co-selective properties of tetracycline and CTAC generated here will facilitate the establishment of safe emission limits. This could guide actions to reduce risks for resistance selection in the environment.
EXPLORING PLASMID-BASED DISSEMINATION OF ANTIBIOTIC RESISTANCE GENES IN ENVIRONMENTAL MATRICES

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Background
Plasmids are important actors of antibiotic resistance gene dissemination that contribute to reduce the effectiveness of our therapeuthic potential.

Objectives
Our objectives are to identify permissive environments and/or conditions driving plasmid transfer in microbial communities.

Methods
Lately, we developed a sensitive molecular approach based on qPCR allowing the detection of rare transfer events of known plasmids in natural microbial communities. Basically, it consists in inoculating microcosms with a donor bacterium and to quantify the relative abundance of both the plasmid and the initial host DNAs over time. Since conjugative transfer is an intercellular mode of DNA replication, the plasmid to donor DNA ratio increases in the community DNA when the plasmid transfers into the indigenous population.

Conclusions
The transfer of the broad host range IncP-1β plasmid pB10 was studied in various environmental matrices. Under low level of inoculation, the transfer of pB10 in complex environments appeared relatively rare and was strongly matrix dependent. In some instance, the transfer of pB10 appeared influenced by eukaryotic predation, which alternatively could promote or inhibit plasmid transfer depending on the environmental matrix considered. An estimation of the IncP-1α/β plasmids abundance by qPCR demonstrated that pB10 transfer tends to be supported by environmental matrices exhibiting a higher content of IncP-1α/β plasmids. This tend to show that the relative abundance of IncP-1 plasmids in a given microbial community reflects its permissiveness to the transfer of plasmids belonging to the same incompatibility group, which seemed to prevail over transfer limitation due to the superinfection immunity (exclusion & incompatibility).
Antibiotic resistance and environment

METAL SPECIFIC MODULATION OF COMMUNITY PERMISSIVENESS TOWARDS BROAD HOST RANGE PLASMIDS THROUGH STRESS

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Background
The extent by which antibiotic resistance encoding conjugal plasmids transfer in microbial communities is of acute relevance in the age of massive antibiotic usage. The occurrence of stressors, like metals, might play a major role in altering the acute permissiveness of a bacterial community, since plasmid transfer is considered a main process in immediate stress response and adaptation to environmental changes.

Objectives
We evaluated: Does the introduction of metal stress alter a soil community’s permissiveness towards plasmids? Is a potential stress response general or a metal specific?

Methods
Hence, using a [3H]-leucine incorporation approach, we measured 20\% and 50\% inhibition concentrations for 5 metals. A \textit{mCherry}-tagged donor carrying the \textit{gfp}-tagged plasmid pKJK5 was mated with a soil bacterial community and exposed to metal stress. Transconjugants were quantified and isolated using advanced microscopy and fluorescent activated cell sorting. Sorted transconjugants were analyzed by pyrosequencing.

Conclusions
The imposed metal stress lowered the plasmid transfer frequency in filter mating assays. The intensity of this effect was metal specific and couldn’t be explained by measured growth inhibition of the recipient community because the exposure was normalized to similar inhibition. Results revealed an effect on transconjugant diversity for the heavy metals, Nickel or Copper, while Arsenic exposure caused no effect. The changes in transconjugal pool diversity couldn’t be directly correlated to a change in community diversity through the introduced stress. Our results suggest that this effect is not general but corresponds to a metal specific stress response of the soil microbial community with regard to plasmid transfer.
EXPLORATION OF THE REMOVAL POTENTIAL OF WASTEWATER ANTIBIOTIC RESISTANCE GENES BY SELECTED PHOTOCATALYTIC AND BIOLOGICAL TREATMENT TECHNOLOGIES

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Background
The pressure applied onto bacteria in aquatic environments by the extensive use of antibiotics, leads to increased incidence of antibiotic resistance (AR) (Volkmann et al., 2004), increasing the prevalence of nosocomial diseases. There is limited research regarding alternative actions of controlling AR in wastewater treatment plants (WWTPs), i.e. advanced oxidation processes (AOPs) and new biological methods which may remove AR more efficiently (Rizzo et al., 2013).

Objectives
The aspects examined were: i) the prevalence of selected AR genes (ARGs) at different steps of the WWTP process ii) the ARGs removal efficiency (vim, vanA, mecA and lak genes) of a pilot scale Membrane BioReactor (MBR) and of two AOPs (solar photo-Fenton treatment and heterogeneous (TiO₂) photocatalysis).

<table>
<thead>
<tr>
<th>Step of treatment</th>
<th>Detector</th>
<th>% change (/100 ng of total DNA)</th>
<th>Detector</th>
<th>% change (/100 ng of total DNA)</th>
<th>Detector</th>
<th>% change (/100 ng of total DNA)</th>
<th>Detector</th>
<th>% change (/100 ng of total DNA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Primary influent</td>
<td>vanA Sybr</td>
<td>-</td>
<td>lak1 Sybr</td>
<td>67</td>
<td>vim1 Sybr</td>
<td>69</td>
<td>mecA Sybr</td>
<td>-</td>
</tr>
<tr>
<td>MBR Mixed liquor</td>
<td>vanA</td>
<td>28</td>
<td>lak1</td>
<td>4400 (increase)</td>
<td>vim1</td>
<td>509 (increase)</td>
<td>mecA</td>
<td>18100 (increase)</td>
</tr>
<tr>
<td>Suspended solids</td>
<td></td>
<td></td>
<td></td>
<td>(increase)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>MBR effluent</td>
<td></td>
<td>7300 (increase)</td>
<td></td>
<td>6500 (increase)</td>
<td></td>
<td>99</td>
<td></td>
<td>-</td>
</tr>
<tr>
<td>Solar photo-</td>
<td></td>
<td>98</td>
<td></td>
<td>96</td>
<td></td>
<td>96</td>
<td></td>
<td>-</td>
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<tr>
<td>Fenton treatment</td>
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<tr>
<td>TiO₂ heterogeneous</td>
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<tr>
<td>photocatalysis</td>
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</tr>
</tbody>
</table>

Table 1. Removal of examined antibiotic resistance genes after the investigated AOPs and biological treatment.

Methods
The AOP experiments took place under bench-scale conditions while the MBR had a capacity of 10 m$^3$ day$^{-1}$. Real-time qPCR assessment followed the filtration and DNA extraction of the samples.

**Conclusions**
The highest prevalence of *lak* and *vanA* genes was found in solar photo-Fenton effluent. The highest *vim* prevalence was found in MBR effluent (Table 1), indicating an increase in the examined ARGs after treatment. Further optimization of the examined technologies must occur to ensure safe disposal of treated effluents into aquatic ecosystems.

This work was funded by COST through a Short Term Scientific Mission (STSM) within the COST scientific programme on 'Detecting evolutionary hotspots of antibiotic resistance in Europe (DARE)'. Nireas-IWRC (ΝΕΑ ΥΠΟΔΟΜΗ/ΣΤΡΑΤΗ/0308/09) is co-financed by the Republic of Cyprus and the European Regional Development Fund through the Cyprus Research Promotion Foundation.
Bacterial spores are ubiquitous in nature. They are stress resistant entities that can withstand high environmental temperatures, chemical insults and physical stress such as radiation or increased pressure. Spores are a concern to microbiological food stability due to these characteristics as upon survival of a preservation process they may start to germinate and grow out in food causing food spoilage. In addition germinating and outgrowing spores at undesired times and places pose a significant health burden. The challenge is amplified due to the heterogeneous germination and outgrowth behaviour of an isogenic spore population. We set out to analyse effects of thermal stress and the presence of a weak organic acid preservative on Bacillus subtilis spores of different maturation stages. Spore germination and outgrowth was assessed using live-imaging including the monitoring of intracellular pH with IpHluorin. Significant heterogeneity in spore germination and outgrowth was observed and monitored using the newly developed Sporetracker image analysis tool. Thermal stress clearly enhances heterogeneous germination behaviour. To analyse whether different levels of spore coat cross-linking may be involved in the (heterogeneous) stress response we set out to identify such cross-links by mass-spectrometry analysis of the insoluble coat fraction of B. subtilis spores of different maturation levels. Heat resistance of spores from all samples was also tested. Using our gel-free proteomic approach and LC-FTICR-MS/MS analysis we monitored the efficiency of tryptic digestion of proteins in the coat during spore maturation over a period of 10 days, using metabolically 15N labelled mature spores as a reference. The results showed that during spore maturation the loss of digestion efficiency of outer coat (for instance CotG, CotC, CotU) and crust (CotY, CotZ) proteins synchronized with an increased heat resistance.
Lactic acid bacteria (LAB) are a functionally related group of Gram-positive bacteria known essentially for their roles in food bioprocessing. The successful biotechnological application of LAB depends to a great extent on their unique phenotypic traits, which among others include fast acidification of the medium, texture and flavour forming abilities, bioprotection, and health promoting properties. However, the highly dynamic food market and ever-changing preferences from end consumers demands constant focus on product development. Fostered by this market push, Chr. Hansen places continuous efforts into the development of new LAB strains and cultures with novel properties and superior performance for the dairy industry. In the 21st century, genetic engineering could make such tasks more ease to accomplish. However, the tight requirements of regulatory agencies and the negative perception by consumers of genetically modified foods impose the exclusive use of natural strain improvement methodologies. Thus, at Chr. Hansen innovative product development includes combining classical strain improvement techniques such as random mutagenesis, dominant selection, adaptive laboratorial evolution, with the insights gained from the molecular understanding of our model systems. This approach will be illustrated with relevant example cases of strain improvement for the dairy industry.
A MOBILE GENETIC ELEMENT IS RESPONSIBLE FOR SUBSTANTIALLY INCREASED HEAT RESISTANCE OF BACILLUS SUBTILIS SPORES

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²Food Safety, NIZO Food Research B.V., Ede, Netherlands

Background
Spores of Bacillus subtilis are able to survive various harsh environmental conditions, including different heat preservation treatments used in food processing. Twenty strains of B. subtilis could be divided into two distinct groups with different spore heat resistances where the high spore heat resistance group required 100 times longer heating at 120°C to obtain the same level of spore inactivation.

Objectives
The goal of this study was to identify a genomic determinant for increased spore heat resistance, and to understand the underlying molecular mechanism mediating this increase in heat resistance of spores.

Methods
Whole genome sequencing was performed for these food isolates of B. subtilis, followed by correlating the genome content to the corresponding spore heat resistance phenotype. A transposon was identified and the insertion location was verified by PCR. Transfer of this element was achieved using filter mating.

Conclusions
A set of genes were uniquely present in the strains producing spores with increased heat resistance, and were located on transposon. The integration site of the transposon was found in the same gene for all strains producing high heat resistant spores. We demonstrated that the transposon was directly responsible for the very high heat resistance of spores; transfer of the transposon to a strain with significantly lower heat resistance (i.e. B. subtilis 168) rendered a phenotype of high spore heat resistance. The exact molecular mechanisms that mediate the increase of spore heat resistance are not known and the role of the genes on the transposon is currently subject of further investigation.
RECONSTRUCTION OF THE METHIONINE TO CYSTEINE CONVERSION PATHWAY IN LACTOBACILLUS PARACASEI FAM18149 BASED ON COMPARATIVE GENOMIC AND WHOLE TRANSCRIPTOME ANALYSIS

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Background

The flavor of cheese results from metabolic activities of bacteria like *Lactobacillus paracasei* during cheese ripening. Among many others, volatile sulfur compounds (VSCs) such as methanethiol, methional and hydrogen sulfide are key flavor compounds. Methionine is the main sulfur source in cheese, and is probably the precursor for bacterial VSC production. We observed that various strains of *L. paracasei* grew in medium that contained methionine as the sole sulfur source. Interestingly, these strains were also capable to produce VSCs.

Objectives

*L. paracasei* strains that grow with methionine as sole sulfur source have the potential to be used as flavor enhancing adjunct cultures in cheese making. A better understanding of the pathways involved in the metabolism of methionine by this species is of importance for proper strain selection.

Methods

*L. paracasei* FAM18149 was grown in medium with cysteine as only sulfur source and in medium with methionine as only sulfur source. The transcriptome of both conditions was measured using RNA-seq. The significantly regulated genes were studied in a gene trait analysis including genomes of 23 additional strains.

Conclusions

We found 55 significantly regulated genes. Among the regulated genes we identified a cluster that is only present in strains that grow with cysteine as only sulfur source. The cluster consists of a cysteine synthase, a cystathionine gamma-/beta-lyase and a serine acetyltransferase.
Based on these results we were able reconstruct the whole pathway of methionine to cysteine conversion and show that the cluster comprises the key enzymes of the pathway.
Background: The interest regarding the genetic variability among *L. casei* group members is increasing. These information are highly relevant in order to improve the knowledge and understand the correlation between genetic traits and biological properties, but also phenotypic characteristics and industrial properties of the strains. Objectives: The study of the correlation between the genetic profile of the strains and the stress response.

Methods: 25 *L. paracasei* strains were selected for MLST analysis; 8 genes related to stress response (ctsR, hrcA, cydD, cydA, nox, npr, pox, dnaK) were chosen as target genes. After the comparison of the obtained sequences, a Sequence Type (ST) number was assigned to each strain. MEGA version 6.06 software (http://www.megasoftware.net) was used to compare and align the sequences and phylogenetic trees were constructed using the UPMGA. DnaSP software version 5.10.1 was used to perform the descriptive analysis.

Conclusions: After PCR of the internal sequences of the selected loci (npr, nox, hrcA, ctsR, cydA, cydD, dnaK, and pox), the amplicon was obtained for all the tested strains, confirming the presence of the target genes. All the studied strains resulted having a different Sequence Type (ST). The mean G + C content of the different gene fragments was higher than 0.45, meaning that the analysed genes are almost stable. Observing the MLST UPGMA trees, the strain’s cluster confirmed a correlation between the different genetic profile of the stress related genes and the different stress resistance of the strains.
LISTERIA MONOCYTOGENES SURVIVAL STRATEGIES: POPULATION HETEROGENEITY AND STRESS RESISTANT VARIANTS

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\textsuperscript{1}Laboratory of Food Microbiology, Wageningen University, Wageningen, Netherlands

Background

The food-borne pathogen \textit{Listeria monocytogenes} is a Gram-positive microaerophilic facultative anaerobic rod and the causative agent of listeriosis. This pathogen is notorious for its robustness and a range of studies have been performed that supplied insight in adaptive stress response and survival strategies.

Objectives

The objectives of this study are the assessment of \textit{L. monocytogenes} population heterogeneity and the genotypic and phenotypic analysis of stress resistant variants obtained after high hydrostatic pressure, heat and acid exposure.

Methods

A range of phenotypes was determined for the different stress resistant variants obtained using \textit{L. monocytogenes} EGDe, LO28 and ScottA, including level of acid resistance, growth rates at different temperatures, and biofilm-forming capacity. Whole genome sequencing analysis of \textit{L. monocytogenes} wild types and selected stress resistant variants was performed, followed by cluster analysis of genotypic and phenotypic parameters to assess population diversity within \textit{L. monocytogenes} strains.

Conclusions

Significant differences in phenotypes among variants were observed that pointed to differences in robustness and growth performance under food-relevant conditions. Whole genome sequencing analysis of \textit{L. monocytogenes} wt and stress resistant variants revealed mutations in \textit{ctsR}, encoding a class III heat shock repressor, and in \textit{rpsU}, encoding ribosomal protein S21. Cluster analysis of genotypic and phenotypic parameters revealed large population diversity even within one \textit{L. monocytogenes} strain and that different adverse conditions select for different variants. The large population diversity of \textit{L. monocytogenes} signifies the organism's genetic flexibility, which in turn may contribute to the survival and persistence of this human pathogen in food-processing environments.
Background

Many lactic acid bacteria strains show extraordinary metabolic abilities, which are interesting for their application in functional food development such as group B vitamin overproduction. In this context, the determination of quality and food safety related traits i.e. antibiotic resistance are required.

Objectives

In this study, 29 Lactobacillus plantarum strains isolated from chicha, a traditional maize-based fermented Andean beverage, were previously selected by their ability to produce folic acid. Thus, we focused on evaluating additional activities relevant to functional food industry and to food safety and quality.

Methods

Lactobacillus plantarum strains were screened for riboflavin production (by reverse phase HPLC with fluorescence detector) and for the presence of riboflavin production related genes (ribA, ribB, ribC, ribG and ribH), antibacterial activity (Gaudana et al. 2010. Brit J Nutr. 103: 1620–1628) and antifungal activity (Magnusson et al. 2003. FEMS Microbiol Lett. 219:129-135). Regarding food safety, antibiotics resistance was tested as recommended by EFSA (2012) for Lb. plantarum.

Conclusions

Most strains strongly inhibited the growth of Listeria innocua CECT 910, Escherichia coli CECT 5947, Salmonella enterica CECT 4138 and showed antifungal activity against Aspergillus oryzae CECT 2094. Some of them, inhibited Aspergillus niger CECT 2807, as well. Among them, four strains exhibited traits of interest for the functional food industry. They produced 400ng/ml riboflavin, determined by HPLC, and showed the amplification products corresponding to genes in the riboflavin synthetic pathway. All of these were sensitive to the antibiotics tested and were able to inhibit the growth of relevant food pathogens and spoiler fungi.
FEMS-1244
Molecular principles of biofilm formation and dispersion

Regulation of functional interferences between *Escherichia coli* type 1 fimbriae and Ag43 phase-variable adhesins

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\(^1\)Department of Microbiology, Institut Pasteur, Paris, France

Bacteria use a large number of adhesins involved in bacteria/surfaces and bacteria/bacteria interactions to colonize their environments. In *Escherichia coli*, there are two essential and prototypic adhesins: the chaperone-usher type 1 fimbriae, involved in mannose-dependent interactions to host cells, and the auto-transporter adhesin Ag43, mediating bacterial auto-aggregation. These two adhesins are regulated by two distinct phase variation mechanisms: a promoter recombination inversion for type 1 fimbriae and an epigenetic mechanism involving the Dam methylase and the oxidative stress regulator OxyR for the auto-transporter adhesin Ag43. Since expression of long type 1 fimbriae physically interferes with short Ag43 adhesins, the functions of these two adhesins are antagonistic, which raises the question of how *E. coli* coordinates their expression. I will describe previously unsuspected regulatory connection between phase variation mechanisms and stress responses and discuss how it contributes to coordinate the expression and exposition of *E. coli* surface adhesins and bacterial colonization capacity.
The process of biofilm dispersion represents an important phenotypic switch that allows cells residing within a biofilm to respond to changing conditions outside and within biofilm structures and to evade environmental stresses. Dispersion occurs in response to a wide array of signals, with dispersion-inducing conditions ranging from environmental cues to self-synthesized signaling molecules. Previous findings indicate biofilm dispersion by *Pseudomonas aeruginosa* to occur in response to various cues, to require BdlA and phosphodiesterases (PDEs, for instance DipA and RbdA), and to coincide with increased PDE activity and a reduction of c-di-GMP levels. However, little is known about dispersion cue sensing, the signaling events translating these cues into the modulation c-di-GMP levels, and the subsequent signaling cascade to enable dispersion resulting in the phenotypic switch. The presentation will focus on recent findings indicating how dispersion cue perception is translated across cellular compartments into the modulation of the intracellular c-di-GMP pool, the molecular events associated with signal transduction, and subsequent events associated with the induction of dispersion.
Background

The human pathogen *Bacillus cereus* is responsible for many recurrent outbreaks of food poisoning. Spores and biofilms are considered the most important reservoirs of *B. cereus* in contaminated fresh vegetables and fruits. Bacterial biofilms are difficult to eradicate specially due to the presence of a protective extracellular matrix made of exopolysaccharides, proteins, and other components. Amyloid-like proteins are essential for the integrity of biofilms of the related bacteria species *Bacillus subtilis*.

Objectives

To investigate the presence of amyloid-like fibers in biofilms of *B. cereus*.

Methods

We identified two genomic loci in *B. cereus*, which encode two orthologues of the amyloid-like protein TasA and a SipW signal peptidase of *B. subtilis*. Mutagenesis in *B. cereus* or heterologous expression of alleles in *B. subtilis* mutants combined with crystal violet staining served to evaluate the formation of biofilm. Electron microscopy let us visualize the presence of fibers on cells.

Conclusions

We demonstrate that the proteins TasA and CalY are necessary for *B. cereus* biofilm formation: i) as pellicle on the air-liquid interphase or ii) adhesion to abiotic surfaces. TasA and to a lesser extent CalY polymerizes in the form of fibers in the cell surface. Our findings of heterologous expression in *B. subtilis* let us propose an amyloid-like
nature of the \textit{B. cereus} TasA-based fibers.

\textbf{\textit{B. subtilis} \text{\textit{\textless}operon}}

\[ + \textit{sipW-to-calY} \quad + \textit{sipW-tasA} \quad + \textit{sipW-calY} \]

\textit{Heterologous expression}
Background
Approximately 80% of Cystic Fibrosis (CF) patients are infected with \textit{P. aeruginosa}. \textit{P. aeruginosa} infected lungs/sputum of CF patients contain pyocyanin up to 27.3 µg/ml. Pyocyanin is a potent virulence factor causing cell death in chronic lung infection in CF patients, thus increasing mortality. Studies suggest that pyocyanin influence \textit{P. aeruginosa} biofilm formation. However, no work was done investigating the mechanism or functions of pyocyanin in biofilm formation.

Objectives
To elucidate the mechanism and functions of pyocyanin in \textit{P. aeruginosa} biofilm formation and identify a novel therapeutic target to modulate pyocyanin functions thereby control \textit{P. aeruginosa} biofilm formation.

Methods
eDNA concentration in \textit{P. aeruginosa} culture was measured using Qubit 2.0 fluorometer. Pyocyanin-eDNA binding mechanisms were analysed using spectrometer techniques. Influence of pyocyanin-eDNA interaction on viscosity of DNA solution and cell surface of \textit{P. aeruginosa} strains were analysed using viscometer and surface thermodynamics approach respectively. Biofilm imaging and analysis of biofilm properties were performed using confocal microscopy. Antioxidants interaction with pyocyanin and inhibition of pyocyanin-eDNA binding were investigated using NMR and spectrometer techniques respectively.

Conclusions
Pyocyanin promotes eDNA release in \textit{P. aeruginosa} wild-type by inducing cell lysis mediated through generation of \textit{H}_{2}\textit{O}_{2}. Pyocyanin intercalates into the nitrogenous bases of DNA and consequently increases DNA viscosity and influences \textit{P. aeruginosa} cell surface hydrophobicity, physico-chemical interactions promoting bacterial interaction and biofilm formation. Pyocyanin deficient strain showed significant reduction in eDNA release and biofilm formation. Ascorbic acid and glutathione modulate pyocyanin functions thus inhibit pyocyanin-eDNA binding and consequently hamper biofilm development.
BISTABLE GENE EXPRESSION IN SALMONELLA CONNECTS VIRULENCE TO PERSISTENCE
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Background
For the human and animal pathogen Salmonella Typhimurium, biofilm formation is correlated with persistence outside a host but the connection to virulence is unknown.

Objectives
To analyze specialized cells of S. Typhimurium that arise from bistable expression of CsgD, the central biofilm regulator, during growth under biofilm-inducing conditions.

Methods
RNA-seq analysis of multicellular aggregates and planktonic cells identified 1856 genes, representing 34% of the genome, that were differentially expressed. Aggregated cells displayed the characteristic gene expression of biofilms, whereas
planktonic cells had enhanced expression of virulence genes, which were previously thought to be activated only by host conditions. Increased synthesis of the *Salmonella* pathogenicity island-1 type three secretion system in planktonic cells correlated with enhanced invasion of a human intestinal cell line and significantly increased virulence in mice as compared to the aggregates. However, when these same groups of cells were exposed to desiccation, the aggregates survived better and the competitive advantage of planktonic cells was lost.
Conclusions

We hypothesize that CsgD-based differentiation is a form of bet-hedging for *S. Typhimurium*, with single cells primed for host cell invasion and aggregated cells adapted for persistence in the environment. This could be a common strategy for pathogens to spread the risks of transmission and ensure a smooth transition between the host and the environment.
FEMS-2654
Molecular principles of biofilm formation and dispersion

GIL, A NEW C-DI-GMP-BINDING PROTEIN DOMAIN INVOLVED IN REGULATION OF CELLULOSE SYNTHESIS IN ENTEROBACTERIA
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Background
Numerous enzymes are involved in c-di-GMP synthesis and degradation in enterobacteria. In contrast, only a handful of c-di-GMP receptors/effectors have been identified.

Objectives
The aim of this study was to identify new c-di-GMP receptors and to further characterize them.

Methods
In search of new c-di-GMP receptors, we screened the Escherichia coli ASKA overexpression gene library using the Differential Radial Capillary Action of Ligand Assay (DRAcALA) with fluorescently and radioisotope-labelled c-di-GMP. We uncovered three new candidate c-di-GMP receptors in E. coli and characterized one of them, BcsE.

Conclusions
The bcsE gene is encoded in cellulose synthase operons in representatives of Gammaproteobacteria and Betaproteobacteria. The purified BcsE proteins from E. coli, Salmonella enterica and Klebsiella pneumoniae bind c-di-GMP via the domain of unknown function, DUF2819, which is hereby designated GIL, GGDEF I-site like domain. The RxGD motif of the GIL domain is required for c-di-GMP binding, similar to the c-di-GMP-binding I-site of the diguanylate cyclase GGDEF domain. Thus, GIL is the second protein domain, after PilZ, dedicated to c-di-GMP-binding.
We show that in *S. enterica*, BcsE is not essential for cellulose synthesis but is required for maximal cellulose production, and that c-di-GMP binding is critical for BcsE function. It appears that cellulose production in enterobacteria is controlled by a two-tiered c-di-GMP-dependent system involving BcsE and the PilZ domain containing glycosyltransferase BcsA.

Host manipulation and bacterial survival Offered by Pathogens and Disease

Salmonella's intracellular toolkit

D. Holden

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Following entry of Salmonella into host cells, this pathogen resides in a membrane-bound compartment called the Salmonella-containing vacuole (SCV). Here, the bacteria sense the acidic pH and poor nutritional status of the vacuole lumen and either enter a viable non-replicating state or begin to divide. Replication is dependent on activation of the SPI-2 type III secretion system (T3SS). The T3SS comprises an envelope-spanning secretion system and associated translocon pore in the vacuolar membrane. Bacteria then sense the near-neutral pH of the host cell cytoplasm; this results in dissociation and degradation of a bacterial membrane-bound regulatory complex, which activates translocation of approximately 30 bacterial effector proteins into the host cell.

We are currently studying the biochemical and physiological functions of various effectors, which have been implicated in several activities, including lysosomal detoxification, SCV localisation and membrane dynamics, interference with immune signalling and the induction of cytotoxicity. In this talk I will discuss our recent progress in these areas.
Host manipulation and bacterial survival Offered by Pathogens and Disease

Manipulation of host membrane transport by Type IV effector proteins

**C. Roy**

*Microbial Pathogenesis, Yale University School of Medicine, New Haven, USA*

Effector proteins delivered into the cytosol of host cells by Dot/Icm type IV secretion systems modulate processes important for creating a vacuole that supports bacterial intracellular replication. Here we describe the biochemical function of *Legionella pneumophila* and *Coxiella burnetii* effector proteins that play specific roles in controlling host membrane transport. These studies reveal new enzymatic activities and protein structures that demonstrate these pathogens encode novel effector proteins that can manipulate evolutionarily conserved host proteins that control membrane transport processes, which provide insight into how bacterial pathogens are able to construct a unique vacuole inside phagocytic host cells.
IDENTIFICATION OF LEGIONELLA PNEUMOPHILA VIRULENCE FACTORS REQUIRED TO SUBVERT HOST AUTOPHAGY

V. Lelogeais, M. Faure, F. Vavre, P. Doublet

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2 Autophagy Infections Immunity, International Center for Infectiology Research INSERM U1111 CNRS UMR 5308 EN S Lyon UCBL, Lyon, France
3 Evolutionary genetic and genomic, Biometry and Evolutionary Biology UMR CNRS 5558, Villeurbanne, France

Background

Legionella pneumophila is the causative agent of legionnaire’s disease, a public health problem leading to at least 10% mortality rate. Via effectors secreted by its type IV secretion system, this intravacuolar bacterium interacts with many intracellular pathways of host cells including autophagy. This highly conserved pathway allows eukaryotic cells to recycle end-life cytosolic components in order to regulate cell homeostasis. Autophagy is also a degradative pathway essential to fight intracellular pathogen infections, but numerous microorganisms have evolved strategies in order to subvert this mechanism. The interaction between L. pneumophila and autophagy has been reported but remains still unclear.

Objectives

Our objective is to better decipher the L. pneumophila/autophagy interplay to determine whether this pathogen is controlled by autophagy or whether it gets benefit from this cellular process to survive/proliferate.

Methods

We used HeLa cells as a model to follow autophagy modulation upon L. pneumophila infection. Cells were infected with the northern Europe endemic strain L. pneumophila Paris.

Conclusions

Our experiments show that L. pneumophila infection induces autophagy in a type IV secretion system dependent manner, early post infection within infected cells but also in neighboring non-infected cells. Otherwise, preliminary data suggest that inhibition of autophagy is linked to a decreased bacterial replication, therefore autophagy could be benificial for L. pneumophila. Thus, the autophagy stimulation could allow the acquirement of nutrients by the bacterium, provide membrane source for the expanding of L. pneumophila vacuole and/or lead to the delay of the fusion between L. pneumophila vacuole and lysosomes.
Host manipulation and bacterial survival Offered by Pathogens and Disease

PASTEURIELLA MULTOCIDA TOXIN MANIPULATES T CELL DIFFERENTIATION

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Background

_Pasteurella multocida_ are gram-negative bacilli, which cause respiratory diseases in a number of wild and domestic animals. Toxigenic serotype A and D strains produce the _Pasteurella multocida_ toxin (PMT), a classical AB toxin containing a deamidase activity in its catalytic domain. The toxin constitutively activates heterotrimeric G proteins and induces downstream signalling cascades involved in cytoskeleton rearrangement, proliferation, differentiation or survival of the host cell. Pathologically, PMT causes porcine atrophic rhinitis characterized by an increased number of osteoclasts and bone resorption at the nasal turbinate bones. However, PMT does not only act on cells of the bone system but we could demonstrate that it also modulates the signalling of many other hematopoietic cells such as B cells, macrophages or monocytes.

Objectives

We investigated how PMT treatment affects the differentiation of primary human T lymphocytes.

Methods

Proliferation studies showed that the toxin amplifies CD3/28 activated proliferation of T cells through the induction of cell cycle progression. Characterisation of transcription factor activation and the release of cytokines showed that PMT manipulates lineage determination towards T_{H17} and T_{reg} cells as PMT-treated T cells are double positive for the lineage specific transcription factors FOXp3 (T_{reg}) and RORγT (T_{H17}) but predominantly behave like inflammatory IL-17-releasing T_{H17} cells.

Conclusions

These results indicate that a bacterial toxin is able to manipulate the direction of T_{H17} cell differentiation and thereby presumably adjusts the sensitive balance of the immune response towards a condition that benefits the pathogen.
**Host manipulation and bacterial survival Offered by Pathogens and Disease**

**R-SPONDIN-2 MEDIATES SUSCEPTIBILITY TO INTESTINAL INFECTION WITH CITROBACTER RODENTIUM**  
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2Human Genetics, McGill University, Montreal, Canada  
3Biotechnology Research Institute, National Research Council of Canada, Montreal, Canada  
4Cytochem, Cytochem Inc, Montreal, Canada

**Background**  
*Citrobacter rodentium* is a mouse pathogen widely used as a model for Enteropathogenic and Enterohemorrhagic *Escherichia coli* infections in humans. While *C. rodentium* causes self-limiting colitis in most inbred mouse strains, it induces fatal diarrhea in hyper-susceptible strains.

**Objectives**  
Here we sought to identify the gene and molecular mechanisms underlying this differential response to intestinal infection.

**Methods**  
We used a forward genetics approach to unambiguously localize the locus conferring this differential outcome to a 4 Mb interval. Systematic evaluation of the genes within this region pinpointed *R-spondin* -2 as the gene underlying infection susceptibility.

**Conclusions**  
Robust induction of *R-spondin* -2 expression during infection in susceptible mouse strains leads to pathological activation of WNT signaling, subsequent loss of intestinal differentiation, and animal death. Conversely, mouse strains that do not induce *R-spondin* -2 expression following infection undergo milder, self-limiting disease with no mortality. Our data demonstrate a previously unknown role of R-spondins and WNT signaling in susceptibility to infectious diarrhea and identify *R-spondin* -2 as a key molecular link between enteric infection and control of intestinal homoeostasis.
Background

Bacterial vaginosis (BV) is a common condition of the human vagina in which the lactobacilli that normally dominate this niche are absent and replaced by a plethora of fastidious anaerobes. BV is associated with an increased risk of acquiring sexually transmitted disease and adverse health outcomes during pregnancy. Factors that drive the microbial dynamics in BV are poorly defined.

Glycogen is deposited in high levels in human vaginal epithelial cells. It has been postulated to be a major carbon source for vaginal bacteria.

Objectives

Here we test the hypothesis that BV-associated bacteria digest and metabolize glycogen.

Methods

We evaluated glycogen degrading enzyme activity in vaginal specimens of women with and without BV. Growth assays were conducted to examine the ability of lactobacilli and BV-associated bacteria to utilize glycogen. We examined evidence for glycogen-degrading enzymes among vaginal bacteria.

Conclusions

These experiments provide strong evidence that BV associated bacteria degrade glycogen and metabolize the released sugar residues, whereas most vaginal lactobacilli do not utilize glycogen. These results provide an explanation of the recent observation by Mirmonsef et al. (1) that glycogen levels in BV are reduced.

Transcriptional networks controlling pathogenicity and polarized growth in U. maydis

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The transition from yeast-like saprophytic growth to the formation of pathogenic filamentous hyphae is a critical step in the life cycle of the basidiomycete Ustilago maydis. The switch is controlled by a pheromone/receptor system and the heterodimeric bE/bW transcription factor complex, encoded by the a and b-mating type loci, respectively. As both pathways control polar growth and cell cycle, they have to be closely interconnected and cross-controlled. The transcription factor Rbf1, a central node for gene regulation during pathogenic development, responds to inputs from both a and b pathways. Coordination is achieved via the interaction of Rbf1 and bW with the Clp1 protein that renders the bE/bW complex inactive and converts Rbf1 to a repressor of the pheromone pathway, releasing both the a and bE/bW mediated cell cycle arrest. We are currently investigating the molecular mechanisms by which the factors independently, cooperatively, and after interaction with Clp1, regulate gene expression during the dimorphic switch, using a combination of reporter gene assays, in vitro DNA binding studies and the genome-wide identification and occupation of promoter binding sites.

Δclp1 strains are capable to infect maize plants, but, as the cell cycle block is not released, development is stalled after formation of the appressorium, before the first cell division. In both wildtype and Δclp1 hyphae, nuclear envelopes start to dissolve before appressorium formation, which is typical for nuclei entering mitosis. In wildtype cells, nuclei migrate into the invading hyphae and divide. However, nuclei of Δclp1 cells remain in the appressorium, suggesting a novel function for Clp1 in the coordination of nuclear division and nuclear migration.
Fungi cause severe crop losses and threaten food security worldwide. The soil-borne fungal pathogen *Verticillium dahliae* causes vascular wilt disease on hundreds of plant species, and disease control is challenging because resistance in plants is relatively rare. Moreover, *V. dahliae* has a flexible genome allowing it to escape host immunity and maintain aggressiveness. So far, knowledge on mechanisms governing this genomic flexibility remains limited.

Through comparative population genomics we have started to unravel mechanisms to establish the genomic diversity that is essential for adaptive genome co-evolution during the continued arms race with host plants. To this end, two *V. dahliae* genomes were assembled from telomere-to-telomere using long-read sequencing technology and optical mapping, and compared these to the genomes of other *Verticillium* spp., revealing a pre-speciation genome duplication event. Comparative genomics using the two finished *V. dahliae* genomes furthermore revealed recent segmental duplications that established lineage-specific regions. Interestingly, these regions are enriched for in planta-expressed effector genes encoding secreted proteins that enable host colonization, and thus contribute to the evolution of virulence. Our evidence suggests that error-prone homology-dependent DNA repair has caused genomic rearrangements, leading to extensive structural variations. Re-sequencing of additional strains showed that independent losses of genetic material favored the escape of host recognition and, likely, host specificity. We propose that evolution of *V. dahliae* is linked to segmental genome duplications mediated by improperly repaired DNA breaks.

In addition to genome evolution, we also study the role of epigenetic modifications on virulence of *V. dahliae* and the biological functions of effector proteins. Collectively, these research lines provide insight in mechanisms that make this fungus such a successful broad host range pathogen.
Fungal pathogens

FGSSP6, A CERATO-PLATANIN PROTEIN WHICH CONTRIBUTES TO FUSARIUM GRAMINEARUM VIRULENCE ON WHEAT SPIKES.

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Background

*Fusarium graminearum* (*Fg*) is one of the main fungal pathogens of wheat crops globally, causing Fusarium ear blight (FEB) disease. Like many other plant pathogens, *Fg* is predicted to produce *in planta* various secreted effector proteins that modulate plant metabolism to suppress or re-programme plant defences. These induced changes enable successful colonisation of plant tissues.

Objectives

To identify *Fg* effector proteins that can suppress host plant defences

Methods

We selected a set of small secreted proteins (SSP) to express *in planta* using the *Barley stripe mosaic virus* over-expression system (BSMV-VOX) and tested whether any enhance *Fg* fungal infection of susceptible wheat spikes.

Conclusions

Amongst the initial set of *Fg* SSP tested, one called FgSSP6, which belongs to the cerato-platanin protein (CPP) family, caused the quicker development of FEB disease symptoms in wheat ears following point inoculation. Bleaching of all spikelets on wheat spikes occurred by 12 days post inoculation (dpi) in the presence of BSMV:FgSSP6, whereas with various BSMV control vectors, full *Fg* infection took 16 dpi or longer. In several other plant pathogenic fungi, CPPs are already known to act as key virulence factors. So far our results suggest that FgSSP6 does not induce necrosis to assist infection, but either suppresses plant defence responses or plays a novel virulence role. This is the first report of the successful use of the BSMV-VOX system in combination with the *Fg*-wheat spike infections to identify novel
candidate \textit{Fg} effectors. Further studies are planned to clarify the precise \textit{in planta} role(s) of this newly identified cerato-platanin effector.
A NOVEL METHOD FOR RAPIDLY ISOLATING FUNGAL SUPPRESSIVE MICROBES DIRECTLY FROM SOIL

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Background
Soilborne fungal root phytopathogens are some of the most damaging and difficult to control pathogens faced by agricultural production in the field, nurseries and greenhouses worldwide. There are many detrimental issues with current chemical control methods and this has increased interest in biological control using soil microorganisms that suppress the growth of phytopathogens. Antibiosis is one of the most important mechanisms responsible for fungal antagonism, with some significant antifungal compounds including, antibiotics, volatile organic compounds, hydrogen cyanide and lytic enzymes (Bhattacharyya & Jha, 2012; Compant et al., 2005). Up to date methods for the isolation of fungal-suppressive microorganisms from the soil are time consuming and tedious.

Objectives
The aim of this study was to establish a simple method for isolating fungal pathogen-suppressive microbes (bacteria and fungi) directly from the soil as well as procedures for confirmation of disease suppression.

Methods
We will report on these methods, which were so far tested with three cotton fungal pathogens Thielaviopsis basicola, Verticillium dahliae and Fusarium oxysporum and a pathogen of button mushrooms Verticillium fungicola.

Conclusions
We have isolated a diversity of T. basicola- suppressive fungi and bacteria from two vastly different soil types. Identification of the antagonistic isolates revealed that they are a diverse lot, some belonging to groups known to be suppressive of a wide range of fungal pathogens, endorsing the power of this technique to rapidly and directly isolate soil-borne microbes antagonistic to a wide variety of fungal pathogens.
EVALUATION OF GENETIC DIVERSITY AMONG PHYTOPATHOGENIC ISOLATES OF FUSARIUM SOLANI COMPLEX CAUSING SHISHAM DIEBACK DISEASE IN PAKISTAN

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Background

*Dalbergia sissoo* Roxburgi, (shisham) is an important and established forest species in south Asia. However, this precious tree is facing many devastating challenges from last six decades that dragging it towards disappearance from the subcontinent. Dieback is one of main diseases in Shisham plantations and caused by *Fusarium solani*.

Objectives

The aim of this study was to investigate the regional dependant genetic variation among different pathogenic *F. solani* isolates and a link to persistence cause of shisham dieback in various agro ecological zones.

Methods

A total of 23 pathogenic representative isolates of *F. solani*, collected from various agro ecological zones were subjected to analysis of genetic variability in terms of DNA polymorphism using RAPD-PCR. Results were obtained with the help of 10 randomly amplified polymorphic DNA markers (OPA1-OPA10) to test genetic variability in *F. solani* isolates.

Conclusions

A total of 238 amplified products generated with the primers and 23 isolates of *F. solani*. The pattern of genetic variability in the isolate was also supported by the analysis of the similarity indices and UPGMA dendrogram. Twenty three isolates of *F. solani* recovered from different areas showed a significant genetic variation during RAPD analysis. It seems that the genetic variability among fungal isolates is regional dependent and reason of persistent cause of shisham dieback in Punjab, Pakistan. Current findings showed the variation among pathogenic isolates of *F. solani* and
revealed that persistent cause of shisham dieback is due to *F. solani* complex in different shisham growing regions in province Punjab, Pakistan.
APPLICATION OF TRICHODERMA STRAINS AGAINST GTD PATHOGENS

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Background

The Grapevine Trunk Diseases are one of the most prevalent diseases of the vineyard worldwide, caused by a variety of fungal pathogens. Mycoparasites, like Trichoderma species are potential biopesticides against GTD pathogens.

Objectives

The aims of this study were taxonomical identification of ten Trichoderma sps. from asymptomatic plants in the Tokaj wine region the determination of mycelial growth rate at different temperatures, the mycoparasitic potential against Diplodia seriata and Neofusicoccum parvum and field experiments with the mixed Trichoderma spore suspension.

Methods

The Trichoderma isolates were identified based on their ITS1,2 and tef1 marker sequences. Their mycelial growth was determined from the two average colony diameters on PDA, measured for each Trichoderma isolates for 4 days. The mycoparasitic ability was detected according the method of Szerekes et al (2006; Journal of Microbiological Methods, 619-622). Some GTD symptomatic plants were treated with Trichoderma spore suspension in 2014. The treated trunks were checked monthly.

Conclusions

The ten isolates belonged to three species: T. harzianum (TR01-05, TR07, TR09-10), T. orientalis (TR06) and T. viride (TR08). The Trichoderma isolates showed different mycelial growth rate on different temperatures (5 - 37°C). The T. orientalis (TR06) and T. harzianum (TR05) isolates showed the highest growth rates within the whole temperature range, and their growth rates were especially high at 30 and 37°C. All Trichoderma overgrew the GTD pathogens, and sporulated on their colonies, therefore their Biocontrol Index was 100%.
The symptoms decreased on all treated samples, except one. Moreover the *Trichoderma* strains could be reisolated from their woody tissues.
DNA substrates for horizontal gene transfer by natural transformation

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The dynamic composition of the genomes of free-living bacteria challenges our understanding of microbial species. Horizontal gene transfer contributes to this dynamics and enables the ongoing dissemination of antimicrobial resistance genes across bacterial populations. Thus, there is a immediate need for a better understanding microbial genome dynamics both on a short-term ecological and long-term evolutionary scale. Natural transformation is a gene transfer mechanism that can both lead to intra-genomic rearrangement and inter-genomic DNA recombination. Here I present some recently experimentally derived results that advance our understanding of how species-foreign DNA fragments can be acquired by the naturally transformable model bacterium Acinetobacter baylyi, and describe the fate of such DNA after initial recombination with the host genome. Extracellular DNA in decaying organisms becomes chemically modified and degraded over time. However, it is known that from paleogenetics that short DNA fragments remain over time in some environments. The newly discovered opportunity for bacteria to acquire very short DNA fragments suggest extinct genomes can be an additional source of DNA for naturally transformable organisms.
Evolution of a superbug: Real-time transfer of antimicrobial resistance in the host

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*Staphylococcus aureus* is a commensal and major pathogen of humans and animals, capable of carrying a wide range of mobile genetic elements (MGEs) encoding antimicrobial resistance (AMR), virulence and host adaptation pathways. Whole genome analysis reveals a population structure consisting of independently evolving lineages, each defined by variation in core genomes. Within lineages, MGEs vary substantially with potential implications for selection in various hosts and under antimicrobial pressures. Little is known about the process of MGE transfer in vivo, nor how *S. aureus* adapt to antimicrobials or new hosts in real time. We co-colonized gnotobiotic piglets with both human- and pig-associated variants of the lineage clonal complex (CC)398, and observed transfer of MGEs including resistance genes from the pig to the human isolate within 4 hours. Extensive and repeated transfer resulted in colonization with isolates carrying a wide variety of MGE combinations. Whole genome sequencing of progeny bacteria revealed no acquisition of core genome polymorphisms. Separate studies of human hospitalized patients colonized with methicillin-resistant *S. aureus* (MRSA) CC22 revealed clonal populations varying in antimicrobial resistance profiles and MGE carriage, as well as free bacteriophage capable of gene transfer by generalized transduction. Our data show that transfer of AMR can occur at very high frequency *in vivo* and at higher frequency than detected *in vitro*. 
PLASTICITY OF EVOLVABILITY DRIVES EVOLUTION TO HIGH ETHANOL TOLERANCE IN ESCHERICHIA COLI

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Background
Ethanol tolerance is a complex trait requiring interaction between different genes and pathways. Exposure to ethanol is toxic and slows down the cellular growth rate. Consequently, evolving high ethanol tolerance is slow and depends on the simultaneous occurrence of different mutations. This is unlikely in wild-type cells with a low mutation rate. In contrast, the odds that different mutations occur simultaneously are higher in hypermutable strains. These mutators are known to have an increased evolvability on a short evolutionary timescale, but display decreased fitness as the genetic load progressively becomes more deleterious.

Objectives
We aim at understanding the evolutionary mechanisms underlying adaptation to ethanol. This can lead to new insights that are also relevant for other complex phenotypes. Additionally, we investigate the role of a mutator phenotype in evolution towards high ethanol tolerance. This can shed light on the way bacteria adapt to severe stress conditions.

Methods
Here we used experimental evolution to study mechanisms underlying evolution to high ethanol tolerance in E. coli. Furthermore, through fluctuation assays and competition assays we determined mutation rates and established the advantage or disadvantage of hypermutation at different times during adaptive evolution.

Conclusions
We show that an increased mutation rate is essential for E. coli to adapt to high ethanol levels under slow-growing conditions. In direct competition under ethanol stress hypermutators outcompete wildtype cells. Moreover, we demonstrate that the population mutation rate decreases again when high ethanol tolerance is reached, indicating that once a population reaches a fitness peak, it strives to preserve the genome.
FEMS-1739
Genomics, evolution, phylogeny

REPLICATION-DEPENDENT GENE DOSAGE OF RIBOSOMAL PROTEINS REGULATES GROWTH RATE OF VIBRIO CHOLERAE.
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Background

The effects of gene order within the bacterial chromosome on cell physiology are poorly understood. In silico approaches have showed that ribosomal protein genes locate near the replication origin (oriC) in fast-growing bacteria, suggesting that such a positional bias is an evolutionarily conserved growth-optimization strategy. Vibrio cholerae, a bi-chromosomal fast-growing pathogen is a well-characterized model organism to experimentally tackle this hypothesis. The S10-spc-\(\alpha\) locus (S10) encodes half of the ribosomal protein genes and is conserved along all life forms.

Objectives

To relocate S10 to different positions along the genome and measure putative physiological effects.

Methods

We used novel recombineering tools to create a set of isogenic strains in which S10 was systematically relocated to alternative genomic positions. Their DNA and RNA was analyzed using NGS.

Conclusions

We show that S10 relative distance to the oriC tightly correlates with a reduction of its dosage, mRNA abundance and growth rate. This is accompanied by a significant reduction in host-invasion capacity of Drosophila melanogaster. Strains bearing two S10 copies far from oriC rescue both phenotypes demonstrating that replication-dependent gene-dosage reduction is the main mechanism behind these alterations. Deep sequencing analyses shows that i) the whole oriC region is less transcribed in the most affected derivatives and ii) Gene Set Enrichment Analyses reveal ‘translation’ as the most altered pathway in such derivatives. S10 positioning connects genome structure to cell physiology in V. cholerae. We show that genomic positioning of genes involved in the flux of genetic information conditions global growth and hence bacterial physiology and evolution.
EVOLUTION AND PATHOADAPTATION OF PSEUDOMONAS AERUGINOSA IN CYSTIC FIBROSIS PATIENTS

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Background
Laboratory evolution experiments have led to important findings addressing to what extent natural selection constrains parallel evolving organisms to follow common evolutionary pathways towards adaptive phenotypes, and in which way evolutionary outcomes become intertwined over time such that future alternatives may be contingent on the prior history of an evolving population. Nonetheless, similar systematic investigations of bacterial populations living in complex natural environments are lacking, limiting our understanding of these processes in situ.

Objectives
To address questions about the role of evolutionary convergence and historical contingency in a “natural evolution experiment”, we investigate the molecular evolution of distinct clonal lineages of *Pseudomonas aeruginosa* from the initial invasion into the cystic fibrosis (CF) airways and onwards as they genetically adapt to a human host after transition from their environmental habitat.

Methods
We sequenced and analyzed the genomes of >500 longitudinally collected clinical isolates of *P. aeruginosa* sampled from >40 CF patients.

Conclusions
Parallel evolution across multiple *P. aeruginosa* strains identified convergent molecular evolution in pathoadaptive genes that revealed host-adaptation to involve remodeling of regulatory networks and central metabolism, acquisition of antibiotic resistance, and loss of extracellular virulence factors. Furthermore, we demonstrated how historical contingency played a role in the evolution of regulatory networks, in which mutations in downstream transcriptional regulators were contingent upon mutations in upstream regulators; suggesting that remodeling of regulatory networks facilitate adaptation. Knowledge of pathoadaptive mutations and evolutionary contingency may help prediction of bacterial evolution in CF patients and design of future intervention strategies.
Mäkelä-Cassell Awardee: Taxon-specific growth rates are related to phylogeny in soil bacterial communities

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Understanding if and how the phylogenetic composition of microbial communities is related to ecosystem function has been a central question in microbial ecology. To date, attempts to unambiguously identify the microbial taxa involved in an ecosystem process of interest have relied on stable isotope probing (SIP), where heavy isotopes are incorporated into the nucleic acids of microorganisms utilizing a compound of interest (¹³C carbon sources) or actively growing (¹⁸O water). Traditionally, this method has been qualitative. However, we have advanced this technique in order to quantitatively assess the amount of isotope incorporation into nucleic acids of specific taxa. This is achieved by comparing the naïve density of a DNA fragment to its density after incubation in the presence of a stable isotope, as the change in density is proportional to the amount of stable isotope incorporation. Using this technique, we have estimated the taxon-specific growth rates of soil bacteria under control conditions and in the presence of an added substrate (¹³C-glucose). The growth rates and ¹³C assimilation rates of the most abundant taxa were compared to their phylogenetic relatedness to determine if there was a phylogenetic signal (measured using Blomber's K test) associated with the growth response. Indeed, closely related organisms were similar in their growth responses. This phylogenetic signal was found after single (K=0.81, p<0.001) and repeated substrate addition (K=0.99, p<0.001), as well as in the absence of added substrate (K=0.73, p<0.001). The different resource conditions elicited growth of distinct, phylogenetically-clustered groups suggesting a microorganism’s taxonomic identity could be a useful indicator of its ecology in the environment. Generally, these data suggest that the growth responses and therefore nutrient transformations of soil microorganisms are significantly influenced by evolutionary history.
Compounds for Overcoming Antibiotic Tolerance and Resistance

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Compounds for Overcoming Antibiotic Tolerance and Resistance

Currently available antibiotics are largely inactive against dormant persister cells produced by microbial pathogens, and are prone to resistance development. Persisters are responsible for the chronic, relapsing nature of the disease. An even more daunting unsolved problem is the seemingly inevitable development of resistance to all known antibiotics. We find that different natural compounds targeting the Clp protease, acyldepsipeptide and lassomycin, kill persisters and eradicate the pathogen population. Teixobactin, a cell wall synthesis inhibitor was discovered in a screen of uncultured soil bacteria. Teixobactin is the first member of a new class of cell wall acting inhibitors and binds two targets, lipid II, precursor of peptidoglycan, and lipid III, precursor of wall teichoic acid. The targets are not proteins, and there is no resistance development to this compound. Soil microorganisms are likely to harbor many more sterilizing and largely resistance-free compounds. Screening uncultured bacteria provides an effective platform for antibiotic discovery.
Infectious diseases are one of the leading causes of death worldwide and the emergence of multi-drug-resistant bacteria, and the lack of new antibiotics in development with new modes of action are widely accepted as one of the major clinical concerns that require to be addressed from the public and private sectors. One of major clinical needs are related with MDR Gram-negative bacteria hospital-acquired infections, frequently associated with high mortality rates and few treatment options. Very limited options have been developed to date against Gram negative, and even last resorts antibiotics such as colistin and polymixin B are facing resistances.

Microbial natural products (NPs) have been one of the most prolific sources of new leads for the discovery of novel antibiotics, with a large number of compounds and analogs successfully introduced in the market in the past decades and still today in the clinic. NPs potency and selectivity is the result of an extended evolutionary selection to create biologically active molecules with the required properties to penetrate bacterial membranes and interact and potentially inhibit bacterial targets.

The strategy developed at MEDINA for the discovery for novel antibiotics is focused on the exploitation of our proprietary NPs collections, one of the most diverse and continuously expanding sources of novel compounds. Most recent efforts have permitted the identification of novel families of compounds that will be discussed in the context of current antibiotic discovery efforts in the academia and the pharma sector.
Antibiotic resistance remains a high unmet medical need. The situation is worsened by the scarcity of investigational new drugs in the antibacterial pipeline, especially against multidrug-resistant Gram-negative pathogens (Enterobacteriaceae, Pseudomonas aeruginosa, Acinetobacter baumannii). Novel antibiotic classes are clearly needed to overcome the current mechanisms of resistance faced by the clinicians. Among the different approaches to discover novel antibacterial classes, the mining of the microbial biodiversity is one of the most promising. In this context, the bacterial genera Xenorhabdus and Photorhabdus represent a high-potential bioresource that remains to be extensively explored. Xenorhabdus and Photorhabdus are symbiotic bacteria of the entomopathogenic nematodes Steinernema and Heterorhabditis. They have an original life-cycle that requires the production of a great diversity of bioactive secondary metabolites, including antimicrobials. This ecological rationale is supported by a high content of PKS and NRPS genes in their genomes. The screening of a collection of Xenorhabdus and Photorhabdus strains led to the discovery of the Odilorhabdins, a novel antibiotic class with a broad spectrum activity against Gram-positive and Gram-negative pathogens, including multidrug-resistant clinical strains. The Odilorhabdins have shown efficacy in murine infection models. They are currently undergoing preclinical optimization by medicinal chemistry. Odilorhabdins are expected to enter into clinical trials in human by 2017.
Srinivas et al., *Science* 2010, 327, 1010.


Cyclic di-nucleotides in bacteria

C-DI-GMP SIGNALING AND E. COLI BIOFILM ARCHITECTURE

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Background

Bacterial biofilms contain matrix-producing and matrix-free cells in distinct zones. In E. coli K-12 the production of matrix components such as amyloid curli fibres and cellulose is promoted by the second messenger c-di-GMP, which is synthesized by 12 diguanylate cyclases (DGC) and degraded 13 phosphodiesterases (PDE). Two PDE/DGC pairs (YhjH/YegE and YciR/YdaM) constitute a bistable switch device that can sharply drive up the cellular c-di-GMP level in slowly growing and stationary phase zones of macrocolony biofilms and thereby turn on the expression of the biofilm regulator CsgD. This transcription factor then activates expression of the curli genes as well as of yaiC, which encodes a DGC specifically required for cellulose synthesis.

Objectives

Recent progress in understanding this c-di-GMP signaling network will be presented.

Methods

Genetics, biochemistry, microscopy, mathematical modelling.

Conclusions

In the central c-di-GMP signaling network that controls E. coli biofilm matrix production, YegE is the 'starter' DGC, which - by producing c-di-GMP that is sensed by the 'trigger enzyme' YciR - promotes YdaM activity and YaiC expression. These two DGCs further increase c-di-GMP levels and thereby for a positive feedback loop. In addition, most of the other DGCs and PDEs are expressed but do not contribute to this scenario under standard conditions, suggesting that these enzymes are inactive. However, specific signals perceived by the various N-terminal sensor domains may activate these DGCs and PDEs and thereby modulate c-di-GMP levels, matrix production and biofilm morphology. As an example a novel subfamily of c-di-GMP-related enzymes will be presented whose activity is regulated by redox signal input.
Signalling nucleotides are key molecules in all domains of life and control fundamental processes including central metabolic or stress response processes as well as biofilm formation. c-di-AMP is one of the more recently discovered signalling molecules and is produced by a number of bacteria including pathogens such as *Staphylococcus aureus*. In *S. aureus*, c-di-AMP is synthesised from two molecules of ATP by the diadenylate cyclase DacA and degraded to pApA by the phosphodiesterase GdpP. c-di-AMP has since been implicated in controlling cell size in *S. aureus*, in helping this microorganism cope with cell wall stress and in the regulation of potassium uptake. Here, the identification and function of c-di-AMP receptor proteins will be discussed. Furthermore, the apo- and c-di-AMP complex structures of PstA, one of the c-di-AMP target proteins, will be presented. This allowed us to propose a mechanism on how PstA functions as a signaling transduction protein. The complex structure also highlighted common features of c-di-AMP binding sites allowing for a more rational prediction of cyclic-di-AMP binding sites.
Cyclic di-nucleotides in bacteria

Chp8, a composite diguanylate cyclase which promotes Pseudomonas syringae pv tomato DC3000 pathogenicity and plant immune evasion during infection of Arabidopsis thaliana

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Pseudomonas syringae (Ps) is a bacterial pathogen which interacts with more than 50 different plants causing diseases in crops including tomato, beans and rice. The associated decrease in crop yields results not only in economic losses, but also threatens global food security. Ps uses type-three-secretion (TTSS) to deliver virulence factors into the plant that promote survival of the bacterium and ultimately results in disease symptoms. The Ps TTSS is a product of the hypersensitive response and pathogenicity (hrp) and hypersensitive response and conserved (hrc) gene cluster under the control of sigma⁵₄, the co-dependent transcriptional activators HrpRS and the alternative sigma factor HrpL.

Here, I will present our findings that in the model strain Ps DC3000 HrpRS also activate expression of the composite diguanylate cyclase Chp8 (Co-regulated with hrp 8) in response to plant derived signals. Chp8 diminishes the hormonal immune response of Arabidopsis thaliana and promotes Ps DC3000 pathogenicity. Chp8 is the first example of a diguanylate cyclase promoting pathogenicity of a bacterial plant pathogen. The study was recently published in mBio (Engl et al., 2014) and appraised as the first complete description of how a diguanylate cyclase can affect bacterial-host interactions in an acute infection model.
FEMS-0860
Cyclic di-nucleotides in bacteria

THE ROLE OF CYCLIC DI-GMP IN STREPTOMYCES DEVELOPMENT
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Background

Streptomyces are non-motile soil bacteria with a unique mycelial, sporulating life cycle and the ability to produce highly valuable secondary metabolites. Streptomyces venezuelae encodes for 10 putative cyclic di-GMP-metabolizing enzymes but the cellular processes controlled by the cyclic di-nucleotide remained elusive.

Objectives

Aiming to gain more knowledge about the regulation of the developmental processes in S. venezuelae, we found that elevated levels of c-di-GMP lead to a pronounced delay of the developmental program. On the other hand, depletion of c-di-GMP results in an enhanced sporulation phenotype due to bypassing the aerial mycelium stage.

Methods

We applied a global pull-down approach using the c-di-GMP Capture Compound (Caprotec, Berlin) and found a c-di-GMP-responsive regulator that connects c-di-GMP signalling pathways and developmental processes in S. venezuelae. Using structural and biochemical analyses we identified a previously unseen oligomeric form of c-di-GMP, which enables the effector protein to dimerize by binding to a heretofore unknown binding motif.

Conclusions

Altogether, we demonstrate that the signalling molecule c-di-GMP controls the hyphato-spore transition in filamentous bacteria.
FEMS-2876
Bacterial persistence and Toxin - Antitoxins

Magic Spot Controls Bacterial Persister Cell Formation by Activating Toxin - Antitoxins
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Background
We showed previously that multidrug tolerance (persistence) of the model organism Escherichia coli K-12, depends stochastic induction of Toxin - Antitoxin (TA) activity by (p)ppGpp in a regulatory cascade that included also Lon protease and polyphosphate (Maisonneuve et al., Cell, 154, 1140-50, 2013). The TAs involved all encode inhibitors of translation, such as RelE, MazF, HicA etc. Remarkably, all antitoxins of E. coli K-12 encoded by type II TA loci are degraded by Lon that, in turn, is activated by polyphosphate. The model explaining stochastic induction of persistence is shown in Figure 1. Sophie Helaine and David Holden (Imperial College London, UK) showed that a similar mechanism allows cells of Salmonella to survive antibiotic treatment within macrophages (Science 343, 204–208, 2014).

Objectives
Many bacterial pathogens contain multiple type II TA genes and rely on (p)ppGpp to be virulent (Figure 2). We will determine the role(s) that (p)ppGpp and TAs play in persistence and virulence of different bacterial pathogens, including, Mycobacterium tuberculosis, Burkholderia cenocepacia and Photorhabdus luminescense.

Methods
In this work, we will employ physiological, genetic, cytological and biochemical methods to reach our aim.

Conclusions
Preliminary results indicate that indeed (p)ppGpp is required for persistence of bacterial pathogens.
FEMS-3162
Bacterial persistence and Toxin - Antitoxins

FIC domains in *Bartonella*: Evolution of diversified host effectors from a widely-spread bacterial toxin/antitoxin system

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The FIC domain being present in thousands of proteins and found in all domains of life mediates post-translational modification of target proteins, typically by transfer of an AMP moiety from ATP onto a target tyrosyl or threonyl side chain by a process called AMPylation or adenylylation. The alpha-proteobacterial genus *Bartonella* comprises facultative intracellular pathogens that utilize a VirB type IV secretion system to translocate a cocktail of bacterial effectors protein into host cells in order to subvert cellular function to the benefit to the bacterial intruder. Diversified copies of the FIC domain are prominently presented in these effector sets, indicating that the various FIC domain variants may mediate diverse effector functions during infection. I will discuss the evolutionary origin of the FIC domain as a bacterial toxin-antitoxin system, its fusion with a type IV secretion signal to constitute an ancestral translocateable effector, as well as the duplication and diversification of the ancestral effector gene in the course of the evolution of the effector cocktails that facilitate host cell infection by bartonellae in diverse mammals.
Prokaryotic toxin-antitoxin modules are involved in the establishment of persister cells. The latter involves complex regulatory mechanisms that link the regulation of protein activity to regulation of transcription. Recent structural and biochemical data shows a plethora of molecular mechanisms to achieve this goal. In the phd/doc module, conditional co-operativity is established via a combination of negative co-operativity through entropic exclusion of and IDP domain combined with a low to high affinity switch for the interaction between toxin and antitoxin. While Doc stabilizes the DNA binding conformation of Phd, Phd likewise inhibits the kinase activity of Doc and simultaneously prevents misfolding of Doc. The same phenomenon of conditional co-operativity is observed for mazEF and ccdAB, but with a different structural basis, although intrinsic disorder in the antitoxin is a common theme. Other TA modules such as higBA are regulated through a seemingly more simple mechanism where the antitoxin acts as the sole repressor, while toxin significantly weakens operator binding. Such a mechanism is observed in several higBA modules as well as in mqsRA and allows for transcription regulation without the specific need of intrinsic disorder.
Bacterial persistence and Toxin - Antitoxins

Chaperone-mediated control of toxin-antitoxins

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Bacterial toxin-antitoxin systems are stress-responsive elements generally composed of a stable toxin that forms an inactive complex with its less stable cognate antitoxin. In response to specific stress conditions the antitoxin is degraded by stress proteases and the free active toxin subsequently targets important cellular processes such as DNA replication or protein synthesis. It is believed that the resulting growth inhibition facilitates adaptation to stress and persistence. *Mycobacterium tuberculosis*, the causing agent of human tuberculosis, encodes 79 putative toxin-antitoxin systems and it has been proposed that persistence induced by active toxins might be relevant for its pathogenesis. The tripartite toxin–antitoxin–chaperone (TAC) system of *M. tuberculosis* is an atypical toxin-antitoxin system strongly induced in persisters and tightly controlled by the molecular chaperone Rv1957, which is related to the canonical SecB chaperone involved in Sec-dependent protein export in Gram negative bacteria. Indeed, in spite of very little sequence similarity, Rv1957 is able to efficiently replace SecB during protein export in *E. coli*, and to specifically control the functional HigB1-HigA1 toxin-antitoxin system of *M. tuberculosis*. How does the mycobacterial SecB-like chaperone respond to stress and control the toxin activation cascade, and to what extent such activation is important for *M. tuberculosis* persistence and virulence are so far unresolved questions.

In this work, we will present recent data concerning the molecular mechanism of TAC complex formation and toxin activation, with emphasis on the role played by the SecB-like chaperone.
Type II CRISPR-Cas9 systems: mechanisms, evolution and applications
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The RNA-programmable CRISPR-Cas9 system has recently emerged as a transformative technology in biological sciences, allowing rapid and efficient targeted genome editing, chromosomal marking and gene regulation. In this system, the endonuclease Cas9 or catalytically inactive Cas9 variants are programmed with single guide RNAs (sgRNAs) to target site-specifically any DNA sequence of interest given the presence of a short sequence (Protospacer Adjacent Motif, PAM) juxtaposed to the complementary region between the sgRNA and target DNA. The system is efficient, versatile and easily programmable.

Originally, CRISPR-Cas is an RNA-mediated adaptive immune system that protects bacteria and archaea from invading mobile genetic elements (phages, plasmids). Short crRNA (CRISPR RNA) molecules containing unique genome-targeting spacers commonly guide Cas protein(s) to invading cognate nucleic acids to affect their maintenance. CRISPR-Cas has been classified into three main types and further subtypes. CRISPR-Cas9 originates from type II CRISPR-Cas that has evolved unique molecular mechanisms for maturation of crRNAs\(^a\) and targeting of invading DNA\(^a,b\) which we identified in the human pathogen Streptococcus pyogenes. On the basis of the discovery of the DNA targeting mechanism, we proposed that RNA-programmable Cas9 could offer considerable potential for genome editing in cells of the three kingdoms of life for biotechnological, biomedical and gene-therapeutic purposes\(^b\). As demonstrated by a large number of studies published in the last 18 months, DNA targeting by CRISPR-Cas9 has quickly been adopted by the scientific community to edit and silence genomes in a large variety of cells and organisms including human cells, plants and mice. I will discuss the biological roles of CRISPR-Cas9, the mechanisms involved, the evolution of type II CRISPR-Cas components in bacteria and the applications of CRISPR-Cas9 as a novel genome engineering technology.
Specific nucleic acid targeting by the Type III CRISPR-Cas complexes

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CRISPR-Cas is a prokaryotic adaptive immune system that provides sequence-specific defense against foreign nucleic acids. Comparative analysis has revealed at least 3 distinct types of CRISPR-Cas systems. During the past few years, much has been learned on the structures and mechanisms of the effector complexes of Type I (Cascade) and Type II (Cas9). Until recently, structural and functional insights on the effector complexes of Type III CRISPR-Cas systems has lagged behind. To address this, the Type III-A (Csm) and III-B (Cmr) complexes of Thermus thermophilus have been selected as model systems. Integrated structural analyses (MS, EM) have revealed that the overall architecture of both complexes strongly resembles that of the Type I Cascade complexes. TtCsm is composed of five different protein subunits (Csm1-Csm5), whereas TtCmr has six subunits (Cmr1-Cmr6). Like Cascade, both Type III complexes have an uneven stoichiometry and a single crRNA guide. Interestingly, the TtCsm crRNA content is similar to the Type III-B Cmr complex, indicating that crRNAs are shared among different subtypes. Unexpectedly, both TtCmr and TtCsm cleave complementary target RNAs at multiple sites. Outstanding questions of the Thermus Type III systems include (i) a motif for auto-immunity protection (PAM analog), (ii) some target quality control (seed analog), and (iii) last but not least the potential of RNA-guided DNA targeting. The latest insights in the TtCsm and TtCmr mechanisms will be discussed.

Staals, R.H. et al. (2013) Mol Cell


An adaptive immune system in prokaryotes called CRISPR (clustered regularly interspaced short palindromic repeats) uses small guide RNAs to neutralize invading viruses and plasmids. In Escherichia coli, immunity depends on a ribonucleoprotein complex called Cascade. This protein complex recognizes double-stranded DNA from invaders by forming basepairs between the guide RNA and invader DNA. The structure of Cascade displays an unusual seahorse-shape that binds target DNA in an unique underwound ribbon like duplex. This recruits the effector nuclease Cas3 which progressively degrades the target DNA in an ATP-dependent manner, neutralizing the virus infection. Apart from giving insight into the crystal structure and function of Cascade, I will also show how viruses are able to escape immunity by point mutagenesis, and unveil how hosts rapidly adapt their immune specificity making use of prior memories to effectively counter escape mutants.


FEMS-1197
CRISPR - biological and technological advances

Studying and fighting bacteria with the help of CRISPR
Establishment of the fungal mycelium

Heterogeneity of the fungal mycelium

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Fungi form colonies that consist of a network of hyphae. Macro-colonies with a diameter > 1 cm are formed on solid media, while micro-colonies with a diameter in the mm-scale are formed in liquid shaken cultures such as bioreactors. Research in our lab has revealed that micro- and macro-colonies are not a collection of hyphae with a synchronized activity. Fungal micro-colonies within a liquid culture are heterogeneous in gene expression and size. Heterogeneity in RNA composition can even be found between and within zones of macro- and micro-colonies. These findings imply that gene regulatory networks will not be correctly predicted by using whole cultures and colonies. This will be illustrated by regulatory networks of amylolytic genes and genes involved in asexual development. The finding of heterogeneity came to a surprise considering the fact that hyphae of higher fungi are compartmentalized with septa that contain large pores. These pores enable cytoplasmic streaming. It has now become clear that cytoplasmic streaming in aspergilli is reduced by dynamic plugging of septa. Modelling has shown that even open septa are a barrier for cytoplasmic mixing.
Establishment of the fungal mycelium

The microtubule cytoskeleton in Aspergillus nidulans

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Polar growth in fungi requires intact microtubule and actin cytoskeletons. Both cytoskeletons consist of filamentous structures composed of alpha/beta tubulin dimers and actin monomers, respectively. The formation of both cytoskeletons is interdependent and relies on a class of so call cell-end marker proteins. The organization of the two cytoskeletons guarantees the directed travel of secretory vesicles. We found in Aspergillus nidulans that hyphal extension occurs in pulses at small areas at the hyphal tip rather than uniformly. Sites of secretion are only maintained for a short period before new sites are established.

A. nidulans harbors two genes encoding alpha tubulin and two genes encoding beta-tubulin. However, 2-D protein analyses revealed early on that tubulin can be post-translationally modified. Such modifications are a common phenomenon in eukaryotes, but their role is largely unknown. One modification is the C-terminal removal of tyrosin from alpha tubulin. The missing tyrosin can be re-added to alpha/beta tubulin dimers. Hence there is a cycle of detyrosination and tyrosination which determines the amount of modified alpha tubulin. Such modifications have not yet been described in fungi. We found that the kinesin 3 motor UncA binds preferentially to a subpopulation of microtubules in A. nidulans and also in N. crassa. We have evidence that the subpopulation consists of detyrosinated alpha tubulin and that the tail of UncA is required for the specificity. However, recent results suggest that the UncA tail harbours a microtubule-binding site and that the motor can interact with the motor domain with one microtubule and with the tail with a adjacent one. Thus UncA could be involved in bundling of microtubules.
Intercellular communication is critical for the survival of unicellular microbes as well as for the development and function of multicellular tissues in higher eukaryotes. Cell-to-cell signaling is also required to develop the interconnected mycelial network characteristic of filamentous fungi and is a prerequisite for symbiotic and pathogenic host colonization achieved by molds. Somatic cell-cell communication and subsequent cell fusion is governed by the MAK-2 mitogen activated protein kinase (MAPK) cascade in the filamentous ascomycete model Neurospora crassa, yet the composition and mode of regulation of the MAK-2 pathway are currently unclear. In order to identify additional components involved in MAK-2 signaling we performed affinity purification experiments coupled to mass spectrometry with strains expressing functional GFP-fusion proteins of the MAPK cascade. This approach identified STE-50 as regulatory subunit of the yeast Ste11 homolog NRC-1 and HAM-5 as cell-communication-specific scaffold protein of the MAPK cascade. Moreover, we defined a network of proteins consisting of two Ste20-related kinases, the small GTPase RAS-2 and the adenylate cyclase capping protein CAP-1 that function upstream of the MAK-2 pathway and whose signals converge on the NRC-1/STE-50 MAP3K complex and the HAM-5 scaffold. Finally, our data imply the COT-1 NDR kinase network as effector module of the MAK-2 pathway to control the chemotrophic behavior of the two communicating cell tips. Taken together, these data have implications for our mechanistic understanding of MAPK pathways and their regulation during intercellular communication in eukaryotic microbes.
FEMS-1744
Establishment of the fungal mycelium

Insights into the early regulation of conidiospore development in Aspergillus nidulans

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Fungal hyphae are capable of undergoing different developmental programmes, thus enabling the mycelium to perform diverse biological functions simultaneously.

Aspergillus nidulans mycelia expand by means of vegetative hyphae at the colony periphery. At sub-peripheral regions, hyphae give rise to conidophores bearing aerial mitospores (conidiospores) suitable for dispersal. At distal regions, hyphae support cleistothecium formation and eventually meiospores (ascospores), with long-term survival capabilities. The relative proportions of these cell-types can be further modulated to suit environmental conditions.

Three decades of molecular genetic research have unveiled the most important factors participating in the regulation of conidiophore development. Those already expressed in vegetative cells prior to the onset of development have been shown to participate in cell fate determination. Moreover, their characterisation has revealed that protein-protein interactions between these factors and their counterparts for cleistothecium formation, into protein complexes, are at the core of cell fate determination. Indeed, the alternative patterns of complex formation may well determine the alternative patterns of development.

In this presentation, we review the interaction patterns described for the velvet family of proteins VeA, VelB, VosA and VelC, the red light photosensor FphA, and the blue light photosensors LreA and LreB. We shall also incorporate interactions involving FluG and FlbA-E, developmental regulators specifically associated with conidiophore formation. An updated model of interactions leading to the determination of alternative developmental programmes will be presented.
Heat is an unavoidable side-product of all life processes. Already in 1784 reported the father of quantitative chemistry – Antoine Laurent de Lavoisier – on the first measurement of the metabolic heat of a guinea pig and made first attempts to offer an interpretation. The main advantages of heat as an indicator for life processes are the reflection of any metabolic changes in real time and the tight correlation to metabolic fluxes and thermodynamic driving forces. Nowadays, even commercially available calorimetric instruments achieve detection limits low enough to measure the heat of 100,000 aerobically growing bacteria or of 100 myocardial cells. Heat can be monitored in reaction vessels ranging from a few nanoliters up to many cubic meters. In contrast to many traditional and emerging microbiological methods, heat flux measurements do not need labeling agents and reactants and also works with optically opaque solutions. It is further possible to assemble the thermal transducer in a protected way that reduces aging and thereby signal drifts.

All of these advantages make calorimetry an interesting tool for many applications in medicine and biotechnology. Unfortunately, metabolic heat signals are currently mainly used to monitor and control biological processes and their full information potential remains unexploited. However, new omics technologies offer a flood of information about complex biological and ecological systems. These detail information open the chance to be combined with calorimetric measurements and thermokinetic modelling to provide a holistic picture of the complex system under consideration. Various methods for quantitative data interpretations and their dependency on the applied calorimetric device and available additional biometric data will be presented and discussed. Also potential misinterpretations are considered.
A novel biochemical trick of anaerobic microorganisms to deal with complicated thermodynamic situations

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Flavin-based electron-bifurcation is a recently discovered mechanism of coupling endergonic to exergonic redox reactions in the cytoplasm of anaerobic bacteria and archaean (1-7). The seven electron-bifurcating complexes characterized to date have in common that they catalyze the reduction of ferredoxin and of a second electron acceptor with NADH, NADPH, dihydrogen or formate as electron donors, the latter having a redox potential between that of ferredoxin and of the second electron acceptor. Via this mechanistic "trick" low potential electrons are generated that are used for energy conservation and biosynthesis in strict anaerobe such as Clostridia, acetogenic bacteria, methanogenic archaean and sulfate reducing bacteria.

References

A thermodynamic theory of microbial growth and its perspectives for a better understanding of community dynamics

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The study of microbes, the simplest form of life, constitutes a fertile thinking ground for a deeper interlinking between physical and biological concepts. In 1922, Alfred Lotka suggested that the similarity between individuals was an invitation to imagine a "statistical mechanics of living beings". As an extension of these ideas, we proposed a thermodynamic theory of microbial growth by showing how systems constituted by microbes in contact with molecules could be likened to ensembles described by the laws of statistical physics. Based on thermodynamic balances established by different authors, we defined the "activation energy" of a microbe and thus the probability for an elementary division act to be triggered. A growth equation could be proposed, which links a flux (the growth of microbes) to a force (the free energy density in the environment). The equation allows adequate modeling of experimental growth data. More importantly, it also allows making new predictions in relation to the microbial isotopic fractionation phenomenon which can be viewed as a kinetic consequence of the differences in energy contents of isotopic isomers used for growth. We showed how recent experimental data actually support these original predictions. In the future, we plan to use our thermodynamic growth equation for modelling microbial community dynamics and studying the convergence of microbial ecosystems. Indeed our law offers a mathematical framework that may help understand the relevance of thermodynamic ecological goal functions such as dissipated power, exergy or entropy production.

Mechanistic basis of equal plasmid spacing by the parABC system

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Low copy number plasmids in bacteria require segregation for stable inheritance through cell division. This is often achieved by a parABC locus, comprising an ATPase ParA, DNA-binding protein ParB and a parC region, encoding ParB-binding sites. These minimal components space plasmids equally over the nucleoid, yet the underlying mechanism is not understood. Here we investigate a model where ParA-ATP can dynamically associate to the nucleoid and is hydrolyzed by plasmid-associated ParB, thereby creating nucleoid-bound, self-organizing ParA concentration gradients. We show mathematically that differences between competing ParA concentrations on either side of a plasmid can specify regular plasmid positioning. Such positioning can be achieved regardless of the exact mechanism of plasmid movement, including plasmid diffusion with ParA-mediated immobilization or directed plasmid motion induced by ParB/parC-stimulated ParA structure disassembly. However, we find experimentally that parABC from Escherichia coli plasmid pB171 increases plasmid mobility, inconsistent with diffusion/immobilization. Instead our observations favor directed plasmid motion. Our model predicts less oscillatory ParA dynamics than previously believed, a prediction we verify experimentally. We also show that ParA localization and plasmid positioning depend on the underlying nucleoid morphology, indicating that the chromosomal architecture constrains ParA structure formation. Our directed motion model unifies previously contradictory models for plasmid segregation and provides a robust mechanistic basis for self-organized plasmid spacing that may be widely applicable.

Although it is often tacitly assumed that gene regulatory interactions are finely tuned, it is unclear how accurate gene regulation could evolve de novo from a state without regulation. Moreover, gene expression noise would seem to impede the evolution of accurate gene regulation, and the fact that expression noise varies greatly across genes suggests natural selection has affected noise levels. To infer how selection has tuned noise levels of E. coli promoters, we designed a novel synthetic biology approach in which we evolved a large library of synthetic promoters de novo. We show that, surprisingly, promoters exhibit low noise by default, and rather than having acted to lower noise levels, selection must have acted to raise noise levels of a substantial fraction of E. coli promoters. In addition, promoters with elevated noise levels are precisely those that are highly transcriptionally regulated.

To explain the general association between expression noise and gene regulation, it has been argued that elevated expression noise may either be directly selected as a 'bet hedging strategy', or that it may be a detrimental but unavoidable side-effect of regulation. We here present a general theory for the interplay of expression noise and gene regulation which shows that this is a false dichotomy: expression noise can be both a necessary consequence of regulation and a beneficial bet hedging strategy at the same time. Moreover, this theory elucidates how regulation can evolve incrementally starting from a state without regulation. Noise propagation from regulators to targets acts as a rudimentary form of regulation, and noisy expression is not a nuisance but rather a stepping stone toward the emergence of accurate gene regulation.
Elucidating the role of molecular stochasticity in cellular growth is important to understanding phenotypic heterogeneity and the stability of cellular proliferation. We used time-lapse microscopy to measure fluctuations in the instantaneous growth rate of single cells of Escherichia coli and in the expression of metabolic enzymes. We show that expression fluctuations of catabolically active enzymes can propagate and cause growth fluctuations. Conversely, growth fluctuations propagate back to perturb expression. Homeostasis is promoted by a noise-cancelling mechanism that exploits fluctuations in the dilution of proteins by cell-volume expansion. Thus, molecular noise is propagated not only by regulatory proteins but also by metabolic reactions. The results suggest that cellular metabolism is inherently stochastic, and a generic source of phenotypic heterogeneity.
How bacteria can maximise their growth rate under stress conditions

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Large population-sizes and the exponential increase in the abundance of fit genotypes makes growth-rate selection a powerful force that shapes bacterial survival strategies. Bacteria have to therefore carefully balance growth and stress tasks to prevent being outcompeted during nutrient-excess and stress conditions. In this talk, I will discuss several principles of bacterial-fitness optimisation, starting from basic relations of biochemistry, steady-state metabolism and growth-rate maximisation. The resulting theory explains how bacteria should balance growth and stress tasks to optimise fitness. Using this theory, several experimental findings can be explained. Those findings include how E. coli regulates its ribosome synthesis via ppGpp, the fitness costs of protein overexpression and the shape of fitness landscapes.
Bio-synthetic routes based on new enzymatic reactions

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Traditionally, new enzyme activities have been discovered by screening microorganisms and mutant libraries. In recent years, these strategies were successfully expanded by a mechanism-based approach. Hereby, the catalytic machinery of known enzymes is harnessed to perform not only the same type of chemistry with different substrates but also to catalyze completely new non-physiological reactions. To identify initial activity and to address the catalytic flexibility, small and functionally rich enzyme libraries have been used. This presentation will highlight hidden as well as uninvestigated enzymatic activities and will illustrate strategies to unmask them. For example, we have recently harnessed the unique protonation machinery of squalene hopene cyclases to generate enzymes for non-natural chiral Brønsted acid catalysis.

The identification of novel enzymatic transformations enables complete new biosynthetic routes to interesting molecules. Examples like menthol, sebacic acid or heterocyclic compounds will be given which also explains how these new reactions can be applied in a retrosynthetic approach.
Microbial biosynthesis and separation of alkanes and alcohols

Economically sustainable conversion of solar energy and CO$_2$ into low value chemical energy (e.g. fuel) using biotechnology is a challenge. Several benefits can be imagined if the chemical product can be immediately separated from the (1) organisms that synthesized the product and (2) biotechnological process. These include enhanced product tolerance, reduced feedback inhibition, and enhanced pathway thermodynamics. In addition, by continuously removing the product, a continuous biotechnological process is possible. This allows the biological organism to act as a catalyst rather than as a substrate for further transformation. Several strategies for immediate product-process separation are possible: (a) a gaseous product spontaneously leaves the liquid phase, (b) excretion and engineered separation of a water-soluble liquid, (c) excretion and spontaneous separation of a hydrophobic liquid. In all above cases, it is necessary for the product to transfer through the cell envelope of the biocatalyst. Towards the objective of producing a useful fuel that can be easily separated, we have constructed several synthetic pathways for alkane and alcohol biosynthesis. The challenges and insight gained from this metabolic pathway and transporter engineering work will be summarised.
Parallel but distinct evolution of Pseudomonas aeruginosa and Acinetobacter baumannii toward pan-drug resistance

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Pseudomonas aeruginosa and Acinetobacter baumannii have emerged as major nosocomial pathogens in the second half of the 20th century, as unexpected consequence of medical progress. These strictly aerobic bacilli share a high metabolic versatility and the ability to thrive in hostile environments. In addition to producing a complex array of virulence factors enabling them to infect fragile patients, both pathogens show a remarkable capacity to become resistant to all the antibiotics available. However, despite these apparent similarities, P. aeruginosa and A. baumannii differ in the mechanisms leading to such a high degree of resistance. Total recalcitrance of P. aeruginosa to β-lactams, the major class of antibiotics used in therapeutics, mostly relies on mutation-driven mechanisms (overproduction of intrinsic β-lactamase AmpC, porin OprD deficiency, activation of efflux systems) operating synergistically with a variety of extended-spectrum β-lactamases collected from environmental reservoirs via horizontal gene transfers. The parallel evolution of A. baumannii to pan-β-lactam resistance involves IS-dependent up-regulation of natural β-lactamase ADC combined with conjugational acquisition of a limited number of OXA-type β-lactamases from closely-related species. Accumulating evidence suggests that the two microorganisms exploit distinct pools of resistance genes (resistomes) to reinforce their own resistance mechanisms, probably as a result of distinct ecological niches.
Antimicrobial resistance: a touchy interface between human and veterinary medicine

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The increasing challenge due to antimicrobial resistance of pathogenic bacteria in human-, and veterinary medicine prompted the general fear that several classes of antimicrobials are becoming more or less useless due to their prophylactic and/or therapeutic use for animals. In light of new knowledge on genomic flexibility and on clonal spreading of pathogens and/or commensal bacteria we aim to re-visit potential links of antimicrobial resistance between animals and man.

Data about possible ways of evolution of resistances against the most critical antimicrobials (third generation cephalosporins, fluoroquinolones and aminoglycosides) show that foodborne enteric pathogens like Salmonella and Campylobacter seem to be trafficking frequently from animals to man, with the occasional opportunity to transfer critical resistance determinants. In contrast to enteric pathogens, the dominant clones of non-enteritis causing antibiotic resistant bacteria - methicillin-resistant Staphylococcus aureus (MRSA), vancomycin-resistant Enterococcus faecium, multiresistant Pseudomonas aeruginosa - usually differ substantially in the human healthcare setting from those occurring in animals. Adaption of distinct clones of bacteria to various species of animals or to humans may have contributed to the observed clonal diversity. The use of fluoroquinolone type antibiotics could also emerge as an enhancer of major international clones of MRSA and of some additional multiresistant pathogens.

Thus, evolution and spread of resistant enteric isolates is frequently driven by the development of some major clones originating and spreading from/within human clinical settings, while the resistant clones of animal/food origin seem to be more diverse with a partial human overlap. Therefore, the differences in spreading antimicrobial resistance within and between differing epidemiological compartments of animals and humans should be considered more analytically.
A NOVEL D-ALANINE-D-ALANINE LIGASE GENE FROM HUMAN ORAL METAGENOMIC DNA CONFERS HIGH-LEVEL RESISTANCE TO D-CYCLOSERINE

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Background

The human oral cavity is a reservoir for antibiotic resistance genes. While investigating integrons (a natural gene capture and expression system) and their associated gene cassettes from the human oral cavity, we identified two variants of D-alanine-D-alanine ligase (ddl) predicted to encode novel variants of the protein. The Ddl enzymes catalyse the synthesis of D-Ala-D-Ala dipeptide during the early stage of peptidoglycan biosynthesis. D-Cycloserine (DCS) is a Ddl inhibitor and is used as a second-line antibiotic to treat multidrug-resistant and extensively drug-resistant Mycobacterium tuberculosis. Overexpression of ddl genes from either Mycobacterium smegmatis or M. tuberculosis has been shown to confer resistance to DCS in M. smegmatis.

Objectives

The aim of this study is to test the hypothesis that ddl, identified on integron gene cassettes from a human oral metagenome may confer resistance to DCS.

Methods

Minimum Inhibitory Concentration (MIC) of DCS in Escherichia coli and Bacillus subtilis overexpressing the ddl variants was determined by agar dilution method according to the guidelines of Clinical and Laboratory Standards Institute (CLSI).

Conclusions

We observed that one variant of ddl confers resistance to DCS in E. coli and B. subtilis with an MIC of 128 µg/ml and 256 µg/ml, respectively. The other variant confers resistance with an MIC of 64µg/ml and 256 µg/ml, respectively.

These results represent the first time that ddl has been found on an integron cassette suggesting that it is important in host adaptation to stress (e.g. cell wall targeting antibiotics). The mechanism by which Ddl confers DCS resistance is currently being characterised.
GUT RESISTOME DEVELOPMENT IN NEONATES MONITORED IN A COHORT STUDY
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Background

The antibiotic resistance of pathogenic bacteria has been studied extensively. However, recently the collection of antibiotic resistance genes (ARGs) in the gut microbiota (resistome) has been acknowledged as an important reservoir of ARGs. The establishment of the human gut resistome is ideally studied in the pediatric microbiota.

Objectives

Our objective was to investigate the presence of various ARGs in the microbiota of newborns in the weeks following birth.

Methods

Within a prospective cohort among neonates born between 2000-2007, fecal samples collected at the ages of 5, 13 and 31 weeks postpartum were analyzed from a random selection of 120 infants. Inclusion criteria included healthy newborns at term, birth weight of >=2500g, and informed consent. Fecal samples were subjected to targeted metagenomics by qPCR for the detection of ARGs cfxA, tetM, tetQ, aac-aph, ermB and qnrS. The microbial composition in all samples was determined in a previous study. Prevalence was compared by using a McNemar’s test for paired samples. Risk factors were analyzed by using logistic regression.

Conclusions

Results show that acquisition of ARGs in the gut microbiota occurs already within the first weeks after birth and that the prevalence of these genes in the microbiota of newborns is highly variable over the course of a few weeks (Figure). Mode of delivery as well as duration of breastfeeding brought about significant changes in the prevalence of ARG. Relating ARG prevalence to microbiome analysis indicated that
changes in microbial composition are partly responsible for shifts in the resistome.
MOLECULAR CHARACTERISTICS OF EXTENDED-SPECTRUM CEPHALOSPORIN-RESISTANT ENTEROBACTERIACEAE FROM HUMANS IN THE COMMUNITY DIFFER FROM THOSE IN BROILERS

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Background
ESBL/AmpC-producing Enterobacteriaceae have frequently been reported in broilers and therefore they have been considered as a reservoir for ESBL/AmpC-encoding resistance genes with the potential to transmit to humans.

Objectives
The objective was to investigate the molecular characteristics of extended-spectrum cephalosporin (ESC)-resistant Enterobacteriaceae collected during a cross-sectional study examining the prevalence of faecal carriage of extended-spectrum β-lactamase (ESBL)-producing Enterobacteriaceae in a population sample of humans living in areas with high or low broiler density.

Methods
ESC-resistant Enterobacteriaceae were identified by combination disc-diffusion test. The presence and composition of ESBL/AmpC/carbapenemase encoding genes were assessed by PCR and sequencing. For Escherichia coli, phylogenetic groups and MLST were determined.

Conclusions
175 ESC-resistant Enterobacteriaceae were cultured from 165 of 1,033 sampled persons, yielding a prevalence of 15.9 %. No carbapenemase genes were identified. Most isolates were Escherichia coli followed by Citrobacter freundii and Enterobacter cloacae. The predominant ESBL genes were blaCTX-M-1, blaCTX-M-15 and blaCTX-M-14, but other allelic variants were also identified as well blaSHV-12 and blaTEM-52. The most common AmpC genes were blacMY-2 and blacMY-48-like. A large variety of E. coli genotypes was found, ST131 and ST10 being most common. ESBL/AmpC genes in Enterobacteriaceae obtained from the community resembled those found in isolates from patients in Dutch hospitals, indicating that healthy humans act as a reservoir for transmission of these determinants to vulnerable people. In contrast, the molecular characteristics of E. coli isolates differed from those of broilers suggesting that
broilers are not an important source of ESBL/AmpC genes for humans living in the community.
Antibiotic resistance: human and veterinary medicine

TRACKING IS26 MEDIATED IN VIVO GENE-SHUFFLING EVENTS ORIGINATING FROM A COMPLEX MULTI-DRUG RESISTANCE LOCUS

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Background

The insertion element IS26 is known to play a crucial role in the rapid evolution of complex drug resistance loci (CRL) of Gram-negative pathogens by shuffling and/or disseminating clustered drug resistance genes. Until recently, the unique molecular mechanisms involved in the mobilization of DNA fragments flanked by IS26 elements were unknown. A model, describing the generation of diverse types of IS26-derived mobile units has now been established using simple artificially constructed resistance units to support the proposed molecular mechanisms. To the best of knowledge, the impact of IS26-mediated gene reshuffling events on naturally occurring CRLs containing multiple copies of IS26 has never been demonstrated.

Objectives

To characterize IS26-mediated laterally mobile resistance subunits that have the potential to originate from a specific complex resistance locus.

Methods

Combinations of illumina and Sanger DNA sequencing, gene cloning and qPCR analytical techniques were used.

Conclusions

We identified a 27kb resistance locus (containing seven copies of IS26 and genes conferring resistance to five different classes of clinically relevant antibiotics) from the genome sequence of an Escherichia coli isolate belonging to sequence type 405, a major emerging uropathogenic E. coli clone implicated in the global dissemination of the blaCTX-M15 gene. We have irrefutably tracked formation of five, of 16 possible, laterally mobile units encoding resistance to different combinations of clinically important antibiotics from the 27kb CRL. Our assays on the CRL documents, for the first time, IS26-mediated horizontal gene mobilisation events in-vivo and experimentally confirm the imperative role played by IS26 in rapid evolution of CRL.
Microbial transformations of metals and metalloids

Ecology and mechanisms of microbial Fe(II) oxidation

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The two most important redox states of iron in the environment are Fe(II) [ferrous iron] and Fe(III) [ferric iron]. Dissolved Fe(II) and relatively soluble Fe(II) minerals are abundant in many environments including pH-neutral soils, sediments and iron seeps. Oxidation of dissolved and solid-phase ferrous iron [Fe(II)] at neutral pH can be catalyzed by microaerophilic, nitrate-reducing and even phototrophic microorganisms and leads to the precipitation of poorly soluble Fe(III) minerals. These minerals influence the environmental fate of many nutrients and contaminants and the precipitation of Fe(III) minerals by these microbial processes has been shown to occur in many modern environments and has been suggested to have been present already on early Earth.

This presentation will summarize the current knowledge and show results regarding mechanisms, physiology, and ecology of microbial Fe(II) oxidation. Special focus will be on microaerophilic Fe(II)-oxidizing bacteria that thrive in gradients of ferrous iron and oxygen and on phototrophic and nitrate-reducing Fe(II)-oxidizing bacteria in freshwater and marine sediments. I will present data on the isolation and quantification of microaerophilic, nitrate-reducing and phototrophic Fe(II)-oxidizers from different habitats as well as results regarding the quantification of their activities under different environmental conditions.
Microbial oxidation of metal sulfur minerals

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MICROBIAL OXIDATION OF METAL SULFUR MINERALS

Microorganisms can significantly impact oxidation rate of sulfur minerals such as pyrite, sphalerite, galena and arsenopyrite and thus regulate sulfur and metals cycles in various environments. A wide range of acidophilic bacteria and archaea are capable of catalyzing sulfur oxidation near the Earth's surface resulting in production of highly acidic, sulfate-rich solutions known as acid drainage which represents extreme geochemical conditions. Recent molecular ecology studies revealed that extreme geochemical conditions such as acid drainage sites that were previously thought lack of life due to limited nutrients and extreme conditions have diverse microbial populations. These DNA based findings open new directions for research dealing with sulfur oxidation by emphasizing interactions between microbe and sulfur minerals.

Elucidating the role of microorganism on dissolution of metal sulfide minerals is the key to understand environmental consequences of metal sulfur oxidation and developing of proper remediation techniques. Dissolution and oxidation of metal sulfur minerals usually generates different sulfur species with a wide range of valence states from sulfide (2-) to sulfate (+6). One way to understand metal sulfur oxidation pathway is to use O and S isotopes of reactants and products that are first witness of dissolution reactions. Biotic and abiotic metal sulfide oxidation experiments (pyrite, sphalerite, galena, arsenopyrite and S⁰) was carried out with different Acidophilic spp. species under various environmental conditions. Modern analysis techniques in addition to δ¹⁸O and δ³⁴S measurements of sulfur species were used to reveal oxidation mechanism (biotic vs. abiotic) of each sulfur mineral. Results will be discussed in detail to step towards understanding of microbial role on the dissolution kinetics of sulfur minerals.

Key Words: Enzyme, metal sulfur, isotope, DNA, dissolution
Background
The prices of some economically strategic elements, e.g. indium, germanium, rhenium, vanadium, and antimony, due to their use for high-tech products have increased to an extent which raises the question, whether these elements can be obtained from complex ores or secondary resources also in Germany.

Objectives
The aim of the studies was to identify bacterial cultures and conditions under which metals mentioned above by microbial activity can be brought into solution. Indium was to be obtained from sphalerite (ZnS) of the ore mountain region around Freiberg, the other strategic metals from "Theisen sludge" which resulted from fine-grained flue dust of former copper smelters and which had been deposited in the Mansfeld area, Germany.

Methods
For indium from sphalerite molecular analyses of the microbial community of acidic waters in the mine "Reiche Zeche" were performed to identify suitable leaching communities. To obtain other elements from "Theisen sludge" appropriate strains were enriched. All leaching experiments were initially performed in shake flaks in several media by pure or mixed cultures, and compared to abiotic controls. Then bioreactor studies were performed. Redox potential, pH, and metal concentrations were determined.

Conclusions
Between 80 and 100% of the zinc were leached from sphalerite in 9K and DSM 882 medium by mixed cultures from various sources including "Reiche Zeche" as well as by Acidithiobacillus ferrooxidans DSM14882T. Due to a co-precipitation of indium with jarosite, an increase in the solids loading resulted in a lower increase of indium as compared to zinc in solution. The other elements mentioned could successfully be leached from "Theisen sludge".
Background

In marine environments, the activity of microorganisms which form biofilms on metallic materials such as Cu, Fe, Al, and their alloys can result in severe deterioration of these materials, a phenomenon known as microbiologically-influenced corrosion (MIC). It is accepted that biofilms comprising diverse microbial populations are more aggressive towards metals than monoculture biofilms. It has also been proposed that corrosion rates observed in the presence of microbial consortia are related to cooperative metabolic activities of biofilm population, as well as the chemical properties of the colonised surface. As a result of their corrosion resistance, thermal conductivity and reported anti-fouling properties, CuNi alloys find many applications in marine environments and are used extensively in heat-exchangers, pipelines and cladding materials. The stability of these alloys in chloride-containing media has been attributed to the presence of a duplex oxide layer on the alloy surface. However, it has been demonstrated the protective oxide structure on CuNi alloys does not prevent, and merely delay the onset of macrofouling and that these alloys are susceptible to MIC.

Objectives

Advanced electron microscopy, surface science and molecular ecology techniques, such as pyrosequencing have been used to characterise the enrichments and resulting biofilms.

Methods

Continuous flowing bioreactors operating at 10 C and 24 C have been employed to expose CuNi coupons with different surface pre-treatments to aerobic and anoxic microbial enrichments obtained from marine biofilm samples recovered from corroding and non-corroding systems.

Conclusions

The study revealed that there was a correlation between the microbial community structure and the extent of localised corrosion attack.
CAN WE EXPLOIT BACTERIA AND FUNGI TO PRESERVE ARCHAEOLOGICAL IRON OBJECTS?

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Background

Without any conservation-restauration intervention, archaeological iron objects are affected by dramatic corrosion that leads to an irreversible deterioration. The main issue with this metallic substrate is the chlorine content within the corrosion layers that, reacting with H₂O and O₂, causes flacks, cracks and loss of shape of the objects after excavation. Conservation-restauration methods nowadays available are expensive, time consuming and often employ toxic substances.

Objectives

Therefore, the MAIA (Microbe for Archaeological Iron Artworks) project aims to develop alternative conservation-restauration methods for iron artefacts that will be based on the use of microorganisms.

Methods

For this purpose, two metabolic activities of specific microorganisms were studied. First we tested iron reduction/accumulation by anaerobic bacteria and fungi, leading to the formation of stable molecules of low molar volume; and second, the capacity of translocating and/or accumulating chlorine by fungi.

In order to identify some microorganisms with the desired iron metabolism, two anaerobic bacterial strains of Desulfitobacterium hafniensis (LBE and TCE1) and several fungal strains were studied. Different iron sources were tested either as soluble (iron citrate) or solid phases (powdered iron compounds from real objects). Spectrophotometric analyses were carried out to ascertain iron reduction and bacterial growth was followed by qPCR. Analyses by SEM allowed studying the iron and chlorine absorption by fungi, and finally, Fourier Transform Infrared (FTIR) and Raman Spectroscopies were used to identify the bio-minerals produced.

Conclusions

The results show that microbes have the potential to stabilize iron objects by both reduction into more stable minerals and by removing chloride from the object.
USE OF ENDOSPORE-FORMING FIRMICUTES FOR THE BIOREMEDIATION OF TRACE-METALS

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Background
In urban environments, run-off water constitutes a sink for metals as it collects dust particles produced by human activities. In the case of the city of Lausanne (Switzerland), run-off and wastewater is not yet entirely separated. As a consequence, the wastewater treatment plant of Lausanne rejects metal contaminated effluent water into the lake of Geneva, leading to an enrichment of lake sediment by metals like copper, cadmium and zinc. In this environment a link between metal contamination and the relative abundance of endospore-forming Firmicutes was clearly observed. Further investigations tend to show that bacterial exudates, massively produced by these bacteria, play an important role in the chelation and immobilization of metals.

Objectives
The objective of this study is to achieve a better understanding of this relation between endospore-forming Firmicutes and metals in aquatic environments. Is it particular to environments under the influence of wastewater treatment plant rejections or could it be generalized to other polluted environments? The chelation mechanism is also of particular interest.

Methods
To achieve this, sediment cores from different location (two different lakes) were analyzed, looking for bacterial communities composition (454-pyrosequencing), but also physico-chemical characteristics, like metals content (ICP-MS measurement). Cultivation experiments in media amended with different concentrations of metals was also initiated, first for evaluating the chelation potential of the different strains tested (ICP-OES), second for isolating and investigating the chelation substances that were produced (SEM and FTIR).

Conclusions
A better understanding of the metal immobilization processes by bacterial biomass could lead to promising bioremediation technologies.
Motility is an important feature for all life forms. Whereas bacteria use the well studied flagellum for swimming motility, archaebacteria employ the archaellum, which is a rotating type IV pilus like structure. The crenarchaeal archaellum is only composed of seven subunits that are essential for the assembly and rotation of the archaellum. Next to the structural filament protein FlaB, the *S. acidocaldarius* archaellum operon encodes for the monotopic membrane proteins FlaX, FlaF and FlaG, whereas the motor complex is formed by the nucleotide binding protein FlaH, the ATPase FlaI and the polytopic membrane protein FlaJ.

Our current functional and structural analysis of the different archaellum components will be discussed as well as their role in assembly and rotation of the archaellum.
This presentation will describe recent highlights of our research examining polar microbial biodiversity and microbial ecology in extreme polar cryoenvironments with a focus on the Archaeal components. In general, we have detected archaeal sequences (haloarchaea, methanogens, Crenarchaeota, Thaumarchaeota) from 16S surveys and metagenome sequencing projects in most of the habitats tested including Arctic and Antarctic permafrost soils, gypsum and sandstone cryptoendoliths, and saline and hypersaline cold perennial springs in the high Arctic of Canada, albeit at relatively less proportions (10 fold or greater) than Bacteria sequences. The Lost Hammer (LH) spring in the Canadian high Arctic perennially discharges subzero (-5°C) hypersaline (~24% salt) brines through thick layers of permafrost (> 500 m), and so far accounts for the only described terrestrial methane seep in frozen settings on Earth and serves as an analogue for possible Martian liquid water habitats. In the LH source sediments, archaeal phylotypes related to signatures from hypersaline deep-sea methane-seep sediments and were dominated by the anaerobic methane group 1a (ANME-1a) clade of anaerobic methane oxidizing archaea while spring channel sediments was dominated by phylotypes most closely related to ammonia-oxidizing Thaumarchaeota. Viral morphologies (fusiform, spherical) characteristic of archaeal viruses are present in the cold saline springs. Haloarchaea and methanogens are readily found in most permafrost soils and their role in CH4 fluxes from melting Arctic permafrost environments will be presented. Overall, the roles and activities of Archaea in cryoenvironments, especially under ambient subzero conditions, remains very poorly elucidated and is significantly hampered by the lack of culturable archaeal representatives capable of subzero growth.
Background

Archaea, however also found in mainstream habitats, are characterized by several extremophilic species. Extremophilic microorganisms have more robust proteins and unique metabolic strategies to survive extreme conditions. *Sulfolobus acidocaldarius*, an aerobic thermoacidophilic crenarchaeon, is a promising host for biotechnological processes.

Objectives

Getting insight in the mode of action of transcriptional mechanisms could lead to a better understanding of archaeal physiology and to an extended transcriptional toolbox for genetic engineering of *Sulfolobus*. Gene regulation in response to environmental conditions is crucial for the fitness and survival. This project focuses on the characterization of a TetR-like transcription factor, SaFadR.

Methods

A variety of biochemical techniques and *in vivo* experiments has been implemented to study DNA-binding behaviour and physiological role.

Conclusions

By analysing the interactions of the SaFadR protein with genomic regions, SaFadR binds *in vivo* ten genomic regions, with the highest affinity to its own promoter region and the one of a juxtaposed, divergently transcribed operon. Other ChIP-seq binding peaks were observed for an operon encoding enzymes involved in the β-oxidation pathway. This binding is confirmed *in vitro* by performing EMSA. Addition of a fatty acyl-CoA reduces the *in vitro* binding.

Using a SaFadR knockout strain, the relative expression levels of SaFadR and the other possible target genes were compared with an isogenic wild type by qRT-PCR.

Based on these results we can conclude that the SaFadR protein is a fatty acid responsive transcription factor that represses genes involved in fatty acid and lipid metabolism in a fatty acyl-CoA dependent manner.
Background
Members of the novel proposed order “Ferrovales” are the first acidophilic iron oxidising bacteria within the Betaproteobacteria.

Objectives
Since neutrophiles and acidophiles seem to use different strategies to maintain pH homeostasis and to cope with acid stress we were interested in the strategies of the novel acidophile “Ferrovum” sp. JA12.

Methods
Therefore we sequenced and assembled the complete genome sequence of “Ferrovum” sp. JA12, derived from the acid mine drainage of a lignite mining site.

Conclusions
The genome analysis revealed the gene repertoire for both the typical strategies of acidophiles and of neutrophiles. Like other acidophiles “Ferrovum” sp. JA12 may be able to create a reverse membrane potential by the increased uptake of potassium ions facilitated by a number of predicted potassium transporters. In “Ferrovum” sp. JA12 cytoplasmic buffering may be accomplished by amino acid decarboxylases or the production of polyphosphates and spermidine. The presence of a urease encoding gene cluster was unexpected since there are so far no similar gene clusters in other acidophilic iron oxidisers. “Ferrovum” sp. JA12 may use the urease to create a cytoplasmic buffering system similar to the pathogenic neutrophile Helicobacter pylori. Furthermore, the genome contains numerous genes encoding chaperones and DNA repair systems to cope with damages to proteins and DNA caused by low pH. Taken together “Ferrovum” sp. JA12 uses strategies to inhibit uncontrolled influx of protons, to cope with excess protons and to deal with cellular damages in order to maintain life in acidic environments.
Background

Archaea are characterized by a complex metabolism with many unique enzymes that differ from their bacterial and eukaryotic counterparts. *Sulfolobus solfataricus* (optimal growth at 80°C, pH 3) is known for its metabolic versatility and is able to utilize a great variety of different carbon sources (e.g. polymers, pentoses, sugar acids, aldehydes). However, the underlying degradation pathways and their regulation are largely unknown.

Objectives

In the SulfoSYS^BIOTEC^ project we analyse growth on different carbon sources using an integrated Systems Biology approach. Besides pathway elucidation metabolic constraints for life at high temperature due to metabolite instability are of special interest.

Methods

Systems Biology approaches (genomics, transcriptomics, proteomics, metabolomics, bioinformatics, modelling, classical microbiology, biochemistry as well as genetic/ molecular biology techniques).

Conclusions

Using a combined modelling and *in vitro* reconstruction approach it could be shown that in *S. solfataricus* the thermal instability of triosephosphates leads to carbon loss and futile cycling. Like in the modified branched Entner-Doudoroff pathway for D-glucose degradation also in the newly discovered pathway for L-fucose the formation of triosephosphates is omitted ((i) L-fucose oxidation to L-fuconate via a promiscuous dehydrogenase , (ii) dehydration to 2-keto-3-*deoxy*-L-fuconate via dehydratase, (iii) 2-keto-3-*deoxy*-L-fuconate cleavage to pyruvate and L-lactaldehyde via aldotransketolase and finally (iv) L-lactaldehyde conversion to L-lactate via aldehyde dehydrogenase). This pathway shows interesting overlaps to the D-arabinose pathway representing another example for pathway promiscuity in *Sulfolobus* species. Thus, circumventing triosephosphate formation appears to be a special metabolic
thermoadaptation strategy in *Sulfolobus* to face the special challenge of metabolism at high temperature.
PURPLE PHOTOSYNTHETIC REACTION CENTERS FOUND IN THE RARE BACTERIAL PHYLUM GEMMATIMONADETES

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Background

Photosynthetic bacteria represent one of the earliest organisms on Earth. Despite over three billion years of evolution, organisms containing (bacterio)chlorophyll based reaction centers have so far been reported in only six out of over thirty bacterial phyla.

Objectives

In search for novel photosynthetic species we conducted a large screening of microorganisms inhabiting several extreme aquatic habitats.

Methods

One of the surveyed lakes was the Swan lake in the western Gobi Desert, Inner Mongolia, China. The screening was performed using a custom made infra-red screening system.

Conclusions

Using this approach we isolated a novel red pigmented strain AP64 belonging to the poorly characterized bacterial phylum Gemmatimonadetes. The new organism contains fully functional type-2 (pheophytin-quinone) photosynthetic reaction centers composed of bacteriochlorophyll a and carotenoids of spirilloxanthin series. Full genome sequencing revealed the presence of a 42.3-kb long photosynthesis gene cluster (PGC) in its genome. This makes Gemmatimonadetes the seventh bacterial phylum containing chlorophototrophic species (after Cyanobacteria, Proteobacteria, Chlorobi, Chloroflexi, Firmicutes and Acidobacteria). The organization and phylogeny of its photosynthesis genes suggested their ancient acquisition via horizontal transfer from purple phototrophic bacteria. It challenges our understanding of the evolution of photosynthesis providing the first evidence that photosynthetic capacity can be transferred between distant bacterial phyla.

For more info: http://www.pnas.org/content/111/21/7795.abstract
Bacterial persisters are generally non-growing cells that are tolerant to antibiotic treatment. They have been studied mainly in *Escherichia coli* *in vitro* and are reported to form stochastically at very low frequency. All bacterial species form persisters and they are particularly relevant in the context of chronic infections that are recalcitrant to antibiotic treatment. *Salmonella enterica* causes acute and chronic infections by replicating and surviving for long periods of time within host cells, notably macrophages. Previously we described a dual fluorescence reporter method, Fluorescence Dilution, which enables heterogeneity of intracellular bacterial replication to be followed at the single cell level. Using this method we found many viable non-replicating bacteria in several organs in the mouse model of typhoid caused by *Salmonella*. We provide evidence that interaction with the host triggers formation of intracellular persisters that are likely to constitute a reservoir for relapse of infection by *Salmonella* and we decipher the host triggering signals and the bacterial actors involved in this phenomenon.
Type II toxin-antitoxin systems: a reservoir for more sophisticated functions?

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Background

Type II toxin-antitoxin systems (TAs) are composed of two genes that encode a toxic protein and its cognate antitoxin. TAs are surprisingly abundant in bacterial and archaean genomes. The reason of this evolutionary success remains largely unknown. Such ‘toxin-antitoxin-like’ functions are also found in complex secretion systems targeting bacterial or eukaryotic cells and relying on secreted toxins.

Objectives

Our hypothesis is that type II toxic domains are recruited by genetic systems encoding sophisticated functions such as type III, IV and type VI secretion systems as well as bacterial contact-dependent growth inhibition system (CDI and Rhs).

Methods

Using bioinformatics approaches, a series of domain of unknown function (DUF) was detected in bacterial genomes and predicted to be bona fide toxins belonging to TAs and to be associated with CDI and/or Rhs or to be homologous to effectors of type III secretion systems. Functionality of putative toxic domains and antitoxin/immunity is tested in E. coli. Toxin activity and targets are investigated.

Conclusions

Our data show that domains from TAs or CDI/Rhs systems or homologous to type III effectors are toxic for E. coli and that toxicity is rescued by co-expression of the antitoxin/immunity cognate protein. These domains were shown to inhibit translation, although through different means, suggesting that new targets might be possibly involved.

In conclusion, recruitment of toxic domains encoded by TAs by secretion systems might constitute a strong selective pressure. This may provide an explanation why TAs are evolutionary successful and widespread in bacterial genomes.
DISTINCT TYPE I AND TYPE II TOXIN-ANTITOXIN MODULES CONTROL SALMONELLA LIFESTYLE INSIDE EUKARYOTIC CELLS

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Background
Salmonella enterica serovar Typhimurium (S. Typhimurium) is a Gram-negative bacteria which can infect and proliferate within several eukaryotic cells lines. This pathogen can persist in the infected organisms during a prolonged period of time and is responsible for human diseases ranging from gastroenteritis to systemic disease. This species, as most bacteria, possesses Toxin-Antitoxin Systems (TAS) in its genome. TAS are operons composed of two genes: an unstable antitoxin and a stable toxin, which can be activated in response to different stress conditions. The toxins targets are important for cell proliferation or viability. In this way, TAS can regulate bacterial cell growth and can induce biofilm formation, persistence, and probably dormancy during infections.

Objectives
To characterize TAS of S. Typhimurium and measure their role in infection.

Methods
TAS search was done using different data bases. Three different cell lines (two types of fibroblasts and HeLa cells) were selected for infection assays. Intracellular TAS expression was monitored by proteomics and RT-qPCR.

Conclusions
We report a systematic survey and functional analysis of TAS found in S. Typhimurium. Our analyses showed that 18 out of 27 identified TAS of S. Typhimurium are bona fide systems. Although we could detect expression of 10 TAS under eukaryotic cell infection, only 5 systems (3 type I and 2 type II modules) contributed to intracellular survival.
FEMS-0739
Toxin-antitoxin systems in bacteria: clinical implications

FIC DOMAIN TOXINS ARE THE ORIGIN OF INTRA- AND INTER-KINGDOM EFFECTORS OF BARTONELLA
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Background
The diverse and abundant proteins containing FIC (filamentation induced by cAMP) domains typically perform AMPylation, the transfer of an adenosine-5’ monophosphate onto target proteins. Research has so far focused on scattered representatives that secondarily evolved into bacterial virulence factors, but their original functions and evolutionary trajectories have remained elusive.

Objectives
We studied the molecular function and biological role of bacterial FIC domain proteins that are related to host-targeted effector proteins of Bartonella in order to understand their evolution.

Methods
We used protein biochemistry and proteomics to identify the molecular function of bacterial FIC domain proteins. Classical microbiology and advanced agarose gel electrophoresis were used to study the effects of these proteins inside bacterial cells, particularly on DNA topology.

Conclusions
One major group of bacterial FIC domain proteins are toxins of a novel toxin-antitoxin (TA) module that acts via the AMPylation of DNA gyrase and topoisomerase IV, the two bacterial type IIA topoisomerases. AMPylation inhibits all activities of the targets and thus causes a disruption of cellular DNA topology that induces bacterial persistence, a dormant state of multidrug tolerance. One of these FIC domain toxins, VbhT, is encoded next to a type IV secretion system (T4SS) in Bartonella and unique
in that it contains a cognate secretion signal. This protein and the associated T4SS are closely related to host-targeted effectors of this widespread mammal pathogen and their secreting machinery. We therefore propose that VbhT is an interbacterial effector protein and constitutes a “missing link” in the evolution of host-targeted effectors from genuine bacterial proteins.
RESPONSIVE DIVERSIFICATION LEADS TO ANTIBACTERIAL TOLERANCE THROUGH ACTIVATION OF A PORE-FORMING TOXIN

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Background

Within a bacterial population, a small fraction of persister cells is transiently capable of surviving exposure to lethal doses of antibiotics. As a bet-hedging strategy, persistence levels are determined both by stochastic induction and by environmental stimuli, called responsive diversification.

Objectives

Our results support a central role for the conserved GTPase Obg in determining persistence in Escherichia coli in response to nutrient starvation. In this work we unravel the underlying molecular mechanism.

Methods

We used single cell methods to correlate Obg levels with persistence and identified the stringent response as a contributing pathway by using knockout strains. Furthermore, we performed a transcriptome study to identify downstream genetic elements. This resulted in the identification of HokB, a pore-forming toxin and part of a toxin/antitoxin module, as an effector of Obg-mediated persistence. To assess the role of HokB in this phenomenon, we showed membrane depolarization upon obg and hokB overexpression. Furthermore, repolarization of the bacterial membrane by means of a light-driven proton pump diminishes the effect of Obg and HokB on persistence, demonstrating a causal link between the membrane potential and the persistence phenotype.

Conclusions

We propose a model for Obg-mediated persistence that requires the stringent response alarmone (p)ppGpp and proceeds through transcriptional control of the hokB-sokB type I toxin-antitoxin module. In individual cells, increased Obg levels induce HokB expression, which in turn results in a collapse of the membrane
potential, leading to dormancy. Combined, our findings signify an important step towards unravelling genetic mechanisms underlying persistence.
FEMS-1957
Toxin-antitoxin systems in bacteria: clinical implications

MOLECULAR BASES OF THE ALLOSTERIC SWITCH IN CONDITIONAL COOPERATIVITY REGULATION OF TOXIN-ANTITOXIN OPERONS
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Background
Conditional cooperativity is the paradigm of transcription regulation of type II toxin-antitoxin (TA) operons and is intricately related to persistence. Conditional cooperativity allows the toxin component of toxin-antitoxin modules to act as a co-repressor at low toxin:antitoxin ratios and become an activator at high toxin:antitoxin ratios. A common feature in the majority of antitoxins is the presence of an intrinsically disordered region (IDR) typically involved in toxin neutralization and repressor complex formation.

Objectives
To address how the antitoxin IDR is involved in transcription regulation, using the phd/doc operon as model system.

Methods
We determined crystal structures of Phd in complex with its operator box. The DNA-binding domain of Phd interacts with DNA in a novel fashion. Moreover, thermodynamic and structural characterization of Phd in complex with the operator region of the operon in different repression states reveals that a phenotypic switch from negative to positive cooperativity, involving the IDR, is the core of the regulatory process.

Conclusions
In absence of Doc, the intrinsically disordered part of the repressor bound to the DNA acts as a "veil" covering the second site and precluding the binding of a second Phd molecule, resulting in strong negative cooperativity and weak repression. The presence of Doc drives the folding upon binding of the additional Phd molecule with the consequent increase in affinity and strengthening repression, up to the point when the excess of Doc overwhelms the system leading to its de-repression. Our model has important implications in the regulation of bacterial operons involving intrinsically disordered transcription factors.
Biodegradation of contaminants in soil is a logistic problem. ‘Logistics’; refers to have ‘the right thing, at the right place, at the right time’. Degrading bacterial communities walk the tightrope of contaminant availability and suitable physical habitats for their metabolic activity; this situation is often aggravated by the concurrence of restricted bacterial mobility and retarded transfer of hydrophobic organic contaminants (HOC). In order to cope with heterogeneous soil environments mycelial fungi therefore have developed a unique network-based growth form. Unlike bacteria mycelia spread efficiently in the soil, penetrate air-water interfaces and cross over air-filled pores between the bacteria and contaminants in the vadose. In air-filled soil, enhanced homogenization of bacteria and contaminants can be achieved by bridging physical air gaps with fungal hyphae thus enabling substrate-directed mobilization of bacteria along chemical gradients. In this contribution we demonstrate the biodegradation-enhancing influence of mycelial dispersal networks on both the microbial transport and the translocation of HOC. Here we show that mycelial networks (i) act as effective dispersal networks for both undirected and targeted mobilization of contaminant degrading bacteria (‘fungal highways’), (ii) act as hotspots for horizontal gene transfer, (iii) increase the mobility of a wide range of HOC due to their translocation in their cytoplasmic streaming (‘fungal pipelines’), and (iv) improve the accessibility of bacteria to soil contaminants and their subsequent biodegradation. Given their ubiquity and length of up to 1000 m g\(^{-1}\) dry soil mycelia may play a significant role for the ecosystem service of contaminant biodegradation. Knowledge of ‘logistic fungus bacteria interactions’ may also be exploited in biotechnological approaches beyond soil remediation, such as in food industry or the production of biofuels.
Fungi-bacteria interactions: from soil functioning to complex behavior

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Although the role of fungi and bacteria in processes such as decaying or nutrient cycling in soils has been established, the part played by the interactions between these microbial taxa in soil functioning is still poorly known. Myriads of potential interactions can emerge as results of the encounter of fungi and bacteria in a shared environment. These interactions range from positive (e.g. mutualism) to negative (e.g. antagonism). I will present results for several of these types of interactions including fungus-driven bacterial dispersal, bacterial farming by fungi, as well as biocontrol and niche exclusion between bacteria and fungi. For those, we have started to understand the underlying principles that have allowed the emergence and maintenance of these interactions within an evolutionary conceptual frame. In addition, we are investigating the molecular and chemical mechanisms that regulate processes such as partner recognition or metabolic activity. All of these elements are addressed in the context of the importance of fungi-bacteria interactions in ecosystem functioning and services, in particular related to pedogenic carbon storage.

In the past few years, the scientific community has started to recognize the importance of fungi-bacteria interactions in fields as varied as human health (e.g. infection and obesity), agriculture (e.g. plant growth promotion), or the environment (e.g. carbon storage and nutrient cycling). In the future it will be important to compare the mechanisms discovered in the case of soil fungi-bacteria interactions with other ecosystems, in order to understand the universality of the processes that allow these phenomena to emerge in nature.
EFFECT OF DISPERSAL NETWORKS ON BACTERIAL DISPERSAL AND BIODEGRADATION AT VARYING WATER POTENTIALS

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Background

In porous media the matric and the osmotic potential contribute to the availability of water to microbes and decisively influence important microbial ecosystem services such as biodegradation. Bacterial motility is considered as a key driver for biodegradation and fungal mycelia have been shown to serve as effective dispersal networks thereby increasing bacterial movement in water unsaturated environments.

Objectives

However, poor knowledge exists on the beneficial effects of mycelia at varying water potentials (Ψw).

Methods

We therefore established experimental microcosms to investigate the effect of mycelia-like dispersal networks on the dispersal and growth of Pseudomonas putida KT2440-gfp at given osmotic and matric potentials and determined their benefit for the biodegradation of benzoate. Using either NaCl or polyethylene glycol 8000 the Ψw of agar was modified between ΔΨw 0 - -1.5 MPa (i.e. water potentials representing completely saturated or plant permanent wilting point conditions).

Conclusions

We found that dispersal, growth and biodegradation rates dropped noticeably below ΔΨw -0.5 MPa in osmotically stressed systems. However, in matric stress treatments this decline occurred at ΔΨw -0.25 MPa due to a complete repression of bacterial movement at this Ψw. The presence of dispersal networks effectively defused the negative effects of lowered matric potentials by enhancing bacterial dispersal. No beneficial network effect was observed in the osmotically stressed systems, likely due to NaCl toxicity rather than the water deprivation effects. We propose that dispersal networks act as an important buffer mechanism and hence may increase the microbial ecosystem’s functional resistance to matric stress.
STRATEGIES OF STREPTOMCETES FOR FUNGAL TARGETING AND INHIBITION

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Background

Streptomyces belong to actinobacteria, have a complex life cycle, and comprise many species. They have a large linear chromosomal DNA, which has been sequenced from many species. They are highly abundant in different soil-types and the rhizosphere of plants. They can play a role in association with selected organisms, and sometimes as symbionts. The knowledge about proteins and compounds facilitating these interactions is still scarce.

Objectives

The aim of the presented investigations was to investigate the role of newly discovered proteins and vesicles during the interaction of streptomyces with selected fungi.

Methods

Characterization of proteins and metabolites using LC-MS, electron-microscopy, fluorescence microscopy, cloning of genes, and analyses of designed mutants.

Conclusions

Streptomyces produce several unique proteins that target selected fungi provoking close interaction. We purified the proteins and elucidated their mode of action. In addition, we discovered firstly that several proteins and metabolites of streptomyces reside in vesicles, which we characterised by several biochemical and microscopic tools. We succeeded to monitor the in vivo generation of vesicles and to propose a model for their biogenesis. The application of vesicles to selected fungi including a plant pathogen, led to pronounced fungal damages. The investigations revealed the sophisticated Streptomyces repertoire to target and to suppress fungi.
including a plant pathogen.
FEMS-2618
Fungal bacterial interactions

ROLE OF VOLATILES IN ANTIFUNGAL ACTIVITY OF A LACTOBACILLUS PARACASEI AGAINST PENICILLIUM STRAINS
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Background
Fungi constitute a major spoilage problem in many foods including dairy products. The use of specific lactic acid bacteria (LAB) with antifungal properties has shown potential for delaying fungal spoilage without compromising sensory acceptability of the products. Much effort has been put into isolating and identifying antifungal metabolites from active fractions of cell free ferments of these bacteria. However, the methods used have been suboptimal for detection of volatiles.

Objectives
To study the antifungal role of volatiles.

Methods
Inhibitory effect of a Lactobacillus paracasei and metabolites was studied in a newly developed chemically defined medium and yogurt against two spoilage-associated Penicillium strains using a gentle sample treatment.

Conclusions
Diacetyl was identified as the major volatile produced. Antifungal activity was observed in both solid and liquid media as well as in yogurt. If cells were removed from the ferment both diacetyl content and antifungal activity was drastically reduced. If formation of diacetyl was inhibited by enzymatic conversion of the pre-cursor antifungal activity decreased markedly underlining the importance of volatile compounds in the inhibitory activity.
Background
Bacterial motility in water-unsaturated soil is usually highly restricted. Previous experiments in vitro however have demonstrated that bacteria may take the ‘fungal highway’, i.e. are able to actively move along fungal hyphae.

Objectives
In order to get better insights into such fungal-bacterial interactions in situ, we developed a column system designed to collect and cultivate fungi and associated migrator bacteria at varying growth conditions.

Methods
Our column system was placed into a soil known to exhibit high metabolic fungal-bacterial interactions within the oxalate-carbonate pathway. After 4 to 8 days, the columns were removed, bacteria and fungi that grew in the columns identified, and the bacteria analyzed for their ability to use different carbon sources, fix nitrogen and to encode the type-III secretion system (as possible indicator for migrating activity along hyphae).

Conclusions
We identified Fusarium sp. and Chaetomium sp. as fungi, and Achromobacter sp., Acinetobacter calcoaceticus, Ochrobactrum sp., Olivibacter sp., Pseudomonas sp., Stenotrophomonas sp. and Variovorax sp. as associated migrator bacteria. We found that the ability of bacteria to move along their fungal hosts was medium-dependent. Most of the bacteria were able to fix dinitrogen. This points to a possible trophic exchange between bacteria and fungi. Except for one bacterial strain genes encoding type-III secretion systems were not found in the isolated bacteria.

Our findings propose that fungal hyphae are not an inert path taken by motile bacteria; ‘fungal highways’ interactions seem to be mediated by trophic exchanges between fungi and bacteria. Therefore, migrator bacteria selection might be based on their metabolic capabilities.
Environmental studies suggest that a large fraction of the total bacterial diversity has so far resisted cultivation. Many of the uncultivated bacteria belong to large taxonomic groups without cultivated representatives and known functional properties. Considering the importance of bacteria for drug discovery, it has been proposed that these elusive organisms represent a massive untapped resource of novel compounds and biotechnologically useful enzymes. To investigate the biosynthetic potential of microbial dark matter, our laboratory uses metagenomic and single-cell methods focusing on host-associated microbiomes. We provide evidence for the existence of a new environmental candidate phylum, termed "Tectomicrobia", members of which live associated with marine sponges and are metabolically highly versatile. These bacteria exhibit a rich specialized metabolism with bioactive compounds not yet encountered in cultivated microbes. Functional studies revealed unprecedented types of biosynthetic transformations that are of significant biotechnological interest.
High-throughput chemotyping and phenotyping using Pseudomonas aeruginosa deletion libraries

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We are currently profiling two P. aeruginosa single-gene deletion libraries (in the clinically relevant strains, PA01 and PA14) in a large number (>200) of chemical and environmental stresses. In these assays, we obtain quantitative measurements both for the fitness, the ability of each mutant to form biofilms and their secondary metabolite profile.

The quantitative phenotypic and chemotypic signature for each mutant will be later used to a) identify novel gene function; b) map metabolic pathways c) map interconnections between functional processes; d) address the role of horizontal gene transfer in network rewiring and gene repurposing; d) understand how chemical perturbations effect the secondary metabolite profile; e) identify novel secondary metabolites; etc.

At the workshop I will present the basis of our methodology, a progress report on the ongoing screen and proof-of-principle examples highlighting the different types of information we are obtaining. Another aspect of the talk will be how to handle and mine such a diverse data set (<16k MS&MS2 profiles, <12800k independent phenotypic data points).
Background

The increase in antibiotic resistance among clinical microbes has created a need to develop antimicrobials with novel modes of action.

Objectives

As part of a marine biodiscovery programme, sponge-associated microbes were screened to identify novel metabolites.

Methods

Metabolomics and genomics are two complementary platforms for analysing an organism as they provide information on the phenotype and genotype, respectively. These two techniques were applied in the dereplication and identification of bioactive compounds from a *Streptomyces* sp. (SM8) isolated from the sponge *Haliclona simulans* from Irish waters. Three distinct bioactives that include an antibacterial fatty acid derivative, and separate antifungal and anti-calcineurin activities were identified. Antimycin compounds were identified in the antifungal fractions. Genomic analysis revealed the presence of multiple secondary metabolism gene clusters, including a cluster for the antimycin family of compounds. When the extract of the mutant strain was analysed, it showed loss of antimycin production and significantly less antifungal activity than the wild-type strain. Butenolides were identified as compounds showing anti-calcineurin activity.

Conclusions

The combined metabolomic and genomic approach used in this study resulted in the identification of antimycins, polyhydroxylated saturated fatty acids and butenolides as
major antifungal and antibacterial and anti-calcineurin metabolites of SM8, respectively. The genome sequence provides a genetic framework to study their biosynthesis and their potential role in regulation of gene expression.

References:

MOLECULAR GENETICS OF SIMOCYCLINONES BIOSYNTHESIS: SURPRISING ASSEMBLY OF THE TETRAENE CHAIN

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Background

Simocyclinone D8 and its’ derivatives were isolated from Streptomyces antibioticus Tü6040. They exhibit antibiotic activity against Gram-positive and Gram-negative bacteria, and are highly potent inhibitors of DNA gyrase supercoiling with an IC₅₀ lower than that of novobiocin¹,².

Objectives

However, biosynthetic studies and bioengineering of simocyclinones were limited because of difficulties in genetic manipulations of the producer strain. We have discovered the new strain Kitasatospora sp. producing novel simocyclinone.

Methods

Kitasatospora sp. genome was sequenced and detailed smc cluster annotation has been performed as well as its alignment to published sim gene cluster from S. antibioticus Tü6040 ³. Inactivation and heterologous expression of several biosynthetic genes have been performed to prove their function.

Conclusions

Several significant differences in the architecture of the new smc gene cluster have been found in comparison to the previously published sim gene cluster from S. antibioticus Tü6040. Most striking, genes responsible for the tetraene chain formation (type I PKS in sim cluster) could not be detected. Instead, three genes with the high similarity to ACP and two beta-ketoacyl-synthases were annotated within the smc cluster (ORF2, ORF3 and ORF4, respectively). ∆ORF4 mutant was deficient in production of the D-type simocyclinones, only angucyclines with the attached sugar could be detected. Subsequent complementation of the mutant with the intact ORF4 restored biosynthesis of simocyclinones. Furthermore, inactivation of the KS domain in type I PKS from sim cluster in S. antibioticus Tü6040 didn’t influence biosynthesis of the simocyclinone D8. Obtained data suggest new unusual pathway for linear polyketide assembly during simocyclinone biosynthesis.
HETEROLOGOUS PRODUCTION OF ANTIBACTERIAL SECONDARY METABOLITES FROM Serratia marcescens IN Pseudomonas putida


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Background

Microorganisms produce a huge variety of secondary metabolites that often exhibit several valuable bioactivities such as antibiosis. This wealth is often accessed by heterologous biosynthesis in a host microbe. Several challenges are typically encountered here, concerning i) the establishment of the genetic information underlying a biosynthetic pathway, ii) the complex biosynthesis and iii) the host’s tolerance towards antibiotic products.

Objectives

Pseudomonas putida is an outstanding host candidate for the heterologous production of substances with antimicrobial activity. The strain P. putida KT2440 is a well-established “safety strain” which is amenable to genetic modification, possesses a versatile metabolic background and withstands many antimicrobial compounds with high tolerance.

Here, we aimed to produce the secondary metabolites prodigiosin and serrawettin W1 from Serratia marcescens. Besides other highly valuable properties such as anticancer activity, both the tripyrrolic red pigment prodigiosin and the lipopeptide serrawettin W1 exhibit antibacterial activity, rendering their heterologous production challenging.

Methods

For heterologous production of prodigiosin, the entire prodigiosin gene cluster was inserted into the chromosome of P. putida. For the production of serrawettin W1, a single synthase gene was expressed from a plasmid in the host. Products were isolated and antimicrobial activities were verified.
Conclusions

We successfully established production of the two antibacterial natural products prodigiosin and serrawettin W1 in the bacterial host *P. putida*, thereby enabling further studies on their bioactivities. This study further highlights that *P. putida* is highly suited for the expression of complex secondary metabolite pathways and the efficient production of antimicrobial compounds.
TILIVALLINE, SECONDARY METABOLITE OF INTESTINAL PATHOBIONT KLEBSIELLA OXYTOCA INHIBITS EPITHELIAL CELL PROLIFERATION

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Background
Klebsiella oxytoca is a natural resident of the human gut microbiota. During antibiotic-induced intestinal dysbiosis colonic overgrowth of this bacterium rapidly leads to antibiotic-associated hemorrhagic colitis (AAHC). We have shown that bacterial production of the secondary metabolite tilivalline, which is structurally related to pyrrolobenzodiazepines, induces apoptosis of epithelial cells, disrupts their barrier function, and is key to the pathogenesis of colitis in an animal model of AAHC (1). The impact of tilivalline on host physiology in healthy carriers of low (commensal) levels of Klebsiella oxytoca is unknown.

Objectives
We synthesized tilivalline chemically to investigate the human cellular processes altered by this enterotoxin in vitro.

Methods
HeLa cells were treated with a range of tilivalline concentrations and cellular stress responses were monitored on the transcriptional level using qRT-PCR, the protein level via Western analysis, through microscopic imaging and FACS as well as by biochemical analysis.

Conclusions
Apoptosis is induced by high doses of tilivalline. Low concentrations of the cytotoxin inhibit HeLa cell proliferation. The position of cell cycle arrest was defined via quantification of cyclin transcripts, DNA damage response markers, modulation of checkpoint pathways, analysis of spindle apparatus formation and centrosome dynamics.

Rapid antimicrobial susceptibility testing: will it be genotypic or phenotypic?

Clinicians are used to get sequential results from the microbiology lab. For blood cultures, it is classical to report as soon as possible the presence of growth and the morphology of the bacteria that are observed under the microscope (e.g. Gram-negative rods), then an identification (e.g. E. coli) and finally the antimicrobial susceptibility profile, AST (e.g. amoxillin R, co-amoxicillin S, etc...). The clinician learns therefore several clinically-relevant elements, on a sequential manner, typically over a few days: i) there is bacterial growth in this sample, ii) the identification of the bacteria may suggest a particular anatomic source (e.g. likely from urinary or digestive origin), and finally, iii) the AST permits reducing the antibiotic spectrum.

MALDI-TOF MS has allowed reducing the time for bacterial identification, but this is typically insufficient to permit targeted antimicrobial therapy. There is therefore a major incentive in knowing as soon as possible how to better target antimicrobial therapy.

Sequencing (and especially with the availability of bench-top NGS systems) can provide an extensive listing of DNA sequences that are present in a sample. Irrespective of the numerous technical challenges (sample preparation, DNA extraction/purification, DNA sequencing, data analysis with or without assembly, etc...), sequence interpretation remains a major hurdle, and can also fail to capture bacterial properties affecting the AST (e.g. the activity of an extrusion pump, ...). Clinically-relevant diagnostic devices may therefore need -at least in the initial phase- to be coupled with some advanced phenotypic information of diverse nature (peptides, mRNA, enzymatic activity, etc...).

This strategy and possible implementation will be discussed during the lecture.
Whole genome sequencing as a tool for clinical diagnostics

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The introduction of Whole Genome Sequencing (WGS) through the Next Generation Sequencing (NGS) technology has great potential as a typing as well as a diagnostic tool in the clinical laboratory. In principal, WGS data contain all relevant information for typing and diagnostic analysis. Existing gene based (local) typing and diagnostic data are often easily extracted from WGS data and many studies have shown the potential of WGS data for (global) analysis of clonal relationship in outbreaks. However, a prerequisite for the transition from current typing and diagnostic methods to methods based on WGS data is to identify the added value of this transition such as increased speed of analysis, decreased costs as well as cheap (free) and user-friendly harmonised WGS-based tools. Also, it will to some extend require the establishment of backward compatibility with existing typing methods, which are currently well-established tools for both clinical and epidemiological investigations. In addition to this, great efforts in relation to standardization and validation of WGS tools as well as harmonization of methods will be a requirement for a successful transition. Finally, WGS has the potential to detect and possibly type pathogens directly in complex samples (the so-called metagenomics), rather than analysing and comparing single isolates. The benefits from this includes decreased time for diagnostics and the ability to detect pathogens, which are not easily detected by standard laboratory methods. However, tools and methods to analyse metagenome data is still in their infancy and will require further maturation before implementation directly in a clinical setting.
FEMS-0870
New diagnostic approaches

AN INTEGRATED ANALYSIS WITH DEEP-SEQUENCING OF BACTERIAL INFECTION AND PROTEOMIC PROFILING OF NEUTROPHILS IMPROVES PREDICTION OF CLINICAL OUTCOMES IN SEPSIS
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Background
Sepsis is a life-threatening complication caused by a serious microbial infection. Due to the complexity of the sepsis response, the prognosis of sepsis is poor and outcomes in individual patients are difficult to predict.

Objectives
This study aims to define molecular signatures of bacterial infection at various stages of sepsis progression and associate them with the clinical features and outcomes of sepsis patients.

Methods
Two thousand three hundred forty-three patients who underwent elective general surgery were recruited prospectively at Jinling Hospital in China from June 2013 to September 2014. A set of 180 patients had six groups that reflected conventional concepts of sepsis progression as a pyramid. Molecular patterns of bacterial infection were elucidated by 16S rDNA-based high-throughput sequencing techniques. We also performed a comparative proteomic profiling on neutrophils derived from peripheral blood of the patients.

Conclusions
The bacterial profiles of patients who were diagnosed as sepsis and ultimately died within 28 days differed markedly from those of survivors with sepsis. The different taxa of bacterial profiling clustered into the following genus: Escherichia/Shigella, Stenotrophomonas, Acinetobacter, Pseudomonas, Corynebacterium and Propionibacterium. A total of 97 proteins were characterized as differentially expressed between nonsurvivors and sepsis survivals, indicating impaired functions of neutrophils in lysosome, chemotaxis, transmigration, phagocytosis and intracellular killing of microorganisms. Our data demonstrate that an in-depth analysis of peripheral blood bacteria with measurements of six protein molecules on neutrophils
would predict survival of sepsis patient. This algorithm may help to guide the
treatment of individual patients with sepsis.
Background
Nine million cases of campylobacteriosis, due to handling and/or consumption of undercooked poultry meat contaminated with *Campylobacter*, occur each year in Europe, with a cost to public health systems of about EUR 2.4 billions. The direct plating and/or enrichment culture methodologies based on the ISO 10272-1B: 2006 are the methods typically used, but the long time required by the classical methods to grow microorganisms, can lead to the distribution of contaminated food. Molecular biology with PCR, qPCR and blotting techniques have greatly improved the techniques by reducing the time required to obtain results, but integrated biochips are the ideal solution for producing portable diagnostic systems.

Objectives
The aim of the work was the development of a miniaturised, highly-sensitive DNA biochip based on a deep-blue organic light-emitting diode (OLED) to analyze real poultry samples.

Methods
Two specific DNA probes (capture and detection) were designed and used to analyze real poultry meat samples. The capture probe was bond to a glass slide while the detection probe was labelled with a Alexa Fluor fluorophore to be detected by the utilization of the OLED.

Conclusions
The results obtained by the biochip were compared with PCR, broth enrichment and colony growth methods, to demonstrate the relevance of this system. The OLED biochip reached a sensitivity of 0.37 ng/μL DNA, which was approximately 20-fold higher than the sensitivity obtained with dot blot assay (25 ng/μL) and didn't need an enrichment step. This system exploits a multi-disciplinary approach, and shows high sensitivity, quick test results and ease of use.
MOLECULAR TYPING OF TREPONEMA PALLIDUM ISOLATED FROM PATIENTES ATTENDING THE UNIVERSITY HOSPITAL OF BUENOS AIRES, ARGENTINA

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Background
Treponema pallidum subsp. pallidum, the causative agent of syphilis, is a sexually-transmitted disease with worldwide occurrence. Molecular typing of T. pallidum provides valuable epidemiological information, including determination of macrolide resistance.

Objectives
This study was aimed to type, for the first time in South America, T. pallidum isolates in clinical samples

Methods
During 2013, 79 samples were collected from patients with clinical signs of syphilis: 72 lesion swabs, 3 whole blood samples and 4 cerebrospinal fluid. DNA was extracted and PCR amplification of polA(TP0105), tmpC(TP0319), TP0136, TP0548, and 23S rRNA was performed(1). A CDC typing scheme was made including the determination of 60-bp repeats in the arp gene, RFLP of tprE-G-J and sequence analysis of TP0548(2). The human MTHFR gene was amplified to test for PCR inhibitors.

Conclusions
Treponemal DNA was detected in 57.3%(43/75) of samples (four were excluded). A total of 36 samples (83.7%) were typed. Nine genotypes were detected using the sequencing-based typing, with SSS being the most frequent. CDC typing scheme
was done in 22 samples, and 7 genotypes were detected, with 14d/d being the most common. In 13.9% of the samples, mutations causing macrolide resistance were detected. In conclusion, during this first molecular typing study, low frequency of macrolide resistant mutations was detected compared to Europe and the US. Although some detected treponemal genotypes were identical to those described in other countries, several genotypes were new, not found outside Argentina.

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FEMS-2303
New diagnostic approaches

RAPID AND SENSITIVE ISOTHERMAL DNA AMPLIFICATION ON A DIGITAL MICROFLUIDIC DEVICE FOR THE DETECTION OF ANTIBIOTIC RESISTANT BACTERIA
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Background
Antibiotic resistance is a growing global threat to human health. There is a clear need to develop rapid and sensitive tests to determine antibiotic resistance as part of diagnosis so that antibiotics can be prescribed to treat infections in a targeted manner. The most rapid, sensitive and specific molecular assays are based on nucleic acid amplification to detect specific genes of interest. Combining this with droplet microfluidic platforms represents the most promising approach for portable point-of-care assays.

Objectives
To develop a miniaturised assay to detect the gene encoding for a CTX-M enzyme, an extended spectrum β-lactamase present in common bacteria such as *E. coli*.

Methods
We perform real time isothermal DNA amplification by recombinase polymerase amplification (RPA) on a digital microfluidic device, which utilises active matrix electrowetting on dielectric (AM-EWOD) technology.

Conclusions
The assay can detect just 10 copies of the *E. coli* CTX-M gene within 15 minutes. Continuous mixing during the assay run on AM-EWOD enhances the sensitivity of the assay by 10-fold relative to a commercial microplate reader. The assay is quantitative, with the time of onset of amplification relating to the logarithm of DNA copies in a linear manner. DNA from *E.coli* containing the CTX-M gene can be detected even when it is mixed 1:100 with genomic DNA from *E. coli* without CTX-M. Our systems ability to rapidly detect antimicrobial resistance genes represents a promising step to a complete and fully automated, sample in – answer out, Point of Care diagnostic assay suitable for directing antibiotic prescribing.
Uwe T. Bornscheuer, Universität Greifswald

Protein engineering has developed in the past decade to a highly important technology (1,2) to create enzymes with desired properties, but also to understand their function. Whereas initially rational protein design based on detailed analysis of structures was the method of choice, directed evolution became an important alternative. In this lecture, the principle strategies and current challenges in protein engineering will be highlighted. We used information from protein databases to create 'small, but smart' focused protein libraries of enzymes from the a/b-hydrolase fold family using the 3DM database (3). This resulted in enzyme variants with enhanced thermostability or enantioselectivity (4). For the synthesis of chiral amines, we developed an in silico analysis and identified a toolbox of novel (R)-selective transaminases (5) as well as (S)-selective enzymes from a structure-guided search (6). More recently, we focused on cascade reactions to convert unsaturated fatty acids to w-hydrocarboxylic acids (7), or to combine alcohol dehydrogenase, enoate reductase and monooxygenase for in vivo biotransformations (8). In another project, we use a chalcone isomerase in the conversion of flavonoids (9). Very recently, we developed a synthesis of e-caprolactone-oligomers through an enzyme cascade (10).


Discovery and engineering of enzyme systems for the transformation and degradation of explosives

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Background. The contamination of the environment with toxic explosives presents a serious and widespread problem at sites across the world. Our recent studies on the biodegradation of energetic compounds have resulted in the isolation of bacteria from explosive-contaminated land that are capable of metabolizing 2,4,6-trinitrotoluene (TNT) and RDX. Our work has targeted the use of genetic engineering for broadening and improving the degradative capabilities of biological systems for the remediation of explosives pollution. Significant progress has been made towards this goal and we have successfully combined the biodegradative capabilities of soil bacteria with the high biomass and stability inherent to plants. In an effort to determine how plant tolerance could be further improved, we have been investigating the enzymology underlying the innate ability of plants to detoxify explosives.

Objectives. 1. To determine the biochemistry and molecular genetics of explosives metabolism in microorganisms and plants. 2. To characterise the enzymes mediating these degradative pathways. 3. To engineer transgenic plant systems for the remediation of explosives contaminated land.

Methods. The work has required a plethora of modern molecular techniques and spans methods in microbial and plant biology through to methods in synthetic biology and enzyme engineering.

Conclusions. These studies have identified a number of interesting enzymes including a unique cytochrome P450 XplA from a strain of Rhodococcus rhodochrous that degrades RDX. We have successfully engineered transgenic plants that can efficiently remove toxic levels of TNT and RDX from contaminated soil and water. As a result of recent advances in our knowledge of the biodegradation of explosives, endogenous plant detoxification systems and the use of genetic engineering, our work on transgenic plants is now poised for testing in the field.
FEMS-1860
White biotechnology

FERMENTATION OF CRUDE SYNGAS FROM THE BIOLIQ® PLANT KARLSRUHE
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Background

The new bioliq® pilot plant at the KIT covers the complete process chain required for producing customized fuels from dry lignocellulosic biomass. For energy densification of the biomass, fast pyrolysis is applied. The liquid pyrolysis oil and solid char obtained can be processed further in the entrained flow-gasifier to tar-free, low-methane raw synthesis gas. Prior to chemically catalysed fuel synthesis a multistep cleaning of raw synthesis gases is performed: Particles, alkaline salts, HCl, H₂S, COS, CS₂, NH₃, and HCN are removed to avoid catalyst poisoning during fuel synthesis. The pilot plant is equipped with an innovative hot-gas cleaning system.

Objectives

Acetogenic bacteria are able to ferment syngas to a variety of organic acids and alcohols. In contrast to the catalysts used in the Fischer-Tropsch process, these biological catalysts can process a broad range of syngas compositions and deal with impurities like sulphur compounds or CO₂. At the moment intense efforts are made to genetically modify C. ljungdahlii to optimize product yields and establish synthesis routes to new products. To assess industrial large scale applicability of these strains it will be necessary to determine their performances with crude syngas, as each purifying step will decrease the economy of the process.

Methods

Our aim is to establish a platform process for evaluating the use of different qualities of raw syngas obtained from the different steps of Karlsruhe bioliq® plant.

Conclusions

Therefore, a setup of multiple 2 L bioreactors with online gas and product analytic is currently developed in our lab.
HIGH LEVEL PRODUCTION OF ETHYLMALONYL-COA PATHWAY-DERIVED DICARBOXYLIC ACIDS BY METHYLOBACTERIUM EXTORQUENS UNDER COBALT-DEFICIENT CONDITIONS AND BY POLYHYDROXYBUTYRATE NEGATIVE STRAINS

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Background

There is a high need for sustainably producible chemical building blocks being applicable e.g. as monomers for novel bioplastics. The ethylmalonyl-CoA pathway (EMCP) harbors several CoA-esters such as ethylmalonyl-, methylsuccinyl- or mesaconyl-CoA whose free dicarboxylic acid derivatives potentially present promising synthons for chemical industry. The EMCP in Methylobacterium extorquens offers the possibility to produce these new building blocks directly from the cheap and non-food competing C-source methanol.

Objectives

Main objective of this work was the optimization of dicarboxylic acid production with M. extorquens strains expressing a thioesterase towards an industrially relevant process.

Methods

Suitable thioesterase enzymes were identified by in vitro assays and the product spectra were analyzed by in vivo experiments. Expression of thioesterases in different M. extorquens strains was increased by optimization of ribosome binding sites. To increase carbon flux into product and avoid acid reuptake we used metabolic engineering and medium optimization strategies. First high cell density fermentations were performed in a bioreactor.

Conclusions

Suitable thioesterases for production of different EMCP-derived dicarboxylic acids were identified and their expression levels in M. extorquens were optimized. Productivities of different strains were compared and increased by different metabolic engineering and medium optimization strategies. In summary, we present the biotechnological production of unconventional dicarboxylic acids from methanol with high efficiency.
PLANT SECONDARY METABOLITE PRODUCTION IN SACCHAROMYCES CEREVISIAE
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Background

The diversity of metabolites of plants with an estimated 200,000 different compounds is unrivaled by any other group of organisms. Many of these metabolites have special features, ranging from scents to antimicrobial agents. The limited availability of secondary metabolites often hinders their exploitation. This can be overcome by implementing synthesis pathways into the yeast \textit{Saccharomyces cerevisiae} that has an excellent track record in white biotechnology.

Objectives

We constructed and characterized \textit{S. cerevisiae} microbial cell factories producing triterpenoids.

Methods

Advanced metabolic engineering including compartment engineering was employed to redistribute carbon flux through the mevalonate pathway and squalene synthesis towards triterpenoids. The product performance of the engineered strains was investigated by bioprocess engineering, while advanced analytics delivered targets for the next round of strain improvement.

Conclusions

\textit{S. cerevisiae} was engineered for triterpenoid production. In fermentations considerable titers of these valuable secondary plant metabolites could be produced. The results will be presented in the context of the required molecular tools and novel engineering targets.
Background

In nature microbes are equipped with enzymes and helper proteins to modify and degrade naturally occurring polymers including polyesters. Due to huge environmental problems, the development of efficient polyester recycling strategies and improvement of biodegradable polyesters that show required material properties is necessary. Learning strategies from microbes enables us to fight against these ecological problems and process polymers efficiently.

Objectives

Here, strategies based on novel microbial enzymes for tailor-made polyester modification and degradation are presented. Mimicking nature, enzymes have been genetically equipped with polymer binding modules [1]. Moreover, mechanistic insights into the enzymatic hydrolysis of synthetic polyesters like polyethylene terephthalate and the compostable synthetic polyester poly(butylene adipate-co-butylene terephthalate) by cutinases and novel “anaerobic” esterases are given.

Methods

The enzymatic hydrolysis of polyesters by cutinases e.g. from *Thermobifida cellulosilytica* (Thc_Cut1) and novel anaerobic esterases from different *Clostridium spp.* was investigated via HPLC-MS. Additionally, fusion proteins of Thc_Cut1 and binding modules of polyhydroxyalkanoate depolymerases and combination with hydrophobins from *Trichoderma spp.* [2] were investigated.

Conclusions

Novel enzymes isolated from anaerobic microorganisms were demonstrated to degrade synthetic polyesters. The adsorption as well as the activity of microbial
polyesterases was increased when fused to different binding modules or when hydrophobins were applied when compared to the native enzyme.

References


FEMS-2838
Microbial symbioses with marine animals

How to eat without a mouth or gut: symbioses between chemosynthetic bacteria and gutless marine worms

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Marine worms without a mouth or gut were first discovered over 35 years ago in coral reef sediments, but the mystery of how they gain their nutrition was not solved until large communities of gutless invertebrates were discovered in 1977 at hydrothermal vents in the deep sea. We now know that these animals gain their nutrition from chemosynthetic bacteria that fix carbon dioxide into organic compounds, as in photosynthesis, but using reduced compounds such as sulfide or methane as energy sources instead of sunlight. Chemosynthetic symbioses occur worldwide and have evolved multiple times from numerous bacterial lineages and at least 9 animal groups (Dubilier et al. 2008. Nature Rev Microbiol). In my talk, I will present an overview of the research in my lab on these symbioses and describe how new tools that range from in situ instruments for measuring environmental parameters to lab-based methods such as ‘omics’ and single cell imaging, provide the opportunity to understand symbiotic associations in their environmental and ecological context.
The lactic acid bacterium *Lactococcus lactis* is of high importance in biotechnology, specifically in the dairy industry. It is used in starter cultures world-wide for the production of safe and nutritious milk fermentation products. Rapid acidification of milk through lactose utilization and proteolytic activity are key parameters in determining product quality. The large-scale industrial processes in which these bacteria are employed put them under a variety of intrinsic and extrinsic (technological) stresses. Consequently, stress response systems in *L. lactis* have been extensively studied, as have industrially important global transcriptional regulators of carbon and nitrogen metabolism.

It is becoming increasingly clear that, in addition to protein regulators, small non-coding RNAs play important regulatory roles in bacteria. Recent developments in next-generation RNA sequencing and in bioinformatics have revolutionized transcriptome analyses. As a result, novel non-coding RNAs have been discovered in many micro-organisms and subsequent studies have uncovered several regulatory mechanisms that operate at the level of RNA.

The presence and roles of non-coding RNAs with regulatory functions have not yet been assessed in *L. lactis*. Because of their presumed importance, we set out to identify novel RNA elements, such as small regulatory RNAs, antisense RNAs and sORFs, by growing the organism under various (stressful) conditions. Using differential RNA sequencing, a transcriptome landscape of *L. lactis* was obtained that was automatically mined using TSSer and verified and refined manually. Thus, we identified hundreds of novel RNA species. Experimental validation was performed for a selection of these potential RNA regulators. The role of one non-coding RNA located in the 3'UTR of the gene for the regulator of *e.g.* the *arc*-operon in *L. lactis*, ArgR, will be further detailed.
Understanding the mechanisms of yeast adaptation to wine fermentations

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The wine fermentation is a complex process produced as a result of the activities of a succession of microorganisms. Although *S. cerevisiae* is the most frequent species in wines, and the subject of most studies, *S. bayanus var. uvarum* and natural hybrids between *Saccharomyces* species such as *S. cerevisiae* x *S. kudriavzevii* and *S. cerevisiae* x *S. bayanus var. uvarum* are also involved in wine fermentations and can be preponderant in certain wine regions. Studies performed in our laboratory, showed that strains of non-conventional *Saccharomyces* species, such as *S. bayanus var. uvarum* and *S. kudriavzevii*, exhibit physiological properties of potential interest in enology because they can respond to the new demands of the wine industry, such as their ability to ferment at low temperatures, their increased production of glycerol, their lower ethanol yield (1, 2, 3). The main objective of this work is to decipher the molecular mechanisms responsible of the overproduction of glycerol in *S. kudriavzevii*, *S. bayanus var. uvarum* and natural strains of *S. cerevisiae* (4, 5), as well as the molecular basis involved in the cold adaptation of *S. kudriavzevii* strains (6).

References.


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**Background**

The RNA chaperone Hfq has been recognized as the principle post-transcriptional regulator of carbon catabolite repression (CCR) in the opportunistic human pathogen *P. aeruginosa* (PAO1). Hfq was shown to act as a translational repressor that prevents ribosome loading through binding to A-rich sequences within the ribosome binding site of mRNAs encoding catabolic enzymes. Furthermore, the non-coding CrcZ RNA was shown to bind to and to sequester Hfq, which in turn abrogates Hfq-mediated translational repression.

**Objectives**

To study the role of *P. aeruginosa* Hfq and of the regulatory RNA CrcZ in anoxic biofilm formation and antibiotic susceptibility.

**Methods**

Biochemical and genetic methods as well as next generation sequencing were employed to address the impact of Hfq and CrcZ on anoxic biofilm formation and antibiotic susceptibility of *P. aeruginosa*.

**Conclusions**

Comparative transcriptome studies identified CrcZ as being up-regulated in anoxic biofilms of PA14 grown under conditions that mimic growth in the cystic fibrosis lung. We show that CrcZ up-regulation suppresses biofilm formation, which most likely results from sequestration of Hfq. Moreover, RNA-seq analyses revealed functions that account for decreased biofilm formation upon CrcZ up-regulation. In addition, we show that CrcZ affects the susceptibility of PA14 towards several antibiotics, which in turn can explain the differential susceptibility of *P. aeruginosa* to certain antibiotics during planktonic and biofilm growth, respectively.
Small regulatory RNAs in type I toxin-antitoxin systems

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Toxin-antitoxin (TA) systems are pairs of adjacent genes in which a stable toxic peptide is neutralized by an unstable antitoxin. A broad variety of type I TA systems are known in which the antitoxin is a small RNA that is convergently transcribed to the toxin mRNA, and inhibition occurs by RNA/RNA binding. The antitoxin either inhibits translation or facilitates degradation of the toxin mRNA. We investigate two type I systems from the B. subtilis chromosome, bsrG/SR4 and bsrE/SR5. The toxins are small hydrophobic peptides, and toxin and antitoxin genes overlap at their 3' ends by ≈120 nt. Whereas BsrG (38 aa) causes cell lysis in the absence of its antitoxin SR4, BsrE (30 aa) is only toxic upon overexpression. The antitoxins SR4 and SR5 promote toxin mRNA degradation by recruiting RNase III. In contrast to SR5, SR4 has a second function: it induces structural changes around the bsrG SD sequence, thus further obstructing ribosome binding which results in translational inhibition.

Secondary structures of toxin RNA, antitoxin and their complexes were mapped, apparent binding rate constants ($K_{app}$) of wild-type and mutated toxin/antitoxin pairs determined, and the results corroborated by in vivo experiments. The binding pathways of toxin mRNA and antitoxin were elucidated by employing time-course experiments. In both cases, binding initiates between the antitoxin terminator-loop and loop 4 of toxin mRNA.

RNases involved in the degradation of toxin mRNAs and antitoxins were identified. RNase R plays the major role in bsrG/SR4, whereas RNase J1 is a crucial player in bsrE/SR5. Antitoxin-induced RNase III cleavage sites in bsrG and bsrE mRNAs were mapped.

Whereas bsrG/SR4 is the first temperature-sensitive TA system, factors affecting bsrE/SR4 expression have still to be identified.
FEMS-2697  
Small RNAs and regulation: basic aspects Offered by FEMS Microbiology Reviews

New players in the field of small regulatory RNAs

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Bacterial small RNAs (sRNAs) are crucial regulators of cellular functions by modulating gene expression in response to various environmental changes. Many bacterial species were shown to express hundreds of sRNAs. These sRNAs regulate target mRNAs by direct base-pairing to positively or negatively affect their translation and stability. In *E. coli* and other Enterobacteriaceae, the sRNA RyhB regulates iron homeostasis. Under iron-rich conditions, the transcriptional repressor Fur blocks *ryhB* transcription. During conditions of iron starvation however, Fur becomes inactive and relieves repression of *ryhB*. Under these conditions, RyhB directly regulates approximately 20 different mRNAs encoding iron-using proteins. By binding to those mRNAs, RyhB shuts down translation and stimulates their rapid degradation through the action of the endoribonuclease E (RNase E). RyhB contributes actively to increase the levels of free intracellular Fe²⁺ (iron sparing) under conditions of iron starvation by reducing the expression of iron-using proteins and by stimulating siderophore synthesis. Surprisingly, although we have used microarrays to successfully determine the effect of RyhB on the transcriptome of *E. coli*, this approach was insufficient to determine the regulation of undetected but important mRNAs. Thus sRNAs can significantly modulate many target mRNAs without interfering with their levels, which unfortunately prevents target detection by classical techniques. To address this challenge, we developed a method based on affinity purification and RNA sequencing, which could identify any RNA molecule interacting with sRNAs, regardless of the regulation. Thus, we used sRNAs tagged with the bacteriophage MS2 RNA stemloops, which are bound with high specificity by the MS2 coat protein. We then analysed all bound target RNAs by RNA sequencing. We will present new targets that suggest an unexpected but important mechanism for bacterial cells.
FEMS-2381
Small RNAs and regulation: basic aspects Offered by FEMS Microbiology Reviews

Regulatory RNAs in the pathogenic Epsilonproteobacteria, Helicobacter pylori and Campylobacter jejuni
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Genome sequencing revealed the potential proteins and a high genetic diversity of Helicobacter pylori, the causative agent of gastritis and gastric cancer, and the related, emerging food-borne pathogen Campylobacter jejuni. So far, little is known about post-transcriptional regulation in these pathogenic Epsilonproteobacteria. However, based on a differential RNA-sequencing approach (dRNA-seq), we had defined a genome-wide map of transcriptional start sites (TSS) and identified >60 small RNAs (sRNAs) in H. pylori. To understand how transcriptome differences could contribute to phenotypic differences among strains, we applied a comparative dRNA-seq approach to multiple C. jejuni strains. Our study revealed that the majority of TSS is conserved among strains, but we also observed strain-specific promoter usage and sRNA repertoires, which could contribute to strain-specific gene regulation.

Based on our transcriptome datasets, we are now using Helicobacter and Campylobacter as new model organisms for riboregulation in virulent bacteria and bacteria without the RNA chaperone Hfq. We are functionally characterizing abundant sRNAs and are especially interested in their roles and underlying molecular mechanisms in stress response and virulence control. For example, we could show that the highly abundant and conserved sRNA, RepG, from Helicobacter directly base-pairs with a homopolymeric G-repeat in the mRNA leader of the chemotaxis receptor TlpB and that length variation of this G-repeat determines the outcome (repression or activation) of RepG-mediated post-transcriptional regulation. More recently, we have identified additional RepG targets, including genes involved in LPS biosynthesis or oxidative stress response, and a potential functional RepG homolog in C. jejuni. The example of RepG shows that identifying and studying sRNAs in bacteria without Hfq can reveal new twists in RNA-mediated regulation and will provide new insights into regulatory mechanisms and virulence control of bacterial pathogens.
The dissimilation of sulfur compounds is likely to have been one of the first energy metabolisms on the early Earth [1]. However, many questions remain about how sulfur-metabolizing organisms obtain energy for growth from reducing, oxidizing or disproportionating sulfur compounds. In the case of sulfate reduction, the terminal reductases involved (APS reductase, AprAB, and dissimilatory sulfite reductase, DsrAB) have long been identified, but how these two reactions are coupled to energy conservation is still not clear. DsrAB, in particular, is a key enzyme in dissimilatory sulfur metabolism, being present not only sulfate/thiosulfate/sulfite reducing organisms, but also in sulfur-oxidizers and sulfur disproportionators.

Our lab has studied several of the key proteins involved in sulfate reduction, including AprAB, DsrAB and two respiratory membrane complexes specific to sulfur-metabolizing organisms, QmoABC and DsrMKJOP, which are involved in the electron transfer pathways with AprAB and DsrAB [2]. These complexes hint at the possibility of menaquinone involvement and chemiosmotic energy conservation, during sulfate reduction. In addition, a genomic analysis of energy metabolism genes in sulfate reducers suggested that the recently recognized process of electron bifurcation may also be involved [3], and that there are conspicuous links between sulfate reducers and methanogenic organisms [1].

In this talk I will present an overview of the role of these proteins in sulfate reduction, with a special focus on the function of the small protein DsrC as a physiological partner of DsrAB and the DsrMKJOP complex [4].

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4- Venceslau SS et al 2014 Biochim. Biophys. Acta-Bioenergetics 1837, 1148
New Insights into the Metabolism of Nitrate-driven Anaerobic Oxidation of Methane

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Methane oxidation is an important process to prevent emission of the greenhouse gas methane and further exacerbating of climate forcing. Both aerobic and anaerobic microorganisms have been reported to catalyze methane oxidation with only few substrates as electron acceptors. New microorganisms were discovered that coupled the oxidation of methane to nitrate or nitrite reduction [1-3]. Here, we investigated such an enrichment culture on physiological, biochemical, transcriptomic and genomic level to establish a metabolic model of nitrate-driven anaerobic oxidation of methane (nitrate-AOM). Nitrate-AOM is catalyzed by an archaeon closely related to methanogens. Methane may be activated by methyl-CoM reductase and subsequently undergo full oxidation to carbon dioxide via reverse methanogenesis. All enzymes of this pathway were present and expressed in the investigated culture. The genome of the archaeal culture encoded a variety of proteins involved in a membrane-bound electron transport chain similar to those found in Methanosarcina species. Nitrate reduction seems to be located in the extracellular space and may be catalyzed by Nxr- or Nar-like protein complexes. The enrichment culture produced mainly nitrite but also some ammonium during nitrate reduction, presumably by the action of an extracellular Nrf enzyme. One of the key questions is how electrons from cytoplasmic reverse methanogenesis reach the nitrate reduction enzymes in the extracellular space. We will present a tentative model, based on genome and transcriptome studies, how electrons from reverse methanogenesis are transported to the nitrate and nitrite reductases, thereby suggesting how central energy metabolism of nitrate-driven anaerobic oxidation of methane could work.

This work was supported by the ERC AG 339880 and the SIAM Gravitation Grant 24002002.

Acetogens constitute an important limb in anaerobic food webs. Synthesis of acetate from carbon dioxide and molecular hydrogen via the Wood-Ljungdahl pathway is thermodynamically at the edge of life since it allows for the synthesis of only a fraction of an ATP. How the pathway is coupled to the net synthesis of ATP has been an enigma for a long time, but recently, new insights have been obtained using the anaerobic, acetogenic bacterium *Acetobacterium woodii* as a model system (1,2). The energy conserving module comprises of a Na⁺ translocating ferredoxin-NAD oxidoreductase (Rnf complex) (3) and an unusual Na⁺-F₅,F₅ ATP synthase (4). Endergonic reduction of ferredoxin with H₂ as electron donor is achieved by energy coupling via electron bifurcation (5). Reduction of CO₂ to formate with NADH would be highly endergonic and this energetic barrier is overcome by a novel hydrogen-dependent CO₂ reductase (6).

Metal reduction in bacteria
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Under anaerobic conditions many microorganisms are able to utilize insoluble metal species during respiration. In Gram-negative organisms this requires electrons that accumulate in the menaquinol pool to be transported across the periplasm, through the outer membrane and ultimately transferred to an extracellular electron acceptor. One of the best-studied organisms capable of metal respiration is the facultative anaerobe *Shewanella oneidensis*, which produces a range of multiheme cytochromes that facilitate electron transfer to extracellular minerals.

During mineral respiration *S. oneidensis* utilizes a porin-cytochrome complex to shuttle electrons from the periplasm across the outer membrane. This complex is comprised of a periplasmic decaheme cytochrome (MtrA), a 28-strand porin (MtrB) and an outer membrane decaheme cytochrome (MtrC) that forms the interface between the cell and mineral surfaces. During anaerobic respiration *S. oneidensis* also secretes trace amounts of flavin mononucleotide (FMN) and this has been linked to effective reduction of insoluble metals. FMN was originally proposed to function as a soluble electron shuttle mediating electron exchange between extracellular cytochromes and the mineral surface, however we have recently shown that FMN binds tightly to the extracellular MtrC, becoming an eleventh cofactor under anaerobic conditions. These findings, coupled with new structural and kinetic information have allowed us to reconsider how the cytochromes of *Shewanella* are used during dissimilatory metal reduction.
Unravelling autoinducing peptide biosynthesis and recognition in staphylococci

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The agr quorum sensing (QS) system plays a pivotal role in the co-ordination of virulence in Staphylococcus aureus and other closely related Gram-positive bacteria. The agr locus consists of two divergent transcriptional units controlled by the agrP2 and agrP3 promoters. The agrP2 operon consists of four genes, agrBDCA. AgrB processes the pre-propeptide AgrD to generate a macrocyclic auto-inducing peptide (AIP). This is exported from the cell and subsequently recognized by AgrC, a transmembrane receptor kinase that activates the response regulator, AgrA. This drives expression of the agrP2 operon completing the autoinduction loop as well as the agrP3 operon which codes for both delta-toxin and a regulatory RNA, RNAIII, the primary effector of the agr response. The agr locus has undergone evolutionary divergence such that the AIPs of four different S. aureus agr groups self-activate but cross-inhibit. Consequently to retain functionality, any changes in the AIP-encoding gene (agrD) that results in modification of the AIP structure must be accompanied by corresponding changes in the AgrC receptor. To better understand the biosynthesis and recognition of staphylococcal AIPs we have exploited both in vivo and in vitro chemical biology approaches to gain new insights into the function of AgrB and the structural features of the AIPs and AgrC that determine activation or inhibition of the sensor kinase. This work is of particular relevance to the design of agr inhibitors as antivirulence agents for the prevention and treatment of staphylococcal infections.
Just like what occurs in humans, plants have been recently recognized as meta-organisms possessing a distinct microbiome which has close relationship with their associated microorganisms. The plant microbiome presents an additional reservoir of genes that the plant can have access to when needed. Plant health is thought to heavily depend also on its microbiome and plants most probably intimately affect bacterial gene expression via a communication highway of signals produced by bacteria and plants. These complex highways will become clearer through the detection of signal molecules and identifying the effects on gene expression of microbial and plant exudates. Although plant microbiome composition is now being well studied, there is hardly any information on these signals.

Interspecies and interkingdom signaling, is now a fast developing field of research; we are using plant associated bacteria to investigate these types of communication. We have identified a new sub-family of bacterial LuxR proteins which is widely present in plant associated proteobacteria which binds and consequently responds to plant signals. In addition we are using a plant disease as model of interspecies bacterial interactions which also highlights the role of these interactions in bacterial plant pathogenicity. Understanding these types of communication between microbiomes and plants are a step in the right direction for disease management.
The DSF family of bacterial cell-cell signal molecules

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The Diffusible Signal Factor (DSF) family of cell-to-cell signalling molecules comprises cis-2-unsaturated fatty acids that differ in their chain length and branching. DSF signalling was first described in the plant pathogen *Xanthomonas campestris* (Xcc) where the signal was identified as cis-11-methyl-dodecenoic acid. The synthesis and perception of the DSF signal in Xcc require products of the *rpf* gene cluster (for regulation of pathogenicity factors). The synthesis of DSF is dependent on RpfF, whereas the RpfC/RpfG two-component system is implicated in DSF perception and signal transduction. RpfC is a complex hybrid sensor kinase whereas the RpfG regulator has a CheY-like receiver domain attached to an HD-GYP domain, which acts to degrade the second messenger cyclic di-GMP, leading to regulation of the expression of virulence factors. Recent work has described a second receptor for DSF in Xcc; the RpfS sensor kinase binds DSF through an N-terminal PAS domain to control factors involved in the epiphytic phase of the disease cycle. Although DSF signalling has now been described in several *Xanthomonas* species and the related genera *Xylella* and *Stenotrophomonas*, it is not restricted to xanthomonads. DSF family signals are produced by the human pathogens *Burkholderia cenocepacia* and *Pseudomonas aeruginosa* where they act to regulate virulence, biofilm formation and antibiotic tolerance. Two sensors for DSF family signals have been described in *B. cenocepacia*: a membrane-associated histidine kinase unrelated to RpfC and a soluble PAS-GGDEF-EAL domain protein that couples DSF sensing to cyclic di-GMP turnover. In addition to intraspecies signalling, DSF family signals have been implicated in interspecies signalling that modulates bacterial behavior to include antibiotic tolerance. An understanding of these diverse signalling mechanisms could suggest strategies for interference, with consequences for disease control.
Quorum sensign in rhizobia: getting attached to their roots

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Background: Most rhizobia have N-acyl-homoserine (AHL)-based quorum sensing regulatory systems. These influence phenotypes such as plasmid transfer, exopolysaccharide synthesis and processing, biofilm formation, motility and root attachment.

Objectives: In Rhizobium leguminosarum bv. viciae, which forms nitrogen-fixing nodules on peas and vetch, quorum-sensing regulation plays an important role in competitiveness for nodule infection. Our aim was to identify the key genes for this and their regulation.

Methods: We identified a small protein (CinS) that is co-expressed in a population-density-dependent manner with the AHL synthase CinI. CinS binds to the repressor PraR, thereby relieving repression of several genes including LuxR-type quorum-sensing regulators. Mutation of praR enhanced biofilm formation in vitro and on legume roots; this was correlated with enhanced infect infection of pea nodules in competition with the wild-type. Analysis of gene expression by microarrays and promoter fusions revealed that PraR represses its own transcription and the expression of several genes including those encoding secreted proteins (including the adhesins RapA2, RapB and RapC, a cadherin and the glycanase PlyB). Single mutations in rapA2, rapB, rapC, plyB, the cadherin or rosR did not affect the enhanced root attachment or nodule competitiveness of the praR mutant. However combinations of mutations in rapA, rapB and rapC abolished the enhanced attachment and nodule competitiveness.

Conclusions: Relief of PraR-mediated repression via quorum-sensing regulation determines a lifestyle switch allowing the expression of genes that are important for biofilm formation on roots and consequently enhances infection of legume roots and formation of nitrogen-fixing nodules.

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Towards new TB vaccines - what are the challenges?

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To control the global threat from tuberculosis, we urgently need new TB vaccines that induce effective protection, as well as new drug regimens and diagnostic tools. However there are significant challenges. A new TB vaccine might replace the current BCG vaccine at birth, or be given as a boosting vaccine following BCG (or a BCG replacement vaccine) to infants or young adolescents. A post exposure vaccine could be given to older adolescents or adults. In addition, a therapeutic vaccine could be given to those with clinical disease to accelerate bacterial clearance. These vaccines could include improved BCG vaccines, safe live mycobacteria, antigens delivered by viral vectors, and proteins delivered with adjuvant. Vaccines could be injected or delivered by the mucosal route. There is a vaccine development pipeline with ongoing international efforts to coordinate and facilitate the development and testing of the most promising candidates. These efforts would be accelerated if we had biomarkers or protective biosignatures that predicted the protective efficacy of a new vaccine. Although there is increasing investment in non-human primate models, at present we also do not have animal models that reliably predict protection in man, so the main challenge is how to prioritise development of the best candidate vaccines. Although some measures of immunity have proved disappointing, there are promising developments exploiting gene expression signatures and novel mycobacterial growth inhibition assays. Identifying protective biosignatures, which may differ for different types of vaccines, would greatly facilitate the prioritisation of new TB vaccine candidates for further development. It would also be useful if we could understand better why the existing BCG vaccine induces such variable protection, to ensure that any new TB vaccine will work irrespective of location.
One of the major challenges in the field of TB vaccines is the lack of a correlation of protection what necessary for efficacy evaluation of new vaccines in clinical trials in endemic countries. Live attenuated vaccine candidate MTBVAC is an attenuated Mycobacterium tuberculosis (MTB) genetically engineered to fulfil the Geneva consensus requirements to enter human clinical trials. We selected a MTB clinical isolate belonging to Euro-American lineage to generate two independent deletions without antibiotic-resistance markers. One is the phoP, essential for virulence in MTB where it controls expression of approximately 2% of the genes, including those for the ESX-1 secretion apparatus, a major virulence determinant and the secretion of the immunodominant Ag85 complex. Deletion in phoP gene lead to compromised production of pathogen-specific cell wall components and attenuation both ex vivo and in vivo. Deletion in fadD26, inactivate phthiocerol dimycocerosates (DIM) operon which, codes for the production of one of the major MTB virulence factors.

MTBVAC exhibits safety and biodistribution profiles similar to BCG and confers superior protection in preclinical studies. A major difference between MTBVAC and BCG, which is derived from the cattle pathogen M. bovis, is that BCG has many deletions in its genome when compared to MTB, and during the attenuation process BCG had lost over a hundred additional genes from its genome including immunodominant antigens. After more than 15 years of preclinical development MTBVAC is the first live attenuated vaccine to enter clinical evaluation in Lausanne.
(Switzerland) with Prof Spertini as Principal Investigator and sponsored by Biofabri and TBVI (ClinicalTrials.gov: NCT02013245).
Background
Tuberculous meningitis (TBM) is a major disease burden in children in TB endemic countries.

Objectives
We use our zebrafish – Mycobacterium marinum infection model for TBM to study early granuloma formation in brain and meninges of zebrafish embryos.

Methods
We dissected the migration mechanism employed by M. marinum to cross the blood-brain barrier (BBB) and cause infection of the central nervous system. To study the role of macrophages in this process, we depleted the macrophage pool in zebrafish embryos with pu.1 morpholinos and clodronate filled liposomes. Embryos lacking phagocytic cells showed an increased overall infection rate and a higher bacterial load in the head at 5 days post infection in comparison to the control group. Detailed analysis showed that in the presence of macrophages, M. marinum leaves the bloodstream and forms early granulomas consisting of mycobacteria and L-plastin positive phagocytes in brain tissue. Virtually all M. marinum inside and outside the blood vessels were present in phagocytic cells, suggesting that mycobacteria use macrophages as a Trojan horse to cross the BBB. As expected, we did not observe granuloma structures after macrophage depletion. However, we still found extravasated mycobacteria in the brain parenchyma. Most of these mycobacteria were found in close vicinity of blood vessels, following the blood vessel shape in a longitudinal manner.

Conclusions
We observed that M. marinum preferably use macrophages as Trojan Horses to cross the BBB. However, in the absence of macrophages M. marinum can adept and seems capable of using additional migration routes.
THE ROLE OF A TYPE VII SECRETION CHAPERONE IN THE SPECIFIC SUBSTRATE RECOGNITION IN PATHOGENIC MYCOBACTERIA

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Background

Type VII secretion (T7S) is employed by mycobacteria, including the important pathogen *Mycobacterium tuberculosis*, to export protein effectors across their highly unusual cell envelope. *M. tuberculosis* contains five paralogous T7S systems, ESX-1 to ESX-5, each having its own role in viability and/or virulence. The mycobacteria-specific PE/PPE proteins are one group of proteins that are secreted via these secretion systems, of which at least some are secreted as heterodimers. We have previously shown that the cytosolic chaperone EspG specifically interacts with cognate PE/PPE dimers and is required for their successful secretion.

Objectives

Our aim is to understand the determinant factors for substrate-specific binding of EspG to T7S substrates.

Methods

The crystal structure of ESX-5 chaperone EspG 5 with the ESX-5 substrate pair PE25/PPE41 was determined and important residues for chaperone-binding of several PPE proteins were mutated. Additionally, EspG-binding domains of ESX-1 and ESX-5 substrates were exchanged.

Conclusions

The structure revealed that EspG 5 interacts with the elongated end of PPE41, a region containing several hydrophobic residues that are conserved in PPE substrates of different secretion systems. Mutating these residues of multiple ESX-5 PPE substrates and an ESX-1 dependent PPE protein abolished the interaction with the cognate chaperone and blocked their secretion by *Mycobacterium marinum*. Subsequently, exchanging the complete EspG-binding domain between an ESX-1 and ESX-5 PPE protein altered chaperone-binding specificity. The effect of system-specific chaperone-binding on rerouting of T7S substrates is currently tested. In
summary, we have elucidated the molecular determinants for EspG-specific binding to PPE substrates and the key role of chaperone-binding in secretion.
MYCOBACTERIUM LEPRAE (M. LEPRAE) CREATES LIPID-RICH INTRACELLULAR ENVIRONMENT BY MODULATING EXPRESSION OF HOST GENES IN INFECTED MACROPHAGES

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Background

*Mycobacterium leprae* (M. leprae), the causative agent of leprosy, parasitizes within lipid droplets stored in enlarged phagosome of histiocytes, which is thought to be important nutrient source and sheltering for the bacillus. However, the mechanisms by which *M. leprae* triggers lipid accumulation within phagosomes remain unclear.

Objectives

To elucidate the mechanism underlying *M. leprae*-induced lipid accumulation in macrophages.

Methods

Real-time PCR, western blotting, histochemical staining and DNA microarray were employed to analyze human premonocytic THP-1 cells infected with *M. leprae* as well as skin biopsies and smears of leprosy patients.

Conclusions

Live *M. leprae*, but not heat-killed *M. leprae*, specifically induced adipose differentiation-related protein (ADRP) and perilipin, which are essential regulators for lipid accumulation, in THP-1 cells. However, live *M. leprae* suppressed and inactivated hormone sensitive lipase (HSL). Clofazimine, one of the drugs for treating leprosy, devastated the *M. leprae*-induced lipid-rich intracellular environment by modulating the expression of ADRP and HSL, and by activating the innate immune response in host macrophages. Staining of skin biopsies and smears confirmed the above results. DNA microarray analysis revealed that live *M. leprae* modulated gene expressions important for lipid accumulation, as well as their common transcriptional factors, *i.e.* peroxisome proliferator-activated receptor (PPAR)δ and PPARγ. The antagonists of PPARδ and PPARγ abolished the effect of *M. leprae* to modulate such gene expressions and lipid accumulation in macrophages. These results indicate that live *M. leprae* creates lipid-rich intracellular environment by modulating gene expressions of host macrophages through PPARδ and PPARγ signaling.
Microbes propel themselves in beneficial directions using cell wall-embedded rotary motors that spin helical propellers. Bacteria swim using bacterial flagellar motors, large nanomachines composed of hundreds of proteins — from approximately twenty protein families — that harness proton-motive force to convey torque to a multi-micron flagellum, while archaea assemble “archaella” that utilize ATP hydrolyse to rotate an analogous filament. How do these function, and how did they evolve? To understand this we are using electron cryo-tomography to perform 3D in situ imaging of these motors. We have recently shown that bacterial flagellar motors are considerably more complex than the ‘normal’ core motor found in *Salmonella* and *E. coli*, thus shedding light on the evolutionary processes used to adapt macromolecular machinery. Why have some bacteria recruited additional proteins to a core that is already fully functional? I will describe phylogenetic and structural work to understand this question. We are currently focusing on a family of large novel structural components found in a variety of proteobacteria including the Vibrio, Campylobacter, and Helicobacter genera. By combining bacterial genetics, phylogenetics and electron cryo-tomography we have identified the proteins that form these large structures. Combining these results we have developed a model for the recruitment of these additional proteins, and have been able to speculate on the effect of these proteins on motor mechanics, in turn highlighting possible evolutionary driving forces behind protein recruitment. I will then describe recent work to study the analogous archaellar motor.
Cryo-EM of the Type VII secretion system in Mycobacterium tuberculosis

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We showed in Cell; 129:1287 that after prolonged infection in macrophages and dendritic cells, M. tuberculosis translocates from phago-lysosomes to the cytosol and killed the host cell a few days later, while the BCG vaccine strain failed to translocate. We found that this process was dependent on a gene in the extended RD1 region (ext-RD1). We then focused on BCG with a knock-in of the entire ext-RD1, which was recently identified by one of our former team member as a novel type VII secretion system (T7SS) (Abdallah AM et al., Nat Rev Microbiol. 2007). We found that these bacteria translocate to the cytosol of the host cell 7 days after infection. We concluded that the ESX-1 system is sufficient for translocation (Abdallah AM et al., J of Immunol. 2011 and Houben et al., Cell Microbiol. 2012).

Since my move to Maastricht University where I was invited to establish a new Institute for Nanoscopy (www.maastrichtuniversity.nl/m4i) we are now using cryo-EM single-particle analysis (SPA), and cryo-EM tomography. We investigate recombinant purified proteins of individual gene products from the T7SS. In addition, we are purifying the entire intact T7SS structure using biochemical methods for 3D reconstruction. We will also generate lamellae of infected cells using cryo-FIB/SEM technology. This should lay the groundwork for the development of novel antibiotics and better vaccines. Since this FEMS meeting is next door to my new laboratory interested colleagues are welcome to contact me for a visit (Universiteitssingel 50, Room G0.201) while attending the meeting (peter.peters@maastrichtuniversity.nl).
FEMS-2842
Imaging of macromolecular structures

The assembly and mechanism of a bacterial drug efflux pump
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Microorganisms encode several classes of transmembrane molecular pumps that can expel a wide range of chemically distinct toxic substances. These machines contribute to the capacity of the organisms to withstand harsh environments, and they help to confer resistance against clinical antimicrobial agents. In Gram-negative bacteria, the pumps comprise tripartite assemblies that actively transport drugs and other harmful compounds across the cell envelope. We describe recent structural and functional data that have provided insights into the architecture and transport mechanism of the AcrA-AcrB-TolC pump of Escherichia coli. This multi-drug efflux pump is powered by AcrB, a member of the resistance/nodulation/cell division (RND) family of transporters, which are energised by proton electrochemical gradients. Crystallographic data reveal how a small protein AcrZ is engaged in a concave surface in the transmembrane domain of AcrB, and we discuss how this interaction may affect the efflux activities of AcrB and other RND family members.
Interactions of a bacterial cell with other bacteria, eukaryotes or archaea are crucial for symbiosis, pathogenicity, competition and defense. These bacterial cell-cell interactions are often mediated by macromolecular machines. In evolution, a phage-derived bacterial contractile structure proved to be very successful in mediating cell-cell interactions in an either contact-dependent or indirect manner. We applied electron cryotomography in an interdisciplinary approach to understand the structure and mechanism of the type VI secretion (T6S) system, arrays of metamorphosis-associated contractile structures (MACs), and other bacterial contractile assemblies. State-of-the-art sample preparation techniques recently allowed us to image bacteria engaging with a eukaryotic host.
Bacterial phosphorylation networks

New insights into the GacS multikinase network controlling virulence in Pseudomonas aeruginosa

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The GacS/A two-component signalling network of Pseudomonas aeruginosa plays a major role in controlling virulence. This network features multiple sensor kinases that work together to decide between acute and chronic modes of infection [1,2]. Signalling by the kinase GacS favours chronic infection, while another kinase, RetS, favours acute infection by inhibiting GacS. We have been investigating how RetS interferes with GacS function and we have found that there are three distinct ways in which RetS can inhibit GacS: 1) RetS can accept phosphoryl groups from GacS-P; 2) RetS has phosphatase activity towards GacS-P; and 3) RetS prevents autophosphorylation of GacS. We can selectively disable each of these mechanisms by introducing specific point mutations into RetS and have tested the effects of these mutations in a mouse model of respiratory infection. We find that loss of either mechanism 1 or 2 dramatically decreases acute virulence, while loss of both mechanisms 1 and 2 abolishes acute virulence. RetS function in vivo therefore depends upon its ability to bring about the dephosphorylation of GacS-P. This study reveals an unexpected degree of complexity in the intricacy of the interactions between the sensor kinases governing the choice between different modes of virulence in this important pathogen.


Bacterial phosphorylation networks

How do you Coordinate signals from two chemosensory pathways to control one flagellar motor?
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1
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The chemotaxis pathway of E.coli is probably the best understood sensory pathway in bacteriology. A kinase, CheA, autophosphorylates in response to a decrease in attractant, and transfers a phosphoryl to either a diffusing single domain response regulator, CheY, which when phosphorylated binds the switch protein of the flagellar motor to cause a change in rotational direction, or phosphorylates a methyl esterase controlling receptor adaptation. However, many bacterial species have more than one chemosensory pathway controlling the direction of flagellar rotation.

The phototrophic bacterium Rhodobacter sphaeroides expresses two pathways, one localised with transmembrane receptors to large patches on the membrane and one localised with soluble chemoreceptors, and associated with the chromosomes, segregating with the chromosomes on division to ensure each daughter inherits a cluster.

In vitro and in vivo methods, including phosphotransfer, fluorescence microscopy, cyro EM, molecular genetics, mass spectometry, live cell tracking and mathematic modelling have been used to develop an understanding of how the direction of rotation of a single flagellar motor is controlled by integrating responses to extracellular and intracellular signals through the two pathways. Recent work suggests that signals from the cytoplasmic cluster dominate and R. sphaeroides tunes its swimming behaviour in response to current metabolic requirements.
Two-component systems (TCS) are the predominant signal transduction mechanism in bacteria, are present in lower eukaryotes such as fungi and plants, but they are absent from mammals. The key players in these systems are two proteins, histidine kinases (HKs) along with response regulators (RRs) which transmit signals through phosphoryl-transfer. HKs are sensor dimeric proteins that, upon signal detection, bind ATP and autophosphorylate in a catalytic histidine residue. Transfer of the phosphoryl group from the HK histidine to a conserved aspartic of its cognate RR propagates the signal and triggers the cellular response. Signal cease by the auto or HK-mediated de-phosphorylation of the RR. The catalytic machinery in charge of carrying out the reactions related with signaling process, either in the HK or the RR, is highly conserved and the couple interaction shows high-fidelity. Despite being the more abundant signal transduction systems in the nature, the molecular bases of these reactions are neither fully understood at biochemical level nor well-characterized at structural level, as HK autophosphorylation reaction is known to be conducted via a cis or trans mechanism in different HKs or the HK-RR phosphoryl transfer and HK-mediated RR de-phosphorylation have been demonstrated not to be reversible. In an effort to clarify the molecular basis of TCS signaling we have worked during the last 10 years with HK853-RR468 and EnvZ-OmpR, two prototypical TCS. Our functional and structural results, which will be discussed in the presentation, shed light on the molecular basis of each reaction and their interrelationship with HK-RR recognition and specificity.
To cause plant disease, pathogenic fungi can secrete effector proteins into plant cells to suppress plant immunity and facilitate fungal infection. Most fungal pathogens infect plants using very long strand-like cells, called hyphae, which secrete effectors from their tips into host tissue. How fungi undergo long-distance cell signalling to regulate effector production during infection is not known. Here, I show that long-distance retrograde motility of early endosomes (EEs) is necessary to trigger transcription of effector-encoding genes during plant infection by the pathogenic fungus *Ustilago maydis*. This motor-dependent retrograde EE motility is controls effector production and secretion during host cell invasion. It involves the mitogen-activated kinase Crk1, which travels on EEs and participates in control of effector production. Fungal pathogens therefore undergo endosome-mediated long range-signalling to orchestrate host invasion. These findings add to recent discovery that EEs distribute the machinery for protein translation. Thus, 15 years after the discovery of motile endosomes in fungi, the biological role of their motility is beginning to reveal.
Peroxisome biogenesis and proliferation in yeast

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Peroxisomes are cell organelles that are present in almost all eukaryotic cells. They can harbor a large variety of enzymes, which highly depends on the organism, tissue and developmental stage. Common peroxisomal functions include the beta-oxidation of fatty acids and hydrogen peroxide metabolism. The importance of these organelles is illustrated by the fact that peroxisome malfunction results in severe disorders and is often lethal in man.

In yeast and filamentous fungi peroxisomes play a role in the metabolism of various unusual carbon and organic nitrogen sources, such as alkanes, fatty acids, methanol, primary amines and uric acid. Notably, in filamentous fungi peroxisomes are crucial for the biosynthesis of commercially important metabolites, such as beta-lactam antibiotics.

We study peroxisome biology using Saccharomyces cerevisiae and Hansenula polymorpha as model organisms. An important and highly debated topic is how new peroxisomes are formed. One model proposes that peroxisomes are semi-autonomous organelles that originate by fission of pre-existing ones, similar to mitochondria. According to an alternative model peroxisomes form de novo from the endoplasmic reticulum, which implies that peroxisomes form a branch of the endomembrane system.

We recently re-investigated both pathways using advanced electron and fluorescence microscopy techniques. Our data suggest that in yeast peroxisome fission is the major mode of peroxisome proliferation.
To define the general principles of genetic networks, our group developed a functional genomics platform called ‘synthetic genetic array’ (SGA) analysis that automates yeast genetics and enables the systematic construction of double and triple mutants. One of our major goals has been to use a simple phenotypic readout of cell growth rate – colony size – to produce the first complete genetic interaction map for any cell, and to empirically delineate the properties of genetic networks. We have now analysed the majority of all possible 18 million yeast gene pairs. The resultant network consists of ~560,000 genetic interactions, spanning 93% of all yeast genes. Analysis of the network has revealed: [1] a central role for and unique properties of essential genes; [2] hubs and pleotropic genes on the network which show a clear association with several fundamental physiological and evolutionary properties that are predictive of genetic interactions in other organisms; [3] functional modules that we use to predict and test conservation of interactions in other systems.
FEMS-2850
Fungal cell biology

The importance of fungi and their movement in crops
Background

Teaching and research are main jobs of a university professor. Both tasks should be nourished by public dissemination of scientific matters. Under the acronym of “microBIO” I have group together several on line tools for the dissemination and spread news and curiosities about microbiology and science in general.

Objectives

In an understandable and entertaining way, with scientific accuracy, the goals of microBIO are: i) make science and microbiology understandable to all kind of people, ii) wake up the curiosity for microbiology and arouse young scientific vocations, iii) avoid the isolation of scientist and increase the visibility of our work.

Methods

microBIO get together several tools on line, open and free: blog, twitter, Facebook, Scoop.it, Storify, YouTube channel, TEDx talks,… We use social networks to disseminate and share images, teaching resources, videos, links and all kind of scientific information of interest.

Conclusions

Two experiences have special attention. 1) Massive Online Open Courses (MOOC) as introductory courses of general microbiology and virology: “Microbes around us” and “Pandemics and new viral infections”, both for people with interest in scientist topics, primarily undergraduate students, first-year university course, science teachers and journalists; ii) microbiology master classes by twitter, during 8 weeks, once a week, we share around 30 “tweets” with the hashtag #microMOOC. In only 140 characters we taught different subjects related with microbiology.

microBIO has thousands of followers from all over the world, 37% from Spain and
the remainder from other countries: México, USA and Latin-America. microBIO demonstrates that microbiology can also be “trending topic”.
DEVELOPING EUROPEAN COMMON CURRICULUM FOR MICROBIOLOGY EDUCATION: IS THERE A NEED?

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Background

Microbiology including General and Applied Microbiology is main subject in Biology (Microbiology) education at European universities. Microbial Biotechnology is important for training in Biology education and Clinical Microbiology - Medicine one. American Society for Microbiology is developing a curriculum. European common curriculum would be of significance to train specialists on the unified high level.

Objectives

The motivations to train microbiologists in European countries would be clarified; the role of Microbiology in Biology education and main parts in Microbiology content should be defined.

Methods

Education practice in Bologna process framework as well as American, European national or university curriculums would be compared and analysed.

Conclusions

To have European common curriculum the following would be of significance:

- for motivations: description and preservation of wide biodiversity, including microorganisms, in different regions in Europe; isolation, identification and study of microorganisms in different ecological environment, discovering their role in nature; monitoring and control of potentially dangerous microorganisms in nature and among habitants, developing new strategy against endemic pathogens; using microorganisms in food and pharmaceutical industry; developing bio-energy production; control of biosafety; biology education at secondary schools.

- Three modules are significant for education: Biodiversity, including biology of microorganisms; Physiology, including microbial physiology; Physico-Chemical
Biology. General (Basic and Systematic) Microbiology and Microbial Biotechnology and/or Clinical Microbiology can be recommended as main topics.

- For teaching issues improving computer and lab works would be efficient in learning Microbiology and developing appropriate practical skills.

To have acceptable curriculum and to implement in different countries, adoption by FEMS (Education Group) would be important.
Background

As stated by the European Charter for Researchers, scientists should ensure that their research activities are made known to society at large in such a way that they can be understood by non-specialists, thereby improving the public’s understanding of science (EC 2005). Science communication is considered a fundamental tool to promote science study, to raise the visibility of universities and their research groups, and to influence policy decisions and funding opportunities.

Objectives

The aim of this work was to create an animated cartoon to raise awareness among young people, citizens, end-users and policy makers about the significance and the broad impacts of the EU funded project ESENCYA “Environmental SENsory perception in CYAnobacterial biofilms: understanding biodeterioration of outdoor stone materials in a changing environment”.

Methods

By working with a multifunctional team composed of character designers, visual development artists, communication specialists and scientists, we created a visually engaging story about complex concepts in new and innovative ways (Fig. 1). We translated complex information and ideas into intuitive and effective visual explanations, and we used a narrative strategy to get our message across, clearly and concisely while reducing misunderstanding.
Conclusions

This animated cartoon wants to encourage and inspire actions directed towards raising public awareness of the importance of sciences for our contemporary society in a cultural and historical perspective. The microbiology community, in Europe and around the world, can find ideas, blueprints and suggestions for activities directed towards raising public awareness of science, technology and other fields where microbiology plays a strong role.
Clostridium botulinum and the safety of modern foods

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Foodborne botulism is a severe and often fatal neuroparalytic intoxication caused by consumption of neurotoxin formed by strains of Clostridium botulinum (and very occasionally by strains of C. baratii or C. butyricum). C. botulinum is defined on the basis of one physiological property, an ability to form botulinum neurotoxin. It is a heterogeneous species that is separated into four distinct phylogenetic/physiological groups (C. botulinum Groups I to IV). The distinction between each Group is sufficient to merit separation into different species. Seven botulinum neurotoxins (types A to G), and >30 neurotoxin sub-types are recognised. The botulinum neurotoxins are the most potent toxins known; 50ng of neurotoxin may cause human illness and possibly death.

C. botulinum Groups I and II and the neurotoxins they form (types A, B, E and F) are responsible for most cases of foodborne botulism. These two physiologically distinct bacteria present different problems. C. botulinum Group I (proteolytic C. botulinum) is a mesophile (minimum growth temperature 10°C-12°C) that forms highly heat resistant spores, while C. botulinum Group II (non-proteolytic C. botulinum) is a psychrotroph (minimum growth temperature 2.5°C-3.0°C) that forms moderately heat resistant spores.

Recent advances in genomics have increased understanding of the biology of C. botulinum Groups I and II, while developments in mathematics have improved risk assessments. The safe development of novel foods and new food processes requires strong control measures for C. botulinum Groups I and II. The extreme severity of foodborne botulism ensures that regulators and industry remain vigilant to minimize the foodborne botulism risk.
NEW INSIGHT INTO BIOGENIC AMINES: TOWARDS SOLUTION AND BIOTECHNOLOGICAL APPLICATIONS


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Background
Biogenic amines (BA) are low molecular weight nitrogenous compounds mainly formed by the microbial decarboxylation of amino acids. The primary relevance of BA is that the consumption of foods or beverages containing high concentration of BA may cause food intoxication. Dairy products are among the foods likely to contain the highest concentrations of BA, being lactic acid bacteria the main BA-producers.

Objectives
Increase our knowledge on BA synthesis in order to adopt strategies for surveillance and prevention of BA accumulation in foods and towards the development of new biotechnological tools.

Methods
Strategies to reduce the BA content in fermented products:

- Avoid the presence of bacteria synthesizing BA through the food chain. The identification of BA gene clusters has allowed the design of molecular methods to identify and quantify the presence of BA producers in cheese samples.

- Identify key factors which favor BA accumulation. Different physicochemical and technological factors have been studied in order to determine their impact on BA accumulation either affecting bacterial growth or BA gene expression.

- Recently, the use of biogenic amine-degrading microorganisms has been proposed. It is necessary to identify and select highly competitive adjunct cultures capable of in situ reduce the content of BA during the elaboration of fermented products.

Conclusions
The study of BA synthesis, is needed to propose new strategies to avoid BA accumulation in foods and can be used for the development of new biotechnological tools.
THE PHAGE ENDOlysIN LYsT712 REQUIRES A FUNCTIONAL HOST MEMBRANE PROTEASE FTSH TO LYSE LACTOCOCCAL HOST CELLS

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Background
Bacteriophages are often the cause of milk fermentation failures and selecting phage resistant starters, namely Lactococcus lactis which is widely use in cheese manufacturing, may reduce the risk. L. lactis lacking the membrane protease FtsH became resistant to the phage TP712. In these mutants, viable phages amassed in the cytoplasm but could not be released.

Objectives
Lysis of the phage host is usually accomplished by the action of two phage proteins: a holin that disrupts the cytoplasmic membrane and an endolysin that degrades the peptidoglycan. Our objective was to demonstrate if the activity of any of these proteins was compromised by the absence of FtsH.

Methods
TP712 lysogens were induced with mitomycin C and treated with ionophores to simulate holin activity. Contrary to the wildtype strain, lysis was not triggered in L. lactis ΔftsH, pointing to a failure of the endolysin to degrade the cell wall. However, zymograms revealed the presence of an active LysTP712 endolysin in L. lactis ΔftsH lysogens. A fusion protein consisting of the fluorescent mCherry protein and the cell wall binding domain of LysTP712 fused to its C-terminus was synthesized in E. coli and purified. Binding experiments to whole cells demonstrated that the fusion protein localized at the septum in both WT and ΔftsH lactococcal cells but the latter bound 40% less protein.

Conclusions
Absence of FtsH does not compromise proper functioning of the TP712 holin.

LysTP712 synthesized in L. lactis ΔftsH is active in vitro but its binding to L. lactis ΔftsH cells is reduced.
Background

Transglutaminase is used in different branches of the industry because of its ability to modify physicochemical properties of proteins. It can be the solution to many technological problems related to appropriate efficiency and food texture. Transglutaminase catalyses the formation of cross-links both within a protein molecule and between molecules of different proteins. In bakery industry it was found that at certain level of transglutaminase concentration it has a significant influence on the dough by increase of relaxation time of doughs. Taking into consideration fact that high relaxation times in general correspond to a high quality flour it is believed that transglutaminase would help improve the quality of weak flours.

Objectives

The aim of the experiment is to determine transglutaminase influence on wheat bread quality.

Methods

Transglutaminase was obtained from *Streptoverticillium mobaraense* strain KKP 2013 obtained from Culture Collection of Industrial Micro-organisms. Preparations were sprayed dryed on maltodextrin as carrier. In the experiments there were used wheat flour type 500, two preparations of transglutaminase in four level of addition: 0.01%; 0.05%; 0.25% and 0.5. All physicochemical parameters were determined according to Polish and European Standards.

Conclusions

Taking into consideration crumb characteristics there has been shown an increase in hardness and chewiness, explained as a result of an improvement in the structure of the crumb in comparison to a control breads. The quality of bread in the organoleptic
evaluation was not very diverse. Although the bread with the addition of transglutaminase received higher number of points than the bread without enzyme.
OXIDATION OF FERULIC ACID TO FORM DIMERS WITH HIGHER ANTIOXIDANT CAPACITY

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1fse, lautech, ogbomoso, Nigeria

Background
Ferulic acid is an abundant and widespread cinnamic acid derivative with antioxidant properties. However, its antioxidant activity is generally low when compared to conventional antioxidants, such as butylated hydroxyanisol (BHA) and other hydroxyl-cinnamic acids, such as sinapic acid. Laccases (EC 1.10.3.2) are enzymes that are capable of catalysing the oxidation of various phenolic compounds with the production of water as the only by-product, and are therefore often referred to as ‘green’ enzymes. The use of laccases in the oxidative coupling and polymerisation of phenolic compounds offers potential for the amplification of the antioxidative capacity of these phenolic compounds.

Objectives
This work employed the use of laccase produced from Trametes pubescens to catalyse the modification of ferulic acid as a way of enhancing antioxidant capacity.

Methods
An airlift reactor was used to produce laccase from free cells of Trametes pubescens and the laccase was purified. Reaction of ferulic acid (10mM) with laccase (10 U) was done at 28°C in a biphasic system of ethyl acetate and 100mM sodium acetate buffer pH 5.0. Products were separated, purified, characterized and antioxidant activities were determined.

Conclusions
This work elucidates the structure of two dimers formed from ferulic acid in organic media one of which is more potent than ferulic acid in terms of antioxidant activity. This new product could find useful application in health and cosmetic industry.
RNA-mediated regulatory circuits link stress response, adaptation and virulence in Staphylococcus aureus


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Staphylococcus aureus is a remarkable versatile pathogen, able to cause a wide spectrum of human diseases, and is one of the main causes of community as well as hospital-acquired infections. The contribution of regulatory RNAs in the establishment of virulence in this pathogen is increasingly appreciated. Our previous data emphasize the multitude of regulatory steps affected by RNAIII in establishing a network of S. aureus virulence factors. We show that RNAIII coordinately represses the expression of numerous mRNAs that encode the transcriptional repressor of toxins, several virulence factors acting early in the infection process, and enzymes involved in peptidoglycan metabolism. Besides RNAIII, we demonstrated that the S. aureus genome likely encodes a high diversity of RNAs including cis-acting regulatory regions of mRNAs, cis-acting antisense RNAs, and small RNAs. One of these sRNAs attenuates the severity of acute systemic infections and enhances chronic catheter infection by regulating MgrA, a transcriptional regulator of capsule formation. Because S. aureus is essentially a human commensal, we propose that RsaA has been positively selected through evolution to support commensalism and saprophytic interactions with the host. We will also illustrate how regulatory proteins and RNAs form complex regulatory circuits to express the virulence factors in a dynamic manner.

RNA-mediated regulation of Legionella virulence

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Legionella pneumophila, a gram-negative intracellular pathogen, has evolved complex mechanisms to efficiently enter, replicate inside, and evade their hosts, aquatic amoebae or alveolar macrophages during disease. L. pneumophila has been shown to cycle between an infectious, non-replicating form thought to promote transmission to a new host, and an intracellular, replicative form, which does not express transmission/virulence traits. This biphasic lifecycle is tightly regulated by a complex regulatory cascade, implicating two component systems, sigma factors, RNA binding proteins and small ncRNAs. Here we have employed RNAseq, transcriptional start site (TSS) mapping and RIPseq combined with molecular biology approaches to better understand this regulatory network and the implication of ncRNAs. We first established a genome-wide map of the TSSs and operons for L. pneumophila, and identified over 500 new small ncRNAs that may act as potential regulators of cis- and trans-encoded target messenger RNAs. Among those a high amount of antisense transcription was observed. A detailed search for targets of the RNA-binding protein CsrA, a key regulator in the expression of the virulence phenotype of L. pneumophila led to the identification of over 50 mRNAs regulated by CsrA, many of which are virulence related proteins. Taken together, we identified the mRNA targets of the RNA binding protein CsrA, ncRNAs regulating CsrA as well as ncRNAs implicated in virulence. Furthermore we showed that tandem promoters and sRNAs with alternative usage are a hallmark of the life cycle depend virulence gene expression of L. pneumophila.
Background

The CRISPR-Cas 'prokaryotic adaptive immune system' represents an intricate defence strategy allowing bacteria to protect themselves against invading genetic elements such as bacteriophages or plasmids. Through the acquisition of short segments of invading nucleic acid sequences into genomic CRISPR loci, this system provides a heritable genetic memory of previous exposure and, in combination with various CRISPR-associated (cas) proteins, a means by which the bacteria can specifically recognise, target and eliminate the exogenous threat. In the past 10 years since the discovery of CRISPR-Cas systems, there has been a heavy focus on functionality; how the acquisition of genetic material occurs and how the system coordinates its targeted response. In contrast, very little is known how about how CRISPR-Cas systems are regulated and of the limited literature available, it seems that a universal mode of regulation is perhaps unlikely.

Objectives

In this study, we focused on elucidating the regulatory control of the model Type I-F system in Pectobacterium atrosepticum.

Methods

Unbiased transposon mutagenesis of a cas1 reporter strain was utilised to identify mutants exhibiting differential promoter activity compared with a wildtype control. CRISPR-Cas activity was also assessed in these mutant strains via conjugation efficiency and spacer acquisition assays.

Conclusions

We have identified a network of regulators of CRISPR-Cas expression, highlighting a link to global regulators that sense the nutritional status of the cell. In addition, we have shown that mutating these regulators drastically affects both the primed acquisition and interference stages within CRISPR-Cas-mediated defence.
PROTEOLYSIS OF ANTI-SIGMA FACTORS IN THE TRANSDUCTION OF THE SIGNAL AND ACTIVATION OF EXTRACYTOPLASMIC FUNCTION SIGMA FACTORS

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Background

Regulation of gene expression allows bacteria to adapt to alterations in their environment. This regulation occurs mainly at the level of transcription initiation by modifying promoter recognition of the RNA polymerase, a recognition that is primarily modulated by substitution of the RNAP sigma subunit. Besides a primary sigma factor responsible of the expression of general functions, most bacteria contain alternative sigma factors that target expression of functions required only under specific circumstances. Of those, the extracytoplasmic function sigma factors (σECF) represent the largest group, and control expression of bacterial functions as important as stress responses, iron uptake and virulence.

Objectives

Activation of σECF is a tightly regulated process that usually occurs in response to environmental signals, a control that is carried out by anti-sigma factors. Binding of the σECF to the RNAP and the anti-sigma factor is mutually exclusive. This work focusses on understanding the molecular mechanism behind σECF activation.

Methods

We have used Pseudomonas as bacterial model and a number of in vivo and in vitro techniques to monitor gene expression and protein levels, stability and structure.

Conclusions

Activation of σECF in response to the inducing signal requires the processing of the anti-sigma factor by a complex proteolytic cascade that involves, amongst other, the Prc and RseP proteases. Interestingly, several anti-sigma factors undergo self-cleavage prior to signal recognition by an enzyme-independent process. This cleavage produces two functional domains that interact and function together to transduce the presence of the signal and to activate the σECF. The biological significance of this widespread mechanism is discussed.
RESPONSE OF YEDQ DIGUANYLATE CYCLASE TO PYRIMIDINE NUCLEOTIDES BY DIRECT INTERACTION WITH THE METABOLIC ENZYME ASPARTATE CARBAMOYLTRANSFERASE AND ITS METABOLIC PRODUCT N-CARBAMOYL-L-ASPARTATE

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Background

In *Escherichia coli*, formation of multicellular communities is mediated by the production of adhesion factors such as curli fibres and the extracellular polysaccharide cellulose. Synthesis of curli fibres and cellulose is co-regulated through the activity of the transcriptional factor CsgD and is under the control of several extracellular and intracellular signals, including the second messenger c-di-GMP. We demonstrated that curli and cellulose production are tightly linked to pyrimidine biosynthetic pathways. Indeed, transcription of curli operons is strongly affected by pyrimidine nucleotide availability; in addition, we found that exogenous uracil can trigger cellulose production independently of CsgD, by increasing c-di-GMP levels via the YedQ diguanylate cyclase.

Objectives

Through a combined in vivo and in vitro approach we studied the molecular mechanism underlying the uracil-dependent regulation of YedQ.

Methods

YedQ protein interactors were co-purified using his-tag pull-down technique and then identified through MudPIT technology. Interactions were validated using bacterial two-hybrid assay. Protein ligand binding was determined through differential scanning fluorimetry, while gene expression was measured with qRT-PCR assay.

Conclusions

We found that YedQ can directly interact with aspartate carbamoyl-transferase (PyrBI), an essential enzyme involved in pyrimidine nucleotides biosynthesis, in a manner dependent on uracil availability, and can also bind N-carbamoyl-L-aspartate the product of PyrBI enzyme. We hypothesise that aspartate carbamoyltransferase
and/or N-carbamoyl-L-aspartate can inhibit YedQ diguanylate cyclase activity by directly interacting with its cytoplasmic domain and this negative effect can be reversed by uracil. The YedQ/PyrBI interaction might represent a model of how activity of metabolic pathways can be coupled to a c-di-GMP signalling system.
FEMS-1441
Signalling, small RNAs and regulation: virulence aspects

A NEW TARGET FOR THE SRNA OMRA: THE VIRF MRNA IN SHIGELLA FLEXNERI
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Background

VirF is a transcriptional regulator which plays a central role in controlling virulence gene expression in Shigella spp, the ethiological agent of bacillary dysentery in humans. VirF protein is present in two different forms, independently translated. VirF₃₀ is responsible for activation of the virulence system and some chromosomal genes, whereas VirF₂₁ exerts negative feedback control on virF expression itself.

Objectives

This work investigates the role of OmrA, a sRNA induced under high osmolarity conditions, on the translation of VirF₂₁ and its impact on Shigella virulence.

Methods

In vitro translation, translational lacZ –fusions, OmrA overexpression were used to evaluate the effect of OmrA on VirF₂₁ translation. RNA footprinting was performed to map the OmrA binding site on virF mRNA.

Conclusions

We have provided evidence supporting the involvement of OmrA in the translational regulation of Shigella VirF. In particular, we have shown that under high osmolarity conditions the expression of a VirF₂₁ fusion is decreased severely in E.coli wt as compared to a defective omrBA derivative. Overexpression of OmrA, but not of OmrB, leads to a strong decrease in VirF₂₁ translation. We also performed RNA footprinting showing the precise binding site of OmrA on virF mRNA. These data suggest that OmrA might interfere with the translation of the virF mRNA, reducing the amount of VirF₂₁ peptide. The result of this complex regulation is a fine-tuned expression of virulence genes, allowing Shigella a prompt adaptation to different environments and conditions.
Components of the fungal cell wall that turn on and turn off the immune inflammatory response

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The fungal cell wall defines the perimeter of its own self and a boundary of communication with other organisms with which it interacts. For a fungal pathogen, every component and layer of the cell wall plays a specific role in its interactions with cells of the human host. We have tried to unpeel the functions of cell wall components by combining genetics, biochemistry and immunology to understand how the wall is assembled and how its constituents are recognized by the human immune system. This work has shown how the polysaccharides of the cell wall activate both the inflammatory and anti-inflammatory responses of the innate immune system.

Chitin is an essential polysaccharide of the walls of all fungal pathogens and the exoskeleton and eggs of invertebrate parasites. We identified NOD2, TLR9 and the mannose receptor as three essential fungal chitin-recognition receptors of innate immune cells and showed that the activation of NOD2/TLR9 receptors by low concentrations of small chitin particles, generated via the action of human chitinases, leads to selective secretion of the anti-inflammatory cytokine IL-10. In contrast, higher concentrations of chitin particles promoted a pro-inflammatory response via dectin-1-TLR2 signalling. NOD2 and TLR9 polymorphisms are associated with susceptibility to inflammatory conditions such as Crohn’s disease, allergy and asthma. Chitin therefore promotes size-dependent anti- and pro-inflammatory immune responses that are critical for immune homeostasis, but can also promote the development of asthma and allergy.

References:

Fungal human pathogens

Pathogenicity and immune evasion of the human-pathogenic fungus
*Aspergillus fumigatus*

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Background

The filamentous fungus *Aspergillus fumigatus* is the most important air-borne fungal pathogen causing 90% of all systemic *Aspergillus* infections in humans. The most severe disease caused by *A. fumigatus* is invasive aspergillosis which almost exclusively occurs in immunocompromised patients. A lack of reliable diagnostic tools and effective treatment options results in high mortality rates despite therapy.

Objectives

We aim at identifying pathogenicity determinants and mechanisms how *A. fumigatus* can overcome the response of immune effector cells.

Methods

Transcriptome, (immune)proteome analyses including systems biological analyses with subsequent generation of deletion mutants and their analysis in processing by the immune system like neutrophilic granulocytes, alveolar macrophages and complement; test of mutants in a mouse infection model.

Conclusions

*A. fumigatus* has developed immune evasion mechanisms which interfere at the different levels of the infection process with the response of the human host. These include recognition of conidia, modulation of phagocytosis, intracellular processing, neutrophil extracellular trap formation, and complement activation. We have identified several molecules including surface components such as proteins and dihydroxynaphthalene melanin as well as secondary metabolites, which manipulate the immune response.

References


FIRST METABOLIC NETWORK OF THE SEBORRHEIC DERMATITIS ASSOCIATED YEAST MALASSEZIA FURFUR
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Background
Yeasts in the genus Malassezia are part of the skin microbiota of humans and other mammals. These commensal yeasts are related to several disorders including dandruff, seborrheic dermatitis and pityriasis versicolor. However, little is known about the pathogenesis process of Malassezia and its metabolism in these skin diseases, especially at the molecular level.

Objectives
In this study, we aim to describe the characteristics of Malassezia furfur genome-scale metabolic network

Methods
For this, a bioinformatics annotation methodology and a (LC/MS)-based shotgun proteomics profiling were used to characterize the 15.7 Mb genome and the subsequent assignment of the reactions and metabolites.

Conclusions
A total of 757 proteins were found in the proteomic profiling and were used with previous reports to annotate the genome, achieving a first draft of the metabolic network of this yeast. Metabolic reconstruction predicts an interesting degeneracy on the objective function, reflecting several mechanism for the yeast survival as we identified a relatively low number of genes related to lipid biosynthesis but several genes encoding lipid-hydrolyzing enzymes. These enzymes that could have a role in the pathogenesis of the yeast.
FEMS-0722
Fungal human pathogens

IMMUNIZATION WITH ANTIGEN DISPLAYED MICROBIAL CELLS BY MOLECULAR DISPLAY TECHNOLOGY AGAINST CANDIDIASIS

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Background
The administration of caspofungin, micafungin, anidulafungin, and amphotericin B are often involved in pharmacotherapy of candidiasis. Unfortunately, mutants of Candida species with reduced susceptibility to these drugs have emerged. Therefore, prevention of infection by vaccination against Candida species is thought to be important complementary method to pharmacotherapy of candidiasis.

Objectives
Candida albicans enolase 1 (Eno1) has been reported a lot as an immunodominant antigen. To develop a convenient and novel type of vaccine against candidiasis, we designed microorganism cells displaying Eno1p as a model antigen on Saccharomyces cerevisiae and Lactobacillus casei surfaces, respectively [1,2].

Methods
The Eno1p-displayed S. cerevisiae and L. casei cells were constructed by molecular display technology [3]. Eno1 protein was genetically fused to each cell wall proteins to be displayed on these surfaces. Delivery of the cells to mice by oral or intranasal administration increased the titers of anti-Eno1p antibodies and prolonged the lives of mice infected with lethal levels of C. albicans [1,2].

Conclusions
In this study, only availability of a genetic sequence that encodes the antigenic proteins is needed to produce oral vaccine based on molecular display technology. Because a vaccine produced using molecular display technology avoids the need for protein purification, this oral vaccine offers a promising alternative to the use of conventional vaccines for preventing a range of infectious diseases.

Background

Nasopharyngeal carcinoma (NPC) is endemic in Southern China. Intensity-modulated radiation therapy (IMRT) for NPC may damage the salivary glands and lead to xerostomia, providing an ideal environment for opportunistic candidial infections.

Objectives

We prospectively investigated the incidence and factors correlating with oral candidial infection in NPC patients treated with radical IMRT.

Methods

Forty five patients with non-metastatic NPC were prospectively recruited. Saliva and mouthrinse samples were saved at baseline and then two-weekly during IMRT. They all received IMRT with 70Gy in 33 to 35 fractions over 7 weeks concurrent with cisplatin chemotherapy. Besides the gross tumour and positive neck nodes, the parotids, submandibular and sublingual glands were contoured on Eclipse Treatment Planning System™ for radiation dose calculation. Fungal culture was performed on Sabouraud dextrose agar and CHROMagar™. In addition, API-32C AUX method and species-specific Taqman probes were used for specific identification of Candida species. Considerable number of patients i.e. 9 (20.0%) and 25 (55.6%) patients had clinical diagnosis of oral candidiasis and positive candidial culture respectively during IMRT. C. parapsilosis (24 patients, 53.5%) followed by C. albicans (18 patients, 40.0%) were most commonly found. Grade 2 and 3 xerostomia were noted in 13 (28.9%) and 32 (71.1%) patients respectively. Univariable and multivariable analysis
revealed that mean radiation dose to the parotids $\geq 45$ Gy was the only factor ($p=0.045$ and $p=0.043$ respectively) correlating significantly with occurrence of oral candidial infection.

Conclusions

Present study demonstrated that IMRT to the parotids may predispose to oral candidial infections which should be taken into account when managing NPC patients.
Fungal human pathogens

RAPID DETECTION OF CYP51A-PROMOTER BASED VORICONAZOLE RESISTANCE IN ASPERGILLUS FUMIGATUS ISOLATES IN A HIGH INCIDENCE POPULATION

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Background

Due to increased azole resistance, first line treatment with voriconazole of Aspergillus fumigatus infections in humans is increasingly difficult. Susceptibility testing is essential but culture based resistance determination takes up to two weeks. Resistance is predominately based on mutations in cyp51A and duplications of the promoter region (TR34 & TR46).

Objectives

To develop a rapid molecular method to screen for azole resistance in A. fumigatus.

Methods

Azole susceptibility profiles and cyp51A genotypes were determined in 105 isolates of 105 consecutive high-risk patients. A probe-based multiplex qPCR was designed and evaluated. This qPCR specifically amplifies the promoter region of cyp51A and uses a cyp51A control probe and probes for the TR34 and TR46 voriconazole resistance associated promoter duplications.

Conclusions

24/105 (22.9\%) of the patient isolates had elevated voriconazole MICs. Sequencing of cyp51A showed that in 18/24 (75\%) isolates the resistance was associated with mutations in the cyp51A ORF and had a tandem repeat duplication in the promoter region, identifying this as the main resistance mechanism. The qPCR detected 18 of the 18 isolates in which ORF mutations and promoter duplication were present. The 6 resistant isolates that lacked duplications in the promoter region, as well as all sensitive strains were not detected by the PCR. The probe-based multiplex qPCR delivers a rapid method for the identification of the majority of azole resistant A. fumigatus isolates, and offers a rapid pre-screen for detection of voriconazole resistance in a routine clinical routine setting.
Quorum quenching acylases for reducing bacterial virulence

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The observation that acylases from the NTN-hydrolase family can cleave the amide bond between acyl side chains and the homoserine lacton moiety of bacterial signalling molecules has raised novel options for interfering with quorum sensing and the onset of virulence. NTN-hydrolases that have initially become known by their paradigm representatives penicillin and cephalosporin acylase appear to be very abundantly encoded in bacterial genomes. Intriguingly the *Pseudomonas aeruginosa* genome contains not less than 4 NTN-hydrolases. The acylase PvdQ is extremely effective in quorum-quenching, hydrolysing bacterial N-acylhomoserine lactones at very low concentrations. Exogenously added PvdQ can reduce the virulence and we have determined its 3D structure. QuiP and HacB are two other acylases hydrolyzing a subset of the signalling molecules of Gram-negatives, C12-HSL and 3-oxo-C12-HSL. The hydrolytic activity of PvdQ is the highest for long-chain AHLs such as 3-oxo-C12-HSL and decreases with shorter acyl chain length, with hardly any activity for C8-HSL. A structure-based approach was applied to evolve this enzyme into an effective quorum quencher of the communication molecule (C8-HSL) of Burkholderia species, opportunistic pathogens commonly isolated together with *P. aeruginosa* from patients suffering from cystic fibrosis. Two point mutations in the active centre of PvdQ were found to increase the activity of this enzyme towards C8-HSL. The combination variant displayed high hydrolytic activity towards C8-HSL when compared to wild-type PvdQ. The effects could be strongly correlated in vivo, as exogenous addition of this mutant led to a significant decrease of autoinducers present in *Burkholderia cenocepacia* cultures, and effectively reduced its virulent phenotype.
Chemical Interception of Bacterial Quorum Sensing: New Languages, New Outcomes

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We are developing chemical tools that attenuate cell-cell communication pathways in bacteria. Many bacteria communicate using small organic molecules and peptides to monitor their population densities in a process called “quorum sensing.” At high cell densities, bacteria use this signaling network to switch from an isolated, nomadic existence to that of a multicellular community. This lifestyle switch is significant; only in groups will pathogenic bacteria turn on virulence pathways and grow into drug-impervious communities called biofilms that are the basis of myriad chronic infections. In turn, certain symbiotic bacteria will only initiate host-beneficial behaviors at high population densities. The molecular mechanisms of these processes are only now being delineated. Our research is broadly focused on the design and chemical synthesis of non-native ligands that can intercept quorum sensing and provide new insights into its role in host/microbe interactions. We have developed a series of efficient synthetic methods that provide us with straightforward access to these ligands. In addition, we have applied both quorum sensing antagonists and agonists in vitro and in vivo to investigate quorum sensing as an anti-infective target. Quorum sensing pathways in Gram-negative and Gram-positive bacteria have been our foci, including LuxI/LuxR-type signaling in Pseudomonas aeruginosa and agr-type signaling in Staphylococcus aureus. This presentation will introduce our research approach and highlight recent results.
AHLEN CYLASE DISRUPTS BIOFILM FORMATION OF NOSOCOMIAL PATHOGEN ACINETOBACTER NOSOCOMIALIS M2

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Background

Acinetobacter is a Gram negative pathogen associated with nosocomial infection and multidrug resistance. Biofilm formation, as one of the most important virulence determinants in Acinetobacter, is regulated by quorum sensing. Disruption of biofilm formation represents an interesting target for antivirulence drug development. One particular species, namely Acinetobacter nosocomialis strain M2 [1] (previously known as A. baumannii) produces a wide range of quorum sensing signaling molecule with 3-OH-C12-HSL as the primary AHL.

Objectives

The aim of this study was to investigate the efficacy of quorum quenching acylase enzymes in reducing biofilm formation of A. nosocomialis M2. Three acylases with different preference of AHLs substrate were used, namely PvdQ [2] (prefer long chain AHLs, such as C10-C14 HSL), PvdQLo146W,Fp24Y [3] (engineered PvdQ, active towards C8) and QqaR [4] (active towards C8-C14 HSL).

Methods

Static biofilm of A. nosocomialis M2 was grown in 96-well plate with or without addition of acylase. The production of biofilm was examined with crystal violet method.

Conclusions

The addition of AHL acylase was shown to disrupt biofilm production of A. nosocomialis M2, up to 50% reduction compare to control. Interestingly, PvdQLo146W,Fp24Y and QqaR are the most effective in biofilm disruption.

References


Background

"Clostridium perfringens" is an anaerobic gram-positive pathogen and causative agent for gas gangrene as well as necrotic enteritis. "C. perfringens" produces several extracellular enzymes and toxins, such as α, θ and κ encoded by plc, pfoA, and colA, respectively. A two component regulatory system (TCS), VirR/VirS, controls production of these enzymes and toxins. Recently, we have identified a 5-membered macrocyclic oligopeptide (CLWFT) as an autoinducer (AIP_{Cp}) of VirR/VirS TCS.

Objectives

The VirR/VirS system is an attractive target for the development of anti-pathogenic chemotherapy of "C. perfringens", because it orchestrates the quorum sensing cascade. To gain effective inhibitor to block the VirR/VirS signal transduction, we begin with drug design and development of peptides antagonist against AIP_{Cp}.

Methods

To develop antagonistic peptides targeting VirR/VirS, each amino acid residue of AIP_{Cp} was substituted to different amino acids and their agonist activity, which induces pfoA transcription in agrBD-null mutant, as well as antagonist activity which inhibits pfoA transcription in wild type strain were examined.

Conclusions

Alanine scanning suggests that whole structure of AIP is involved in agonist activity. Especially, Z-AIP_{Cp}-W3A showed neither agonist nor antagonist activity, while AIP_{Cp}-F4A solely showed weak but significant antagonist activity, suggesting that Trp-3 and Phe-4 are crucial for receptor binding and receptor activation, respectively. Interestingly, exchange of Trp-3 and Phe-4 yielded a potent antagonist with IC_{50} = 0.4 μM. Alternately, Z-AIP_{Cp}-L2A-T5A also showed a potent antagonist activity of IC_{50} = 0.3 μM. High potential of these oligopeptides is suggested that these can be significantly used for therapeutic applications.
NON-SOCIAL ADAPTATION DEFERS A TRAGEDY OF THE COMMONS IN PSEUDOMONAS AERUGINOSA QUORUM SENSING
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Background
In a process termed quorum sensing (QS), the opportunistic bacterial pathogen Pseudomonas aeruginosa uses diffusible signaling molecules to regulate the expression of numerous secreted factors or public goods that are shared within the population. But not all cells respond to QS signals. These social cheaters typically harbor a mutation in the QS receptor gene lasR and exploit the public goods produced by cooperators.

Objectives
Here we show that non-social adaptation under growth conditions that require QS-dependent public goods increases tolerance to cheating and defers a tragedy of the commons.

Methods
The underlying mutation is in the transcriptional repressor gene psdR. This mutation has no effect on public goods expression but instead increases individual fitness by derepressing growth-limiting intracellular metabolism. Even though psdR mutant populations remain susceptible to invasion by isogenic psdR lasR cheaters, they bear a lower cheater-load than do wild-type populations, and they are completely resistant
to invasion by lasR cheaters with functional psdR. Mutations in psdR also sustain growth near wild-type levels when paired with certain partial loss-of-function lasR mutations. Targeted sequencing of multiple evolved isolates revealed that mutations in psdR arise before mutations in lasR, and rapidly sweep through the population.

Conclusions
Our results indicate that a QS-favoring environment can lead to adaptations in non-social, intracellular traits that increase the fitness of cooperating individuals and thereby contribute to population-wide maintenance of QS and associated cooperative behaviors.
QUORUM SENSING-DEPENDENT REGULATION OF RHAMNOLIPIDS IN BURKHOLDERIA GLUMAE IS NUTRITIONALLY CONDITIONAL

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Background
Burkholderia glumae is a phytopathogenic bacterium that uses the LuxR/LuxI-type TolI/TolR Quorum Sensing (QS) system to regulate various functions, including the major virulence determinants toxoflavin and flagella. B. glumae also releases surface-active rhamnolipids which, along with flagella, are required for the social behavior called swarming motility in Pseudomonas aeruginosa and Burkholderia thailandensis.

Objectives
We investigated the role of rhamnolipids in swarming motility and the interplay between QS and nutrition-based regulation of rhamnolipids production in B. glumae.

Methods
Swarming and swimming assays were carried out to test the motility of various mutants. Analyses of rhamnolipids and N-acyl-homoserine lactones were performed by LC-MS/MS. rhlA expression in wild-type and QS-defective tofI strains was monitored using a lux reporter fusion.

Conclusions
We show that QS positively regulates the production of rhamnolipids in B. glumae and these are also necessary for swarming motility in this species. Unexpectedly, the C8-HSL-negative tofI mutant, in contrast with growth in a minimal medium, regains the ability to produce rhamnolipids in a complex nutrient broth medium. Moreover, decreasing nutrient concentrations leads to a major induction in rhlA expression in the wild-type strain. However, rhlA expression in the tofI mutant is induced to a lesser extent by nutrient limitations. Under nutrient-limited conditions, QS mediates rhamnolipids biosynthesis, presumably to promote swarming motility. These results indicate that, in contrast with the situation in P. aeruginosa, QS regulation of rhamnolipids in B. glumae is dispensable and nutritionally conditional.
Jorgen Lehmann and Gerhard Domagk event on Mycobacterium tuberculosis: New insights into the pathogenesis of MTB impacting on the host’s immune response

TB vaccines; on the role of Th1, memory and immunity in the lung?

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TB vaccines; on the role of Th1, memory and immunity in the lung

*Mycobacterium tuberculosis* and humans have co-evolved since the most early human origin and this has allowed the pathogen to adapt and develop a refined set of countermeasures to the human immune response during infection. The result is a pathogen that is rarely cleared by the natural immune response but instead establishes a long-term chronic infection. Although IFN-γ is clearly insufficient for the control of TB in individuals that develop disease, most TB vaccine research have still been focused on immuno-dominant antigens administered in delivery systems that promote strong Th1 responses. Recent breakthroughs in our understanding of requirement for immunological memory and the recruitment and expression of immunity to TB in the lung now provide a framework for reconsidering that strategy.

I will discuss recent data from mouse and non-human primate models of vaccine promoted protective T cells. Our data suggest that instead of boosting Th1 and IFN-γ, supplementing BCG with *T*CM that are insufficiently induced by BCG may be a feasible avenue towards TB vaccines that efficiently protect the TB infected lung. The data from animal models will be compared to the immune profile obtained with the H1/H56 vaccine in ongoing clinical trials.
THE ROLE OF THIOLS IN PROTECTION AGAINST OXIDATIVE STRESS IN MYCOBACTERIA

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Background
Low molecular weight thiols maintain the reducing environment in cells and are responsible for protecting cells against oxidants and toxins. In mycobacteria such as Mycobacterium tuberculosis, the major low molecular weight thiol is mycothiol (MSH), which has similar protective functions as glutathione (GSH), the LMW thiol present in eukaryotes and gram-negative bacteria. Other LMW thiols present in mycobacteria include ergothioneine (ESH), whose sulfur atom is most stable in solution in the thione form, and thioredoxins, small redox proteins containing a dithiol-disulfide active site. The relative contributions of these LMW thiols in protecting mycobacteria against oxidative stress is not clear.

Objectives
The objective of the study was to characterize the mutants to evaluate their role in oxidative stress defense.

Methods
Mycobacterium smegmatis mutants lacking MSH, ESH, and disrupted in thioredoxins and thiol peroxidases were evaluated for sensitivity to oxidative stress. Quantitative real-time PCR was also performed to determine the expression of the thiol associated genes. Also, the levels of thiols, protein carbonylation and lipid oxidation were determined for the mutants.

Conclusions
A majority of the mutants were sensitive to oxidative stress and a significant increase in expression of genes involved in thiol biosynthesis and genes coding for thiol peroxidases occurred on oxidant exposure. Protein carbonylation and lipid oxidation levels were higher in MSH mutants as compared to mutants lacking ESH and thiol peroxidases. Our data suggests that MSH rather than ESH plays an important thiol in protection against oxidative stress and there is redundancy in protection against oxidative stress.
Background
Hypoxic conditions are known to induce a state of dormancy in mycobacterial species. To what extent phage growth is affected in mycobacterial cells exposed to hypoxia is not known.

Objectives
We attempted to investigate this by studying the growth properties of mycobacteriophage D29 in Mycobacterium smegmatis hosts exposed to hypoxia.

Methods
*M. smegmatis* cells were subjected to hypoxia either by limiting O$_2$ supply (Wayne Model) or by treatment with vitamin C, an O$_2$ scavenger. Treated cells were then infected with D29 and phage growth was monitored by plaque assays.

Conclusions
1. The results indicate that the phage grows poorly under hypoxia but its proliferation is enhanced during the recovery phase.

2. Hypoxia sensors DevR-DevS/DosT may have a role to play in the process as DevR deletion mutant did not support the observed increase in phage growth.

3. The growth properties of the *wt* and *devR* *M. smegmatis* strains were compared in presence and absence of O$_2$ and was observed that the growth of *wt* cells was retarded by hypoxic conditions but the *devR* strain was unaffected. So, DevR plays an active role in retarding the growth of Mycobacterial cells when they are exposed to hypoxia.

4. Following the return to normoxia the retarded cells were found to resume growth in synchrony.

5. Synchronous growth was found to benefit phage development because most of the cells were in the same metabolic state. D29 thus indirectly uses DevR-DevS signalling system to its own advantage when their hosts were exposed to hypoxia.
MSMEG_2433 OF MYCOBACTERIUM SMEGMATIS EXHIBITS BOTH DD-CARBOXYPEPTIDASE AND BETA-LACTAMASE ACTIVITIES

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Background

DD-carboxypeptidases (DD-CPases) are low-molecular mass (LMM) penicillin-binding proteins (PBPs), mainly involved in the peptidoglycan remodelling, but little is known about the DD-CPases of mycobacteria. In this study, a putative DD-CPase of *Mycobacterium smegmatis*, MSMEG_2433 has been characterized.

Objectives

*Physiological characterization of the membrane bound form of MSMEG_2433
*Biochemical characterization of the soluble form of MSMEG_2433

Methods

The gene for the membrane-bound form of *MSMEG_2433* was cloned and expressed in *E. coli* in its active form as revealed by its ability to bind to the Bocillin-FL (fluorescent penicillin). Interestingly, *in vivo* expression of MSMEG_2433 could restore the cell-shape oddities of the septuple PBP mutant of *E. coli*, which was a prominent physiological character of DD-CPases. Moreover, expression of MSMEG_2433 *in trans*, elevated the beta-lactam resistance in the PBP deletion mutants (ΔdacAdacC) of *E. coli* that strengthened its physiology as a DD-CPase. To confirm the biochemical reason behind such physiological behaviours, soluble form of MSMEG_2433 (sMSMEG_2433) was created, expressed and purified. In agreement to the observed physiological phenomena, the sMSMEG_2433 exhibited the DD-CPase activity against artificial and peptidoglycan-mimetic DD-CPase substrate. The enzymatic analyses of MSMEG_2433 revealed the efficient deacylation for beta-lactam substrates, which is unique for beta-lactamases. In addition, *in silico* molecular analyses predicted the presence of an active-site that favours DD-CPase activity of MSMEG_2433 and an omega-loop like region that determines its beta-lactamase activity.

Conclusions

Based on the *in vitro*, *in vivo* and *in silico* studies we conclude that MSMEG_2433 is a dual enzyme, possessing both DD-CPase and beta-lactamase activities.
FEMS-2852
The diversity of bacterial nanomachines

Unfolded protein transport across membranes: The Injectisome in Action
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We are focusing on two fundamental aspects in biology (1) assembly of multicomponent systems into functional nanomachines and (2) transport of proteins across membranes. Both are found in type three secretion systems (T3SS), which are required of bacterial secretion of proteins into the host that subvert the host antibacterial response in order to promote infection. These systems utilize a 3.5 MDa syringe-like, membrane embedded “injectisome” containing an ~800 Å long needle complex thought to cross multiple membranes to connect the intracellular compartments of the bacteria and the host. The striking resemblance of the injectisome to a bona fide syringe, revealed by electron microscopy has led to the hypothesis that bacterial proteins are secreted through the needle complex into the host. I will discuss how and where substrates are passed through the T3SS and across two membranes by visualizing injectisomes in action and in situ.

(www.marlovitslab.org)
The diversity of bacterial nanomachines

A new class of rotary complexes promote motility and sporulation in bacteria

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Bacteria move over solid surfaces by so-called gliding motility, a process whereby the bacterium moves smoothly along its long axis in absence of extracellular organelles. Due to the small size of the bacterial cell the gliding mechanism has long remained enigmatic. In recent years, we identified the Myxococcus xanthus motility engine and showed that it localizes to discrete sites along the cell long axis. However, the exact propulsion mechanism remained undetermined. In this presentation, I will present the motility machinery and recent work combining genetics, biophysics and modeling that allowed us to elucidate the motility mechanism.
Background

Many bacterial species possess appendages that allow them to move while staying attached to the surface. One of the most ubiquitous cell appendages that mediate movement is the type IV pilus system.

Type IV pilus dependent movement has been known as ‘twitching motility’ for a long time, but only recently we were able to demonstrate experimentally that Neisseria gonorrhoeae moves with directional persistence und proposed that its multiple pili are coordinated through a tug-of-war mechanism. [1]

Objectives

We addressed the question whether the pilus dynamics was coordinated mechanically and thereby promoting persistent movement, a mechanism that has been shown to be efficient in other processes.

Methods

We used a combined theoretical and experimental approach to test this hypothesis. Using laser tweezers we measured detachment rates and pilus retraction speeds at the level of single pili. Using this quantitative data we build a tug-of-war model.

Conclusions

Unexpectedly, we found that the two-dimensional tug-of-war is less efficient in generating persistent motion than the one-dimensional tug-of-war studied previously in the context of cytoskeletal transport. [2] The systematic comparison of the patterns of motility with the corresponding experimental data showed that a form of directional memory is required to explain experimental observations. Experimentally, we confirmed memory in the form of bursts of pilus retraction.

Overall, type IV pili are essential for two opposing phenotypes in N. gonorrhoeae, namely motility and clustering into microcolonies and biofilms. Directional persistence is likely to bias the behaviour towards motility.
The diversity of bacterial nanomachines

MICROCYSTIS AERUGINOSA NANOWIRES: AN ELECTRIFYING TALE

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Background

Microorganisms are known to produce extracellular, proteinaceous nanofilaments called type IV pili (TFP) which are known to be involved in different functions like twitching motility, adhesion, biofilm formation etc. These TFP from Geobacter sulfurreducens were found to be electrically conductive and termed as microbial nanowires.

Objectives

As early studies on microbial nanowires focussed on anaerobic microorganisms, this study was undertaken to identify, optimize and structurally characterize microbial nanowires in Microcystis aeruginosa.

Methods

For electrical characterization, conductive atomic force microscopy (CAFM) was used while for structural characterization, transmission electron microscopy (TEM) and in-silico methods were employed. For optimal production of pili, cells were subjected to different stressful culture conditions i.e. carbon source limitation and high light intensity and their piliation level were qualitatively checked using TEM.

Conclusions

TFP in M. aeruginosa was observed to be 8.5-11nm in diameter and 2μ to several μ long. They seem to be either composed of two distinct subfilaments or may be hollow cylinders. The piliation level in M. aeruginosa was sensitive to the culture conditions used. CAFM study showed that TFP in M. aeruginosa are electrically conductive in nature and can act as a microbial nanowire. in-silico studies suggested that aromatic amino acids may be involved in electron transfer through these TFP. The role of such nanowires in toxic bloom formation needs to be investigated further. This study significantly increases our understanding about aerobic cyanobacterial nanowires and also gives impetus for further research in this field and potential environmental applications.
Background

Salmonella swims through liquid environments by rotating a helical organelle, the flagellum. This flagellum is closely related to virulence-associated injectisome systems. A type-III protein secretion (T3S) apparatus is utilized to secrete building blocks of the flagellum, structural subunits of the injectisome complex, as well as virulence effectors into host cells.

Objectives

The T3S apparatus utilizes both the energy of the proton motive force (PMF) and ATP hydrolysis to energize substrate unfolding and translocation. However, the role of the T3S ATPase and other export apparatus components remained unclear.

Methods

We used bacterial genetics, fluorescent microscopy and reporter secretion assays to probe the function of the T3S apparatus in vivo.

Conclusions

We report formation of flagella in the absence of T3S ATPase activity by mutations that increased the PMF and flagellar substrate levels. We additionally show that increased PMF bypassed the requirement of the Spi1 virulence-associated T3S ATPase for secretion. We thus demonstrate that the actual export process is energized by the means of the proton motive force and does not require ATP hydrolysis.

Finally, we performed a genetic screen to dissect the minimally essential components of the flagellar T3S apparatus. We show that most integral-membrane components are essential and all cytoplasmic components are dispensable for export, emphasizing the importance of the core inner-membrane export apparatus components for the function of type-III secretion systems.

In summary, our data support a role for T3S ATPases in enhancing secretion efficiency under limited secretion substrate concentrations and reveal the essential core components of the T3S apparatus.
FEMS-1990
The diversity of bacterial nanomachines

THE STRUCTURE OF THE TYPE SIX SECRETION SYSTEM CONTRACTILE SHEATH SOLVED BY CRYO ELECTRON MICROSCOPY
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Background
Bacteria use rapid contraction of a long sheath of the Type VI secretion system to deliver effectors into a target cell.

Objectives
The mechanism of contraction is still unclear, mostly due to a lack of a high-resolution structure of the sheath.

Methods
We used cryo-electron microscopy and helical reconstruction to solve an atomic resolution structure of a native Vibrio cholerae sheath. Fluorescence microscopy and bacterial competition assays confirm central findings.

Conclusions
The localization of the outer domain of the sheath facilitates unfolding by ClpV. Structural alignments with phage sheaths indicate a conserved mechanism of interaction. In the innermost layer, a 4-stranded β-sheet from three proteins links the protomers in the same strand as well as neighboring strands. This β-sheet is crucial for polymerization and stability.
New insights in syntrophy in methanogenic communities

In methanogenic environments, syntrophic communities form the core of the organic acid degrading population. Organic acids are only efficiently converted if the syntrophs maintain a close relationship, driven by interspecies electron transfer. The eco-physiology of syntrophs is poorly understood as only a few syntrophs have been thoroughly characterized. To clarify their eco-physiology, it is important to study syntrophs in situ but also understand the biochemical mechanisms they have to adapt their metabolism to changing environmental conditions.

Based on genome analysis of our model syntrophic propionate-degrading community *Syntrophobacter fumaroxidans* and *Methanospirillum hungatei* and expression of genes involved in interspecies electron transfer we revealed mechanisms that enable the coculture to adapt\(^1\).

Single-cell microbiology enables microbiologists to observe who does what, when, where and next to whom\(^2\). Single cells taken from the environment can be identified and their genome sequenced. Single-cell based gene expression analysis could be a valuable tool in interpreting gene functions and microbial metabolism in various syntrophic communities and reveal genes and gene clusters that are involved in syntrophy\(^3\).

This presentation will discuss syntrophy and methanogenic syntrophic communities and emphasize on recent developments and discoveries.

References


Syntrophic interactions in anaerobic communities

Studying microbial interactions using electrochemical tools

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Microbial collaboration for effective electron transfer

Research on extracellular electron transfer has accelerated in the past ten years. Whereas the process itself was described in 1986, only in the 2000s major interest emerged leading to the discovery of direct electron transfer to and from electrodes, and more recently the discovery that pili or pilus-like appendages are involved in long range electron transfer. The electrical interactions are not only occurring between a microorganism and an electrode or mineral, it is now clear that microorganisms interact among each other via electrons. It is likely that they do so to link their overall respiration, settling charge balances via solid state transfer rather than relying on diffusive processes. Taking this information on board, earlier findings on the microbial ecology of electroactive biofilms make more sense and it appears that in anaerobic environments the physiology related to electron transfer needs to be included when considering aspects such as syntrophic interactions. In my presentation, I will give an overview of the findings by many researchers worldwide, and try to assess how this impacts particular applications such as microbial electrosynthesis, microbial electrochemical systems and anaerobic digestion processes.
FEMS-1988
Syntrophic interactions in anaerobic communities

PROTEOMICS OF SYNTROPHOMONAS ZEHNDERI AND METHANOBACTERIUM FORMICICUM GROWING ON LONG-CHAIN FATTY ACIDS
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Background
Conversion of long-chain fatty acids (LCFA) in anaerobic digesters relies on syntrophic relationship between acetogenic bacteria and methanogenic archaea. Conversion of unsaturated- and saturated-LCFA has been previously shown by a co-culture of Syntrophomonas zehnderi and Methanobacterium formicium. Degradation of unsaturated-LCFA is rare among Syntrophomonas species; the best studied fatty-acid oxidizer, S. wolfei, can only grow on saturated-LCFA.

Objectives
Major differences are expected in the pathways and enzymes involved in the degradation of unsaturated-LCFA. In this work we used proteogenomic approach to study these differences.

Methods
A draft genome of S. zehnderi was obtained by IlluminaHiSeq sequencing. Genomes of S. zehnderi and S. wolfei (available at NCBI) were compared. S. zehnderi and M. formicicum co-cultures grown on oleate (unsaturated LCFA, C18:1) and on stearate (saturated LCFA, C18:0) were further studied using a proteomics approach.

Conclusions
Genomic comparison of S. zehnderi and S. wolfei revealed approximately 900 different proteins and 1200 common proteins. In the genome of S. zehnderi, two replicates of the unsaturated acyl-CoA dehydrogenase genes were identified, one of which differs considerably from the acyl-CoA gene found in S. wolfei. Proteomic analysis of S. zehnderi and M. formicicum co-cultures revealed high expression levels of proteins related to the β-oxidation of LCFA (up to 30% of total proteins identified). Different protein expression levels were observed during the degradation of oleate (44% unique proteins) and stearate (23% unique proteins). In addition, proteins involved in electron transfer were highly expressed, including electron transfer flavoproteins, ATP synthases and a number of hydrogenases and formate dehydrogenases.
SYNTROPHIC COCULTURE OF GEOBACTER SULFURREDUCTENS AND SYNTROPHOBACTER FUMAROXIDANS GROWING WITH PROPIONATE AND Fe(III)

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Background
The Fe(III)-reducing G. sulfurreducens is restricted to simple organic substrates like acetate as the electron donor and unable to metabolise propionate. S. fumaroxidans on the other hand cannot use Fe(III) as electron acceptor.

Objectives
Syntrophic coculture of the electroactive bacterium Geobacter sulfurreducens and the propionate oxidising bacterium Syntrophobacter fumaroxidans with propionate and Fe(III) were constructed.

Methods
G. sulfurreducens was first grown in pure culture with single and different combinations of the electron donors of acetate, formate and hydrogen as the products of propionate oxidation by S. fumaroxidans. All single electron donors supported the growth of G. sulfurreducens where 6.44 ± 0.02, 2.43 ± 0.01 and 1.63 ± 0.41 mole of Fe(II) was produced per mole of acetate, formate and hydrogen, respectively. When different combination of electron donors applied, no acetate consumption was noted in the presence of formate. Moreover, in the presence of hydrogen, reduced amount of acetate/formate was needed for Fe(III) reduction compared with similar conditions without hydrogen. Next, our coculture results showed for the first time the successful growth of these two bacteria with propionate and Fe(III) in batch cultures, where neither of bacteria was independently capable of such growth.

Conclusions
The exceptionally high ratio of Fe(II) production per propionate consumption (17.3 ± 2.52) showed high potential of the applied coculture for iron reduction using propionate as the electron donor. Further molecular and microscopic studies are underway to illustrate the spatial structure of the coculture and the abundance of the two partner bacteria.
SYNTROPHIC ETHANOL OXIDATION - A LONG WAY TO A CONVINCING SOLUTION
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Background
Methanogenic conversion of ethanol to acetate plus methane has been described by Wolin et al. in 1967 as the first case of syntrophic cooperation between two species in methanogenic degradation. In the co-culture “Methanobacillus omelianskii”, the so-called S-organism cooperated with a hydrogen-oxidizing methanogen. The oxidation pathway as conceptualized with this culture included alcohol dehydrogenase, acetylating acetaldehyde dehydrogenase, and subsequent ATP formation via phosphotransacetylase and acetate kinase. This concept has been confirmed later with Pelobacter acetylenicus and other Pelobacter species.

Objectives
Pelobacter spp. have to invest part of their ATP gain into a reversed electron transport to release electrons as hydrogen or formate. The overall energy balance leaves less than one ATP per reaction run for these bacteria. We wanted to elucidate the mechanism of electron flow from ethanol to hydrogen.

Methods
Studies included enzyme assays in cell-free extracts as well as genomic and proteomic analyses.

Conclusions
Genomic studies indicated that Pelobacter carbinolicus contains a comproportionating hydrogenase oxidizing NADH and ferredoxin simultaneously. Methanogenic ethanol-oxidizing cocultures maintained formate and hydrogen levels equivalent to a standard redox potential of E° = -360 mV. Combined with further genomic information, our results provide a convincing concept for electron flow and ATP synthesis in syntrophically ethanol-oxidizing Pelobacter species.
Background

Mixed microbial communities are complex and dynamic systems. Integrated omics (combined metagenomics, metatranscriptomics, metaproteomics and metabolomics) are currently gaining momentum for detailed descriptions of community structure, function and dynamics in situ. These top-down analyses also allow inference of fundamental ecological properties of constituent populations and allow a description of the flow of matter through these systems.

Objectives

To define niche breadths of constituent populations and to infer the flow of matter through a mixed microbial community from integrated omics data.

Methods

We have developed an integrative workflow comprising wet- and dry-lab methodologies that enables systematic measurements of microbial communities, and the integration and analysis of the resulting “multi-meta-omic” data including in the context of reconstructed metabolic networks. Here we apply this methodological framework to our model microbial communities (lipid accumulating microbial consortia from a biological wastewater treatment tank) sampled over space and time.

Conclusions

By resolving multi-omic data at the population-level, we are able to infer lifestyle strategies (generalists versus specialists) of the respective constituent populations. More specifically, through our analyses, we have uncovered patterns which suggest that, under rapidly fluctuating environmental conditions, the dominance of a microbial generalist is linked to finely tuned resource usage, and that genetic variation within generalist populations is constrained. Furthermore, by reconstructing population-resolved metabolic networks and interfacing there, we are able to partially reconstruct the flow of matter through the community. These results bring us closer to being able
to infer key ecological characteristics of constituent microbial populations as well as matter and ultimately energy flows through communities.
Type IV secretion (T4S) systems are molecular machines used for the transport of macromolecules across the bacterial cell envelope. T4S systems are highly versatile. Conjugative T4S systems translocate DNA from a donor to a recipient bacterium and contribute to bacterial genome plasticity, spread of antibiotic resistance or other virulence trait among bacterial pathogens. In some bacteria such as Helicobacter pylori (Cag PI), Brucella suis (VirB/D), or Legionella pneumophila (Dot, Icm), T4S systems are directly involved in pathogenicity as they mediate the secretion of virulence factors (DNA or toxins) into host cells. The archetypal T4S system, the VirB/D system, was defined in Agrobacterium tumefaciens where it is naturally responsible for the delivery of the T-DNA to the plant host-cell. The A. tumefaciens VirB/D system comprises 12 proteins (VirB1 to 11 and VirD4). Recently, structures of large complexes formed by several of these proteins have become available shedding unprecedented light on T4S system secretion mechanism.

Key recent references


On the role of photosensory receptors in (chemo) trophic bacteria

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Oxygenic photosynthesis and its applications in (energy) biotechnology will have to have a key role in a sustainable future. It is therefore significant that this process can be straightforwardly engineered in oxygenic phototrophic organisms, such as cyanobacteria, using the tools of synthetic biology. This has led to the emergence of cell factories that catalyze the (sun)light-driven conversion of CO$_2$ and water into products like ethanol, butanol, or other bio-fuels, commodity chemicals like lactic acid, a bio-plastic precursor, and even artificial sweeteners and flavor compounds. Only oxygen is produced as a by-product. For successful application of this approach it is of key importance to optimize such cell factories to maximal efficiency. This holds for their light-harvesting capabilities, e.g. under circadian illumination in large-scale photo-bioreactors, but equally well for the ‘dark’ reactions of photosynthesis, i.e. the conversion of CO$_2$, NADPH and ATP into a (biofuel) product.

After an introduction of the versatility of this approach, an analysis based on metabolic control theory is presented of the amount of the enzymes catalyzing the heterologous metabolic pathway for product formation with which such cyanobacterial cell factories have to be equipped for maximal rates of product formation. For this engineered L-lactic acid producing *Synechocystis* sp. PCC6803 strains are used to identify the relation between actual production rate and expressed enzyme capacity. The analysis shows that engineered cell factories (e.g. for L-lactic acid) require high capacity of the heterologous product-forming pathway, before control in other parts of the cell factory can be detected and relief of limitation in those other parts becomes relevant.

**Reference**

Photoprotection in cyanobacteria and the photoactive Orange Carotenoid Protein

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The water-soluble orange carotenoid protein (OCP), a cyanobacterial protein of 35kDa which binds a single keto-carotenoid was first described and isolated by Holt and Krogman in 1981. During long time its role was unknown. In 2006 we demonstrated that OCP is essential in photoprotection in cyanobacteria. Cyanobacteria protect themselves from intense sunlight by dissipating excess absorbed energy as heat at the level of their extramembranal antenna, the phycobilisomes. This decreases the formation of dangerous species of oxygen and diminishes photodamage. The process is induced by the absorbance of blue-green light by the OCP, which acts as the light intensity sensor and the excitation energy quencher and is a new member of the family of photoactive proteins. It is the first photoactive protein containing a carotenoid as the photoactive chromophore and its photocycle is completely different from those of other photoactive proteins. The absorbance of blue-green light by the OCP induces structural changes in the carotenoid and the protein, converting its dark stable orange form (OCP_o) into a relatively unstable active red form (OCP_r). OCP_r form is essential for the induction of the photoprotective mechanism. Only OCP_r is able to interact with phycobilisomes and to quench excess energy. A second protein, the Fluorescence Recovery Protein, is needed to detach the OCP_r from phycobilisomes and converted to the inactive OCP_o form. All the known details of this mechanism will be described in the conference. We have recently demonstrated that OCP has another essential role in the cells: it’s a very good singlet oxygen quencher. Thus, OCP has dual and complementary photoprotective action as energy quencher and as singlet oxygen quencher.
Phytochrome is a multidomain red-light photoreceptor that allows cells to respond to the quality and intensity of incoming light. Bacterial phytochromes adopt a deep figure-of-eight knotted polypeptide. They bind the linear tetrapyrrole biliverdin covalently via a thioether link to cysteine. Photon absorption causes a rotation about the C15=C16 double bond of this chromophore, leading to signaling through the conserved PHY domain to a modular output domain. To advance our research goals of understanding the structure/function relationships of bacterial phytochromes, we are studying the biochemistry and structural biology of the signal transduction pathway of the bacterial phytochromes of *Deinococcus radiodurans* and *Ramilbacter tatouinensis*. Each contains a histidine kinase domain, and each is predicted to be a light-regulated enzyme that phosphorylates across a homodimeric interface and subsequently transfers phosphate to a response regulator protein. We have established a robust *in vitro* kinase assay and have shown that *R. tatouinensis* bacterial phytochrome is an autokinase that interacts specifically with at least one cognate single domain response regulator (Brr). Kinase activity is markedly lower under red light. We have also refined the X-ray crystal structure of the Brr protein. A C-terminal extension of the canonical response regulator fold creates a unique dimer interface not seen outside the phytochrome-associated response regulators. We are engaged in determining the functional consequences of the knotted phytochrome topology and the oligomeric response regulator. We imagine both could limit motions of phytochrome domains upon illumination to steer signal transduction along a productive trajectory.
The human virome: new tools and concepts

Increasing the knowledge of the human virome at homeostasis and disease by deep sequencing of clinical samples

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The composition, interindividual and temporal variability of the human virome are not precisely known but new questions arise in the context of the rapidly increasing knowledge regarding its composition and function. Its impact on human health has received less attention than that of the bacterial microbiome, but is likely to be as important, both in homeostasis and disease. In fact, it is increasingly apparent that viral pathogens are only the salient members of a larger group of viruses associated with humans that are not directly linked to disease, analogous to the bacterial component of the microbiome. In particular, viruses abound at the body interfaces with the external environment, on the skin and in the mucosa. In contrast to this, the presence of viruses in the blood and organ parenchyma of healthy people is rather the exception than the rule, although a limited number of viral species establish persistent, albeit mostly silent, systemic infections, more often in immunocompromized individuals. With the ever-extending use of next generation, rapid progress on its impact on human health is to be expected in the coming years including the identification of viruses with roles in chronic diseases. We will show how deep sequencing of human clinical cases samples reveals, rather than new viral species, an unexpected role of known viruses in human diseases particularly in immunocompromised patients. On the other hand, whilst cross transmission from animals is preeminent in emerging viral diseases, close or identical new viral species are found in human and animals at homeostasis, which suggests more circulation between species than anticipated.
Microbes drive the biogeochemical cycles that fuel the planet, and their viruses play important roles ranging from mortality and nutrient cycling to gene transfer in environments ranging from the oceans and soils to industrial fermenters and humans. However, our inability to “see” viruses in the environment challenges our broader and quantitative understanding of viral roles in ecosystem processes. My lab specializes in developing new experimental and informatic approaches to better “see” and interpret the diversity of wild viruses across myriad data types and environments. Together these methods link viruses to their hosts en masse, provide both single-cell and population perspectives on virus-host interactions, or help elucidate Viral Dark Matter that dominates surveys of viruses in nature. In natural communities, we have mapped much of the global virome, shown that viral genome sequence space is clustered (into ‘populations’ formally delineated as species), and provided hypotheses on the role of viral metabolic reprogramming in modulating ecosystem function. This ability to now ‘count’ biologically meaningful units for viruses in nature – both genes and ‘organisms’ – opens watershed possibilities for applying ecological and evolutionary theory and developing predictive modeling across diverse ecosystems. Beyond the oceans, microbes are emerging as major players in terrestrial and atmospheric biogeochemical and energy cycling, as well as many diseases humans. Viral roles in modulating these microbial impacts are virtually unstudied, but likely to be large. To this end, my lab is now optimizing and applying these novel ocean-derived experimental and ‘big data’ analytical approaches to viral communities across diverse environments including sediments, the human lung and gut, and climate-sensitive thawing permafrosts, with aspirations to explore roles of viruses in the atmosphere and the Built Environment.
Viruses and animal viruses identified through metagenomics: pathogens, commensals, or contaminants?

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Viral metagenomics is the unbiased amplification and sequencing of viral particles-associated nucleic acids and their classification using sequence similarity to known viruses. The research and diagnostic opportunities provided by this approach include the development of universal and sensitive virus identification tools and viral pathogen discovery. Our methodology and bioinformatics pipeline, together with their limitations, will be briefly described. Examples of virus and viral DNA contamination and the detection within mammalian samples of viral sequences originating from diverse cellular hosts will be shown. New viral genomes from human and animal samples with unexplained diseases of suspected infectious origins will be described together with the experiments performed or required to demonstrate their pathogenicity. Some of these newly described viruses are highly prevalent, particularly in infants from developing countries, both sick and healthy, with frequent co-infections. The rapid rate of viral discovery provides many opportunities for epidemiological and in vivo studies to explore the role of these “new” viruses in both health and disease.
FEMS-2864
Why we need to worry about bacterial infections?

Collaborating with the innate immune system to combat multidrug-resistant bacterial pathogens

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Antibiotic resistance presents an ever increasing challenge to the public health with a dearth of new drugs in the development pipeline. Conventional screening paradigms in antibiotic discovery are based on MIC/MBC testing in conventional bacteriologic media, and similar tests on patient isolates are used to guide physician management. Economic factors have favored development of broad spectrum agents, which exert "collateral damage" on the normal microflora, now increasingly recognized to have adverse health consequences. A single-minded focus on direct antimicrobial activities overlooks the fact that significant infections are really a disease of the host-pathogen interaction. Indeed, before the patient has even seen a doctor, their infection is already being treated by multiple antimicrobials - namely the cellular and molecular components of the innate immune system. We see value in exploring potential novel therapeutic approaches for drug-resistant bacteria that aim to tip the host-pathogen interaction back in favor of the host. This talk will illustrate three such classes of novel therapeutics: (A) Inhibitors of bacterial virulence factors that re-sensitize the pathogen to innate immune killing; (B) Drugs that directly boost the antibacterial killing capacity of host phagocytic cells; (C) Antibiotics that synergies with endogenous antimicrobial peptides to effect bacterial killing. These studies will reveal how standard MIC testing can be misleading, and overlook potent antibiotic activities that are recognized only the context of the normal innate immune system. In this new discovery and treatment framework, drugs used in medicine for other indications, or antibiotics otherwise deemed ineffective, can be "repositioned" for treatment of multi-drug resistant pathogens such as methicillin-resistant Staphylococcus aureus (MRSA), vancomycin-resistant Enterococcus spp. (VRE) or carbapenemase-resistant strains of Gram-negative pathogens including Acinetobacter baumannii, Klebsiella pneumoniae and Pseudomonas aeruginosa.
FEMS-2343
Why we need to worry about bacterial infections?

Meningococemia a disease of the endothelial cells

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*Neisseria meningitidis*, (also known as the meningococcus), is a common inhabitant of the human nasopharynx, and as such is a normal, saprophytic organism that is transmitted from person to person. These bacteria can therefore spread in a population by direct contact. Paradoxically, in a small proportion of colonized subjects, they disseminate from the nasopharynx into the bloodstream, leading to clinical features, which are rarely observed in bacterial septicemia. First, meningococci usually cross the blood brain barrier, invade the meninges and cause meningitis. Secondly meningococcemia is usually associated with a thrombotic syndrome, which causes limited skin purpuric lesions, in addition to the meningeal inflammation in its milder form, or extensive thrombosis and necrotic purpura associated with massive vascular leakage and shock, known as *purpura fulminans*, in its more severe form. The epidemic nature of meningococcal transmission and the high mortality rate, especially in case of *purpura fulminans*, are responsible for the fear of meningococcal infections in communities. *In vivo* observations have clearly shown that *Neisseria meningitidis*, once in the bloodstream, interact closely with the microvessels. This interaction is mediated by the type IV pili. The mechanisms and consequences of this interaction will be analyzed in the light of the clinical consequences.
Buruli ulcer is an infectious disease transmitted by arthropod vectors harboring *Mycobacterium ulcerans*, a mycobacterium which belongs to the same family of bacteria causing tuberculosis and leprosy. The infection causes painless swelling and severe skin lesions. One key feature of *M. ulcerans* bacterium is its ability to secrete a necrotic toxin, the mycolactone within small lipophilic vesicles, which are critical for the bacterial induced pathogenicity. The biological knowledge as well as the preventive and therapeutic means for this invalidating disease is still very limited.

The extensive skin lesions, despite their severity, are not accompanied by pain. It was previously thought that this remarkable analgesia is ensured by direct nerve cell destruction. We demonstrate that *M. ulcerans*-induced hypoesthesia is instead achieved through a specific neurological pathway triggered by the secreted mycobacterial polyketide mycolactone, leading to potassium-dependent hyperpolarization of neurons. By use of a combination of siRNA and chemical library image based screening, we showed that mycolactone elicits signaling through type 2 angiotensin II receptors (AT2Rs) leading to the activation of cyclooxygenase I and TRAAK potassium channels, which causes neuron desensitization. We further validate the physiological relevance of this mechanism with in vivo studies of pain sensitivity in mice infected with *M. ulcerans*, following the disruption of the identified pathway. Our findings shed new light on molecular mechanisms evolved by natural systems for the induction of very effective analgesia, opening up the prospect of new families of analgesics derived from such systems.
FEMS-2866
Why we need to worry about bacterial infections?

The use of antibiotics drive the evolution of resistance by activating competence for DNA uptake: Molecular mechanisms involved in Streptococcus pneumoniae

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The rapid spread of antibiotic resistance, combined with a near absence of new antibiotics, are leading to a public health threat. One of the leading bacterial causes of morbidity and mortality worldwide is Streptococcus pneumoniae (the pneumococcus). Frighteningly, inappropriate antibiotic treatments can accelerate the occurrence of multidrug resistance by activation of a developmental process called bacterial competence.

In this presentation I will discuss how antibiotics affects pneumococcal physiology and how antibiotics promote competence development. Molecular insights into the mechanisms driving bacterial evolution and resistance will advance the quest for novel treatment strategies.
Bacterial predators

*Bdellovibrio bacteriovorus*- biology of invasive bacterial predation

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*Bdellovibrio bacteriovorus* is a predatory delta-proteobacterium, with a 3.8Mb genome inside a vibroid cell only 1µm x 0.3µm. Locomotion by a single flagellum or by a gliding apparatus delivers *Bdellovibrio* to encounter other Gram-negative cells at their outer membrane. There is an attachment (via the non-flagellar pole) and recognition phase and then invasion by *Bdellovibrio* and intraperiplasmic growth, inside the, now-dead, but intact, prey. Growth in the periplasm requires modification of the prey cell wall to prevent premature lysis as the predatory life history strategy of *Bdellovibrio bacteriovorus* is not to share nutrients with bacteria outside the prey. The nutrients available from the prey are finite and dependent on cell volume and even or odd numbers of *Bdellovibrio* progeny are produced. *Bdellovibrio* motility resumes at the end of predator replication to escape the exhausted prey cell by flagellar or gliding motility. The non flagellar pole is specialised for prey-interaction and this involves Type IV pili and associated regulatory proteins, functions for which are becoming understood. This presentation will review the prey-modification and predatory growth processes of *B. bacteriovorus* and touch on their evolution.
Bacterial predators

When cheese gets the flue: how dairy lactic acid bacteria succumb to bacteriophage attack

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Bacteriophages are a constant threat to commercial dairy plants, where their presence may lead to inconsistent, delayed or even disrupted fermentations. Over the last 50 years bacteriophages infecting lactic acid bacteria, and in particular dairy strains of Lactococcus lactis, have been isolated and studied to learn about their diversity and biology, as well as to find ways by which their negative impact on their host can be reduced or managed. In recent years, a wealth of information has been generated related to lactococcal host recognition by phages with (non-contractile) tails, representing the vast majority of (dairy-associated) phages. This presentation will present data on recent work in this field, highlighting the role of the saccharide-recognizing base plate located at the tip of the lactococcal phage tail, while also discussing the (diversity of the) phospho-polysaccharide receptor, which is recognized by the base plate.
Bacterial predators

Structural Biology as a Tool to Study Predatory Bacteria

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Bdellovibrio bacteriovorus is a predatory bacterium that enters other Gram negatives, residing in their periplasm and killing them from within. The goal of my lab is to understand this process on the molecular level, detailing how predation-specific proteins function by use of structural biology. We are currently investigating cell wall metabolism (entry into prey), cyclic-di-GMP signalling (control of predation events), host-independence (conversion to a form that can grow/replicate outside of the prey cell) and a variety of cryptic genes (Bdellovibrio has many unique genes, forming the “predatosome”). This presentation will provide examples of our progress in these areas and what we can learn about the toolkit required to live a predatory lifestyle.
Because of their outspoken ubiquity and versatility, bacterial viruses (phages) constitute an inevitable and integral part of microbial ecology, and are now being recognized as an important selective pressure and/or advantage that continues to drive microbial evolution and behavior. Moreover, the ongoing evolutionary arms race between bacterial hosts and their viral parasites has endowed phages with fine-tuned strategies to interfere with or even reshape microbial physiology. Understanding bacteriophage infection and the impact it has on the bacterial host greatly depends on insights at the systems biology level. Our research focuses on the phage-based identification of new antibacterial targets and indicates a close interplay at the RNA, proteome (interactions & post translational modifications) and metabolome level of Pseudomonas phages.

Lytic bacteriophages inject their genome in the bacterial cytoplasm and immediately redirect the host metabolism towards phage genome replication and the production of virions. However, the mechanisms behind the molecular hijack are largely unknown. While bacteriophages infecting Pseudomonas aeruginosa have been known and studied for a long time, over 80% of their genes have an unknown function. Recent ‘omics’ techniques can lead to a drastic increase in the understanding of the struggle between virus and host to dominate the cellular metabolism during all stages of infection. The exploration of four levels of control (transcriptional, posttranscriptional and – translational, protein-protein interactions) and their influence on the bacterial metabolome are be the key focus of this presentation. We are currently exploring the potential for biotechnological applications and antimicrobial design strategies of these phage-encoded gene products.
FEMS-2809
The Type VI secretion system

Recent insights into the export and activity of type VI secretion toxins
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Though they are by definition single-celled organisms, bacteria can exhibit properties more often ascribed to multicellular life. Our group aims to identify, characterize, and eventually, exploit, pathways that bacteria use for intercellular communication and competition. In this presentation, I will discuss our recent progress toward understanding the diversity of mechanisms by which antibacterial toxins delivered by the type VI secretion system act upon recipient cells. I will also describe our discovery that a conserved signaling pathway allows cells of the Gram-negative opportunistic pathogen \textit{Pseudomonas aeruginosa} to mount a coordinated response to antagonism by competing bacteria.
Gram-negative bacteria have evolved various strategies to compete in hostile environments. Among them, secretion systems have attracted a lot of attention because of their clinical importance but also because of their complex architecture and regulation. The conserved bacterial type VI secretion system (T6SS) was initially associated with bacteria-host interaction but subsequently found to inject toxins into bacterial targets. These toxins have various activities, the best characterized of which are amidases, phospholipases and DNases.

*Pseudomonas aeruginosa* is an opportunistic pathogen, which has three T6SSs called H1- to H3-T6SS. We used a combination of structural and *in vivo* protein-protein interaction approaches to characterize molecular aspects of the H1-T6SS assembly of *Pseudomonas aeruginosa*. We also characterized several T6SS toxins and studied their mode of delivery and their impact on bacterial competition.
The bacterial Type VI secretion system (T6SS) is a large dynamic organelle that is functionally analogous to contractile tails of bacteriophages. T6SS is used by Gram-negative bacteria to inhibit adjacent cells via translocation of toxic effector proteins and thus play an important role in bacterial ecology. Time-lapse fluorescence light microscopy revealed that T6SS sheath, which powers the secretion, cycles between assembly, quick contraction, disassembly and re-assembly. Single cell analysis of subcellular localization of T6SS assembly revealed that T6SS organelle encoded by Pseudomonas aeruginosa H1-T6SS cluster is assembled and aimed to specifically retaliate against attack by other bacteria. I will present latest update on the structure, function and dynamics of T6SS as well on mechanisms of effector delivery. I will focus on the structure of the T6SS sheath and the implications for T6SS dynamics and assembly.
Germination of spores of *Bacillus subtilis*: nothing but surprises

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Background

Previous work has identified proteins essential for *Bacillus subtilis* spore germination as the: i) germinant receptors (GRs) that recognize nutrient germinants; ii) GerD protein essential for rapid GR-dependent germination; iii) SpoVA proteins that likely comprise a channel for dipicolinic acid (DPA) release early in germination; and iv) lytic enzymes that degrade the peptidoglycan cortex later in germination. Previous work has determined these proteins’ locations in spores, some of their structures and some knowledge of how their function is regulated.

Objectives

Identify and understand the earliest spore germination events.

Methods

To help answer these latter questions we have examined germination of multiple individual *B. subtilis* spores under various conditions.

Conclusions

Surprising recent findings from this work include the following. 1) At approximately when spores become committed to germinate, spores’ inner membrane permeability decreases and DPA begins to leak out, with this leakage more evident at higher temperatures. Following leakage of 10-20% of spores’ DPA, the remainder is released in 1-2 min. The change leading to slow DPA leakage is unknown, but could be changes in spores’ inner membrane, the DPA channel, or both. 2) Spores have memory of exposure to nutrient germinants, such that a 1ˢᵗ germinant pulse results in more germination following a 2ⁿᵈ germinant pulse. This memory is established before commitment, as memory is lost upon extended incubation, especially at elevated temperatures. Memory is formed by stimulation of any GR and can be accessed by stimulation of any GR. However, how spore memory is generated, stored and accessed are not known.
A general problem in developmental biology concerns the process by which cells of one type divide to give dissimilar daughter cells. Even though these daughter cells may be genetically identical, they can differ morphologically and physiologically and have different fates. As one of the simplest differentiation processes, *Bacillus subtilis* sporulation represents an excellent model system for studying cell differentiation. Several decades of study of this process have provided insight into cell cycle regulation and development. The first clear morphological step in sporulation is formation of division septum at the polar site, which is fundamental to the development of highly resistant endospores formed by bacterial species as Bacilli and Clostridia. The earliest visible event in this asymmetric cell division is the formation of a Z-ring by FtsZ, a tubulin like protein, and E-ring by SpoIIE at the future sporulation septum site.

There are many important advances in our understanding of asymmetric gene expression during spore formation with an emphasis on developmental stages that lead to asymmetric septum formation and especially to activation of the first compartment specific sigma factor - sigF. It is clear that SpoIIE is a crucial constituent of the sporulation septum formation and an import determinant of the proper activation of sigF. However, we understand these processes only partially and there are few fascinating unanswered questions as: how SpoIIE localises to the polar septum; how SpoIIE causes FtsZ to relocalise from mid-cell to the polar site; how SpoIIE’s phosphatase activity is controlled so that sigF activation is delayed until the septum is completed; what role SpoIIE plays in septal thinning or what is the role of other proteins and Min system in this stage of sporulation.
The role of spores in C. difficile infection

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Clostridium difficile infection (CDI) is an important nosocomial infection in the developed world and one for which no vaccine currently exists. In the USA alone about 28,000 people die each year and of these most occur from nosocomial infection. It is universally accepted that extensive antibiotic usage promotes germination of C. difficile spores in the GI-tract followed by secretion of toxins (A and B) from the live bacterium that then produce symptoms of disease (gastroenteritis and diarrhea). C. difficile is a spore-forming bacterium and both humans and animals can be asymptomatic carriers. It is also accepted that C. difficile can be acquired from the environment and there is evidence of transfer of animal strains (e.g., the 078 ribotype) to humans.

While the vegetative bacterium produces symptoms of disease there is compelling evidence that other virulence factors exist and that the spore plays an important role in infection. For example, oral administration of non-toxin producing spores confer protection to CDI in humans implying that other correlates of infection exist.

This talk will focus on two aspects of CDI. First, the characterization of proteins found on the surface of C. difficile spores and second, strategies we are using to vaccinate against CDI using mucosal delivery of C. difficile antigens and that are now in clinical trial.

Analysis of spore coat proteins has revealed a number of proteins that carry enzymatic properties of which one can be linked to inflammation. A second, BclA1, has been shown to play a distinct role in the initial stages of infection potentially by recognition of a specific receptor molecule in the gut mucosa.
A key event in the germination of spores of *Bacillus* and *Clostridium* species concerns depolymerisation of the thick layer of cortical peptidoglycan that surrounds the spore core. Cortical dissolution is achieved via the activities of a group of peptidoglycan lysins referred to as cortex lytic enzymes (CLEs). CLEs can be further sub-divided into those that cleave intact peptidoglycan, and those that function as cortical-fragment lytic enzymes (CFLEs). The molecular mechanisms that underpin the transition of inactive CLEs and CFLEs in the dormant spore, to catalytically active enzymes during spore germination, are poorly understood. In the current work we have used X-ray crystallography to examine CFLEs from both *Bacillus* (*SleL*) and *Clostridium* (*SleM*) spores in an attempt to gain structural insights to the regulation of CFLE activity. To date, high-resolution structural information has permitted relatively facile identification of the active sites and catalytic residues in the respective enzymes, and has revealed why *SleL* from *B. cereus* spores is significantly more catalytically efficient than the orthologous *B. megaterium* enzyme. Determination of the *C. perfringens* *SleM* crystal structure has additionally revealed unusual structural features that are probably involved in substrate or protein interactions. Finally, comparison of *SleL* and *SleM* crystal structures with the *Bacillus* CLE, *SleB*, has permitted the generation of hypotheses regarding the structural basis for CFLE substrate selectivity, which will be discussed during the presentation.
Small RNA-based antiviral immunity in insects: defense and counter-defense


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The RNA interference (RNAi) pathway senses virus infection to induce an antiviral response in insects. The ribonuclease Dicer-2 detects viral double-stranded RNA and generates viral small interfering RNAs (vsiRNAs), which are then used to restrict viral replication. Recent small RNA profiling studies demonstrated that both RNA and DNA viruses are targets of the RNAi response and identified the viral RNA substrates that trigger vsiRNA biogenesis. The importance of the antiviral RNAi pathway is underscored by the observation that viruses have evolved sophisticated mechanisms to counteract this immune response. More recently, it was proposed that another small RNA silencing mechanism, the piRNA pathway, also processes viral RNAs in Drosophila and mosquitoes. I will present recent insights into antiviral RNAi and discuss the dynamic interplay between host antiviral RNAi responses and virus counter-defense strategies.
Insect-microbes interactions

The Drosophila-Spiroplasma interaction as a model to dissect the molecular mechanisms underlying insect endosymbiosis

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Virtually every species of insect harbors facultative bacterial endosymbionts that are transmitted from females to their offspring, often in the egg cytoplasm. These symbionts play crucial roles in the biology of their hosts. Many manipulate host reproduction in order to spread within host populations. Others increase the fitness of their hosts under certain conditions. For example, increasing tolerance to heat or protecting their hosts against natural enemies. Over the past decade, our understanding of insect endosymbionts has shifted from seeing them as fascinating oddities to being ubiquitous and central to the biology of their hosts, including many of high economic and medical importance. However, in spite of growing interest in endosymbionts, very little is known about the molecular mechanisms underlying most endosymbiont-insect interactions. For instance, the basis of the main phenotypes caused by endosymbionts, including diverse reproductive manipulations or symbiont-protective immunity, remains largely enigmatic. To fill this gap, we are dissecting the interaction between Drosophila and its native endosymbiont Spiroplasma poulsonii. This project will use a broad range of approaches ranging from molecular genetic to genomics to dissect the molecular mechanisms underlying key features of the symbiosis, including vertical transmission, male killing, regulation of symbiont growth, and symbiont-mediated protection against parasitic wasps. We believe that the fundamental knowledge generated on the Drosophila-Spiroplasma interaction will serve as a paradigm for other endosymbiont-insect interactions (ex. Wolbachia) that are less amenable to genetic studies.
How gut microbes interact with animals: insights from Drosophila

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Background: Animal surfaces and cavities provide multiple habitats for those microorganisms that can tolerate the conditions and exploit the resources provided by their animal hosts. Many of these microorganisms display adaptations for the host habitat, some of which are co-evolved with metabolic or immunity-related traits of the host. Teasing apart these interactions represents a major challenge, especially in humans and other mammals with very complex physiological systems and diverse microbial communities.

Objectives: To use the relationship between Drosophila and its gut microbiota as a model system to investigate the fundamentals of interactions between animals and their gut microbiota.

Methods: We have quantified the impact on the association of systematically varying the diet composition, microbiota composition and host genotype, using multiple methods including analysis of (meta)genome-wide associations

Conclusions: Key traits that are influenced by host and microbial genetic factors are the abundance of the microorganisms in the gut and the energy status of the host, including metabolic rate and fat content. Our first datasets also point to the importance of among-microbe interactions in shaping the total function of the microbial community. Because the microbial community in Drosophila is relatively simple, it is feasible to investigate these interactions systematically.
Termites degrade lignocellulose with the help of their intestinal symbionts. The general role of the gut microbiota in the digestive process is slowly emerging, but the specific functions of individual populations and their evolutionary origin are still in the dark. Deep sequencing of the hindgut community revealed strong differences among the major host groups, and dramatic changes in the abundance of particular taxa coincide with major events in termite evolution. The acquisition of cellulolytic protists by an ancestral cockroach provided new habitats for bacterial symbionts, which colonize the surface and cytoplasm of the gut flagellates in all evolutionary lower termites. After the loss of flagellates in higher termites, the wood particles became available for bacterial colonization, providing new niches for fiber-digesting populations. While some lineages in the core microbiota may be cospeciating with termites since the early Cretaceous, others seem to be specifically selected by the microenvironmental conditions in insect guts. Experiments addressing the role of the host in the selection of its gut microbiota are hampered by the elaborate social system of termites and their obligate dependence on intestinal symbionts. However, the controlled inoculation of germ-free cockroaches with normal and foreign microbiota confirmed that host factors are important drivers shaping the gut microbial community.
The diversity of shapes of organisms is one of the most fascinating aspects in the field of biology. While bacteria display a myriad of morphologies, the mechanisms that control morphogenesis and the evolution of bacterial morphology are not understood. I will describe the mechanisms that control morphological diversity in species related to *Caulobacter crescentus* that synthesize appendage-like extensions of the cell envelope at distinct sub-cellular positions. I will show that stepwise evolution of a specific domain of a developmental regulator led to the gain of a new function and localization of this protein, which drove the sequential transition in morphology. Our results indicate that evolution of protein function, co-option, and modularity are key elements in the evolution of bacterial morphology. In addition, I will show how evolutionary consideration of the mechanism of growth in the alphaproteobacteria led to the surprising discovery that polar growth, rather than the previously assumed binary fission, is the predominant mode of growth in a large group of the alphaproteobacteria that includes the plant pathogen *Agrobacterium tumefaciens* and the human pathogen *Brucella abortus*. Finally, I will describe new methods of peptidoglycan labeling that allow the detection of sites of peptidoglycan synthesis in live cells and in real time, and their use to study the mechanisms of peptidoglycan synthesis and to show for the first time that pathogenic Chlamydia have peptidoglycan, ending 50 years of speculation and debate concerning the chlamydial anomaly.
Bacterial chromosome segregation at the single-molecule resolution

How does a bacterial cell without a nucleus, without mitosis, and without extensive sister cohesion, organize, replicate, repair and segregate its chromosomes? We use live-cell super-resolution PALM and 3-D SIM imaging, alongside classical biochemistry and molecular genetics, and in vitro single-molecule biophysical techniques to address these questions. The presentation will focus on the sequential and coordinated action of Topoisomerase IV and the SMC complex, MukBEF, in E. coli chromosome segregation.

References

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Regulation of cell division in bacteria

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Background: Accurate positioning of the division site is essential for generating appropriately-sized daughter cells with the correct chromosome number. In bacteria, cell division initiates with assembly of the tubulin homologue FtsZ into a ring-like structure, the Z-ring. Thus, the position of Z-ring formation dictates the cell division site. Accordingly, all known systems that regulate positioning of the division site control Z-ring positioning.

Objective: We analyzed how the rod-shaped Myxococcus xanthus cells regulate positioning of the Z-ring at midcell.

Methods: Genetics, microscopy and in vitro analyses of proteins.

Conclusions: We report that PomX, PomY and the ParA-like ATPase PomZ are important for formation and positioning of the Z-ring. PomXYZ colocalize in a dynamically localized cluster and depend on each other for localization in this cluster. Early during the cell cycle this cluster localize on the nucleoid away from midcell. Subsequently, the cluster relocates by directed motion to midcell and localize here before and in the absence of FtsZ. PomZ stimulates the movement of the PomXYZ cluster to midcell in an ATP hydrolysis-dependent manner. In vitro PomX and PomY interact and stimulate PomZ ATPase activity. Moreover, PomY and PomZ separately inhibit FtsZ filament formation. We conclude that M. xanthus uses a novel system to precisely mark midcell for Z-ring formation and cell division. We propose that Z-ring formation is inhibited throughout cells by PomY and PomZ. At midcell, PomXYZ establish a zone that is permissive to Z-ring formation either by inhibition of PomYZ function and/or by direct stimulation of Z-ring formation.
The PALM nanostructure of FtsZ along the cell cycle of a pathogenic coccus

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Ovococci form a morphological group that includes several human pathogens (streptococci and enterococci). Their shape results from two modes of cell wall insertion, one allowing division, and another one allowing limited elongation. Both cell wall synthesis modes rely on a single cytoskeletal protein, FtsZ, since MreB is absent in ovococcal bacteria. Despite the central role of FtsZ in ovococci, a detailed view of the in vivo nanostructure of ovococcal Z-rings has been lacking thus far, limiting our understanding of their assembly and architecture.

We have developed the use of PhotoActivated Localization Microscopy in the ovococcus human pathogen Streptococcus pneumoniae by engineering spDendra2, a photoconvertible fluorescent protein optimized for this bacterium. Labeling of endogenously expressed FtsZ with spDendra2 revealed the remodeling of the Z-ring morphology along the division cycle at the nanoscale. Changes in the ring axial thickness and in the clustering propensity of FtsZ correlated with the advancement of the cell cycle. We observed double-ring sub-structures, suggestive of short-lived intermediates that may form upon initiation of septal cell wall synthesis. The nanostructural details provided in this work have been integrated in a model describing the morphology of the ovococcal Z-ring along the cell cycle.
Globally, each year about 270 Gt CO₂ are net fixed via photosynthesis into biomass, mainly lignocelluloses. Of this biomass about 2% end up in anoxic environments low in sulfate, Fe³⁺, Mn⁴⁺, nitrate and/or nitrite, where biomass is fermented via syntrophic associations of anaerobic protozoa, fungi, bacteria and archaea to methane and CO₂. The methane thus generated either accumulates in the sediments forming methane hydrates or diffuses into zones where the electron acceptors sulfate, Fe³⁺, Mn⁴⁺, nitrate, nitrite and/or O₂ are available and where methane is oxidized by anaerobic archaea and aerobic bacteria to CO₂. Part of the methane also escapes into the atmosphere, where it is photochemically converted to CO₂, and where methane is a potent greenhouse gas, which is of concern since the methane concentration in the atmosphere has doubled within the last 100 years. Many of the microbes involved in the methane cycle display a unique biochemistry, which my group helped to unravel within the last 50 years. In my lecture I will focus on the function and structure of the nickel enzyme methyl-coenzyme M reductase, which catalyzes in methanogenic archaea the methane forming step and in methanotrophic archaea the methane oxidizing step. The enzyme contains uniquely the nickel porphinoid F₄₃₀ as prosthetic group and, highly conserved, six posttranslationally modified amino acids within the active site, amongst them a thioglycine, two C-methylated amino acids and an only recently discovered didehydroaspartate. The function of the posttranslational modifications has remained enigmatic and therefore also the catalytic mechanism of MCR.

Background

The human gut harbors a densely populated microbial ecosystem, termed the gut microbiota. The gut microbiota can serve as a reservoir for antibiotic resistance genes (‘the resistome’) and drug-resistant opportunistic pathogens.

Objectives

In this study, we determined the dynamics of the microbiota and the resistome in thirteen patients that were hospitalized in Intensive Care Units. Because gut-associated bacteria and their resistance genes may be shed into the environment through sewage, we also assayed the levels of antibiotic resistance genes in hospital sewage and in subsequent stages of wastewater treatment.

Methods

16S rRNA profiling, functional metagenomics, metagenomic sequencing and nanolitre-scale quantitative PCRs were used in this study.

Conclusions

The gut microbiota of hospitalized patients contained genes that are predicted to confer resistance to fourteen classes of antimicrobials. Levels of antibiotic resistance genes increased during ICU hospitalization and were associated with changing patterns of the composition of the microbiota. Sequencing of vector inserts in drug-resistant fosmid clones indicated that resistance genes were mainly carried by anaerobic gut commensals (*Clostridium* and *Bacteroides*). Resistance genes were associated with mobile genetic elements, which could facilitate horizontal gene transfer of resistance genes from commensal bacteria to opportunistic pathogens. Antibiotic resistance genes were abundant in hospital sewage but the influence of hospital sewage on the abundance of resistance genes in the wastewater treatment...
plant was limited. Levels of resistance genes and human-associated bacteria further decreased during wastewater treatment, limiting the dispersal of antibiotic resistance genes from hospitals into the environment.
Background

The number of multi drug resistant pathogens is constantly growing and novel antibiotic substances are desperately needed in order to at least maintain the status quo. Ribosomally synthesized antimicrobial peptides are not yet exploited for human applications despite an indisputable potential. Among those, lantibiotics, a class of posttranslationally modified and thioether-ring bearing antimicrobial peptides, show desirable properties as high but specific antimicrobial activity and excellent stability.

Objectives

We follow a synthetic biology approach for the generation of novel lantibiotics employing the blueprint of natural lantibiotics. Based on their structural and functional features, natural lantibiotics are dissected into modular subunits. These subunits are then shuffled to generate thousands of novel, putative active, chimeric lantibiotics. The lantibiotic genes are subsequently produced by combinatorial DNA de novo synthesis. These libraries will then be screened for molecules with high antimicrobial activity.

Methods

To enable for high screening rates we developed a platform based on nL-sized reaction vessels (nL-reactors) that are used for peptide production and activity-screening in a single step and at rates of $10^5$ variants per day. Library cells are grown to microcolonies within nL-reactors along with a sensor strain. Cells secreting an active antimicrobial peptide will deactivate the sensor cells. Clearance of an nL-reactor from the sensor thus indicates the presence of a strain secreting a highly active peptide.

Conclusions
$10^4$ different chimeric variants derived from the module-shuffling of ten natural lantibiotics were analyzed. Several interesting antimicrobials have been isolated, demonstrating for the first time the generation of new-to-nature lantibiotics by module-shuffling.
Background
Lantibiotics and microcins form different groups of posttranslationally modified bacterial peptides mainly produced by Gram-positive and Gram-negative bacteria, respectively. Most of these peptides exhibit a potent antimicrobial activity, more active even than some antibiotics. Maturation of a peptide begins with a propeptide which is guided throughout specific modification events due to leader peptide (an amino acid sequence recognized by particular modification biomodule). The leader peptide is proteolitically removed in the last steps of peptide modification making the modified peptide active. Posttranslational modifications endorse lantibiotics and microcins with high target-specific activities and stability against proteolysis.

Objectives
To design plug-and-play biomodules for production of novel peptides containing both, lantibiotic- and microcin-specific posttranslational modifications.

Methods
We used a synthetic biology approach to design biomodules. We utilized peptide modification systems cloned under different controlled promoters to assure gradual and temporal activity of designed biomodules. Codon optimized synthetic genes of designed model peptides to be modified were employed together with modification machineries.

Conclusions
This is a unique case where posttranslational modifications from two different peptide classes are fused together into a single peptide chain in vivo. Here, we also show the possibilities of employing chimeric leader peptides for substrate recognition and procession by particular posttranslational modification machineries.
Background
Rapid development of antibiotic resistance of bacteria necessitates the search for new natural compounds inhibiting the growth and functioning of pathogenic microflora.

Objectives
To study the antibacterial mechanisms of warnerin, a new peptide from lantibiotic family that demonstrate the sequence of events causing the death of attacked bacterial cells.

Methods
Outcomes of warnerin attack of *Staphylococcus epidermidis* 33 GISK (Moscow) were analyzed using the combination of different methods such as polarography (respiration), spectrography (intracellular potassium loss), fluorescence assay of membrane potential dynamics and hydroxyl radicals’ formation, electron and atomic-force microscopy of cell structure damages

Conclusions
Peptide introduction into the bacterial incubation medium results in the increase in their oxygen consumption followed by rapid cellular release of potassium ions and sharp decline of intracellular ATP concentration. Apparently, the disturbance of ATP-linked electron transport along the respiratory chain favors their accumulation on iron-sulfur centres. This results in increase of free Fe$^{2+}$ ions and as a consequence, to the elevation of hydroxyl radical concentrations. Peptide inhibition of ATP formation is accompanied by marked proton accumulation on the outer side of bacterial membrane, particularly in its adjacent layers of peptidoglycane. This facilitates the autolytic hydrolase extrusion from the binding sites and to uncontrolled total bacteriolysis that is revealed by microscopic data. Results of investigation evidence for the pluripotent antibacterial action of warnerin that point to expressed prospects for its practical use.

The work was supported by RFBR grant No. 14-04-00687_a.
Background

With the emerging threat of infections caused by multidrug-resistant bacteria and scarce prospects of newly introduced antibiotics in the future, bacteriophages (phages) are currently being reconsidered as alternative therapeutics. However, the full breath of phage diversity suitable for treatment of bacterial infections is still largely unexplored.

Objectives

This study aims at providing an overview of the general occurrence of bacteriophages (with therapeutic potential) in natural environments and at devising strategies for their rapid characterization and classification at molecular level.

Methods

Novel phages were isolated from various aquatic systems (e.g. from general and hospital wastewater, activated sludge samples from sewage plants, streams, rivers, ponds and lakes). Environmental samples were pre-incubated with clinical isolates of multidrug-resistant bacteria. Enriched bacteriophages were subsequently obtained with the double agar layer plaque assay. “PhiSigns” (http://www.phantome.org/phisigns/) was used for phage identification based on the detection of signature genes. Genomic comparison of phages was performed based on digitized fluorescent restriction-fragment-length-polymorphism-analysis (fRFLP).

Conclusions

For roughly 50% of tested bacterial strains lytic bacteriophages were found against multidrug-resistant Pseudomonas aeruginosa, Klebsiella pneumoniae and Enterobacter sp. with waste water samples harbouring the highest phage diversity. By contrast no bacteriophages were isolated against multidrug-resistant Staphylococcus aureus and Acinetobacter baumannii. PhiSigns combined with fRFLP proofed to be a useful tool for rapid phage identification, which is the base for their proper selection for ultimate whole genome sequencing. S. aureus and A. baumanii phages require additional isolation steps such as FeCl₃-precipitation and/or use of
subinhibitory antibiotic-concentrations for enhancing plaque visibility on agar plates.
Background
The globally spread multi drug resistant *E. coli* clone ST131-O25b:H4 is responsible for a significant proportion of multi-drug resistant extraintestinal infections. By definition, isolates of this clone express the O25b antigen, whose unique structure has been resolved recently.

Objectives
We aimed to develop humanized monoclonal mAbs against the O25b antigen that may provide an alternative adjunct or stand-alone therapeutic option against this *E. coli* lineage.

Methods
Murine mAbs against the unique O25b antigen were generated by standard hybridoma technique. Following a primary screening, selected antibodies were subjected to humanization. Specificity and binding characteristics were assessed by immunoblots and ForteBio using purified antigens. Surface staining of ST131-O25b isolates irrespective of the capsular type expressed was shown by flow cytometry. Upon surface binding, the humanized antibodies induced complement mediated bactericidal activity in vitro. Passive immunization with the humanized mAbs elicited high level of protection at low mAb doses in a murine model of bacteremia. All whole cell assays were performed with bacteria grown in standard culture medium as well as in depleted pooled human serum in order to mimic *in vivo*-like conditions.

Conclusions
Humanized mAbs against *E. coli* O25b antigen were developed that elicited bactericidal activity *in vitro* and showed protection *in vivo*. Prophylactic passive immunization of colonized individuals or adjunct therapy of infected patients by mAbs may replace/substitute antibiotic therapy against this drug resistant clone. Relevant cases could be identified by the co-developed companion diagnostic tool.
LACTOCOCCUS GARVIEAE’S VIRUSES: AN INSIGHT INTO TEMPERATE AND VIRULENT BACTERIOPHAGES

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Background
Lactococcus garvieae, one of the most important pathogens in the aquaculture sector, has been recently recognized as an opportunistic human pathogen. However, little is known about the factors controlling of its growth.

Objectives
The aim of this project is to isolate and characterize temperate and virulent phages of L. garvieae.

Methods
Forty-five L. garvieae strains isolated from various ecological niches were tested for the presence of inducible prophages. Virulent phages were also isolated from different environments. Morphological and genomic characteristics of the new phages were studied through transmission electronic microscope (TEM), DNA restriction profile, protein profile, host range, and in few cases, genomic sequences.

Conclusions
Twenty temperate phages belonging to the Siphoviridae family and five new virulent phages able to infect L. garvieae strains have been isolated and partially characterized. Temperate phages, integrated into the genome of the host cell, promoted genome plasticity and appeared to be involved in gene mobilization. Virulent and strain–specific phages may be useful as alternatives to antibiotics to treat L. garvieae infections.
ALGAL VIRUS PRODUCTION IS NEGATIVELY AFFECTED BY PHOSPHORUS AND NITROGEN LIMITATION, BUT THIS MAY BE COUNTERACTED BY NUTRIENT ASSIMILATION DURING INFECTION

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Background
Viruses are a main cause of phytoplankton mortality, thereby controlling their community composition and driving biogeochemical cycling in the marine environment. Phytoplankton growth and production is often limited by nitrogen (N) and/or phosphorus (P), which can affect the virus growth characteristics upon infection. The influence of nutrient limitation on phytoplankton – virus interactions are, however, still poorly studied. As climate change will lead to an expansion of stratified and consequently nutrient limited oceanic regions, it is important to understand how phytoplankton mortality rates in the future marine environment will be affected by these changes.

Objectives
Our goal was to study the effects of N- and P-limitation on virus host interaction and the potential relieve of these effects by the uptake of nutrients during viral infection.

Methods
Therefore, cultures of the the phytoplankton species Micromonas pusilla and Phaeocystis globosa were grown semicontinuously under limitation of either P, N, or NP until the moment of infection. Upon infection the abundances of host cells and viruses were monitored to obtain one-step growth curves.

Conclusions
Nutrient limitation led to elongated viral latent periods (up to 3-fold) and strongly reduced (up to 8-fold) burst sizes, whereby the outcome was dependent on both the species and the limiting nutrient involved. Furthermore, the negative effects of P-limitation on virus production in the picophytoplankter M. pusilla could be overcome by spiking the infected and P-limited cultures with small pulses of inorganic and organic P, simulating bacterial remineralization and supply by lysis of neighboring cells, respectively.
REMARKABLE DIVERSITY AMONG NOVEL DENSOVIRUSES FROM CRICKETS

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Background
Densoviruses are members of the Parvoviridae family infecting invertebrates. Since 2009, a cricket densovirus, AdDV, caused a severe epidemic in the $600-million cricket industry in North-America.

Objectives
During the last 18 months, we also received samples of diseased crickets from North America, Europe and Japan that were negative for AdDV. Electron microscopy demonstrated that these samples contained densovirus-like particles. Objectives of this study included characterization of these virus(es), cloning, sequencing, and X-ray crystallography.

Methods
A SISPA method was used to detect the pathogens (J Virol 88:12152). Sequencing of the amplicons identified those with identities to known viruses allowing by primer extension to obtain the complete genomes and cloning in suitable vectors to obtain infectious clones. Sequencing revealed that some were novel circoviruses and a novel ambisense densovirus (Genome Announc. 1(2):e00079-13, 1(3):e00328 and 131(6):e00914-13). However, the greatest surprise was a densovirus with a segmented genome (AdSDV). NS- and VP- coding sequences are in separate segments of about 3.3 kb. Both NS- and VP- segments are flanked by identical hairpin terminal sequences but lack ITRs. The two ORFs for VP arose through recombination between a mosquito brevidensovirus and AdDV. The downstream ORF with AdDV-VP1up-like phospholipase A2 could be spliced to the C-terminus of the major capsid protein. Currently, the capsid structure (X-ray crystallography) and transcription strategy of AdSDNV.

Conclusions
A brevidensovirus that normally infects only mosquitoes has adapted itself through recombination with AdDV to crickets. These findings necessitate a revision of the definition and taxonomy of parvoviruses.
FEMS-1562
Poster Discussion: Virology

VACCINE-TYPE ATTENUATING MUTATIONS IN HIGHLY PASSAGED STRAIN OF VARICELLA ZOSTER VIRUS

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Background: Varicella-zoster virus (VZV) is causative agent for chickenpox and zoster. Live attenuated vaccines have been developed based on Japanese Oka strain and Korean MAV/06 strain. Comparison of the complete sequences between vaccine and wild-type strains suggested at least 24 mutations important for attenuation in vaccine strains. Certain strains with high in vitro passage history such as the strains Ellen and 32p72 contained some of the vaccine-type mutations.

Objective: In this study, we attempted to understand vaccine-type attenuating mutations in highly passaged VZV strains.

Methods: Genome sequences of clinical strains with different passages were determined by next-generation sequencing and compared with those of other VZV strains.

Conclusion: Vaccine-type mutations were found at position 106262 (T→C) in the Korean clinical strain YC01 passage 32 and at positions 560 (T→C), 105169 (A→G), 106262 (T→C) and 107252 (T→C) in YC01 passage 61. Similar mutations were also found in high-passaged another clinical strain YC02 at positions 560, 106262 and 107252. Same mutations at 106262 and 107252 were found in the strains Ellen and 32p72. Thus, in the course of in vitro passaging of VZV, mutations at 106262 and 107252 seemed to occur first, and followed by mutation at 560. Direct PCR sequencing of these positions in YC01 and YC02 strains at various passages identified the passage numbers when the attenuating mutations occurred. Further studies of in vitro passaging under attenuating conditions such as low temperature or guinea pig cells will help to understand the mechanism of attenuating mutations in VZV.
Background
Dengue virus (DENV) infection causes viral haemorrhagic fever that is characterized by extensive activation of the immune system.

Objectives
The aim of this study is to investigate the kinetics of the transcriptome signature changes during the course of disease and the association of genes in these signatures with clinical parameters.

Methods
Sequential whole blood samples from DENV infected patients in Jakarta were profiled using affymetrix microarrays, which were analysed using principal component analysis, limma, gene set analysis, and weighted gene co-expression network analysis.

Conclusions
Time since onset of disease associates with the shift in transcriptome signatures from immunity and inflammation to cell cycle and repair mechanisms in patients with non-severe dengue. The strong association of time with blood transcriptome changes hampers both the discovery as well as the potential application of biomarkers in dengue. Clinical diagnosis (according to the WHO classification) does not associate with differential gene expression. However, network analysis did show that that key clinical markers, including platelet count, fibrinogen, albumin, IV fluid distributed per day and liver enzymes SGOT and SGPT, strongly correlate with gene modules that are enriched for genes involved in the immune response. The expression level of these gene modules may support earlier detection of disease progression as well as clinical management of dengue.

CHARACTERIZING THE VIROME OF THE ENTOMOPATHOGENIC FUNGUS BEAUVERIA BASSIANA

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Background

Entomopathogenic fungi are of scientific interest because they enable analysis of virus-host interactions and can be used as biocontrol agents of insect pests. The deuteromycetous fungus Beauveria bassiana has a widespread geographical distribution and a wide host range. Mycoviruses have been described mostly in phytopathogenic fungi and are occasionally associated with hypovirulence, while in entomopathogenic fungi presence of mycoviruses has been reported only rarely.

Objectives

The aim of the present study is to detect and analyse the viruses present in a collection of B. bassiana isolates sourced from worldwide locations.

Methods

Population studies, RT-PCR, cloning and sequencing, hybridization experiments, time-course studies and transmission electron microscopy (TEM) were used to characterize the virome of B. bassiana.

Conclusions

A population study revealed that 17/75 (22.7%) B. bassiana isolates harbor dsRNA elements. Two members of the Partitiviridae family, B. bassiana partitivirus-1 and -2 (BbPV-1 and BbPV-2), have been sequenced, while TEM revealed the presence of virus-like particles. Hybridization experiments revealed that BbPV-1 and BbPV-2 are present in seven B. bassiana isolates derived from different hosts and geographical origins. Additionally, four members of the Totiviridae family, originating from the Iberian Peninsula and the Canary Islands, have been partially sequenced. Furthermore, two previously uncharacterized viruses, B. bassiana polymycovirus-1 (BbPmV-1) and B. bassiana non-segmented virus-1 (BbNV-1), have been detected and fully sequenced and the prevalence of further PmV-like viruses in B. bassiana is currently under investigation. Finally, a time-course study revealed a strong negative
correlation between the copy numbers of viral dsRNA and the developmental stages of the fungus.
CHARACTERIZATION OF VIRULENCE FACTORS AND IMMUNE-STIMULATING ANTIGENS IN VIBRIO SALMONICIDA

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Background

*Vibrio salmonicida* is the cause of cold-water vibriosis in farmed Atlantic salmon. The disease manifests as a hemorrhagic septicemia, with a mortality ranging from 3-90%. In the 1980s, the disease was a major challenge to the Norwegian aquaculture industry, but it is now being effectively controlled by the use of oil-based vaccines. Nevertheless, in recent years there have been some reports of disease outbreaks. In 2013, 13 Norwegian fish farms were diagnosed with cold-water vibriosis, all localized in the three northernmost counties of Norway.

Objectives

The knowledge on virulence factors of *Vibrio salmonicida* and the subsequent immune response of the fish is limited. As bacteria adapt to their current environment, a bacterium isolated from a diseased fish will not be identical to the same species cultivated *in vitro*. The aim of this study was to explore the phenotype of bacteria present *in vivo*.

Methods

We have grown *Vibrio salmonicida* in semi-permeable implants in live fish and analyzed bacterial protein expression by two-dimensional gel electrophoresis and tandem mass spectrometry. The experiment was approved by the Norwegian Animal Research Authority (approval no. ID6228).

Conclusions

Our findings may contribute to the knowledge on disease progression in cold-water vibriosis, as well as to provide novel targets for more effective vaccines. Also, the
study may provide insights in the course of disease for other bacterial infections in fish.
IDENTIFICATION AND TYPING OF LACTOCOCCUS GARVIEAE BASED ON MALDI TOF MS

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Background

MALDI-TOF-mass spectrometry (MS) has emerged as a powerful technique for the routine identification of microorganisms in clinical microbiology laboratories allowing an easier and faster diagnosis than conventional phenotypic and molecular methods. We have evaluated the suitability of MALDI-TOF for the rapid identification of Lactococcus garvieae, the etiological agent of lactococcosis, a septicemic infection affecting different wild and farmed fish species, and recognized also as an opportunistic emerging human pathogen.

Objectives

Assessment of the reliability of MALDI-TOF MS for the identification of Lactococcus garvieae

Methods

A total of 35 isolates from different host and environments were obtained from the culture collection of VISAVET were used in the study. The identification results obtained by MALDI TOF were compared with those obtained by biochemical identification (API 32 Strep) and by a species-specific PCR assay (Zlotkin et al. 1998. J Clin Microbiol 36, 983-985). Mass spectra acquisition and analysis was performed on a Bruker UltraFlextrem platform (Bruker Daltonics) using MALDI Biotyper™ 3.0 software in the automatic mode using a matrix of saturated solution of α-HCCA. In addition, a subset of isolates was subjected to PFGE.

Conclusions

The majority of peaks were obtained in the range from m/z 2000 to 10000. The proteomic results matched with those of genotypic approach. Our results also showed some differences in MS spectra in L. garvieae isolates recovered from different origins/hosts suggesting the possible identification of molecular biomarkers for L. garvieae. Our study demonstrates that proteomics identification using MALDI TOF MS could be a reliable approach for identifying and discriminating this microorganism.
O SEROGROUPS OF ENTEROPATHOGENIC (EPEC) AND SHIGATOXIGENIC (STEC) ESCHERICHIA COLI FROM
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Background

*Escherichia coli* producing the attachment-effacement (AE) lesion (EPEC) and/or Shiga toxins (STEC) cause enteritis and (bloody) diarrhoea in young calves and in humans, and are also present in the intestines of healthy cattle. Besides the O157:H7 serotype, EPEC and STEC can belong to more than sixty O serogroups. Of them, 8 have been most frequently identified worldwide: O5, O26, O103, O111, O118, O121, O145 and O165, with some also causing diarrhoea in young calves.

Objectives

This study aimed at identifying the pathotypes and the O serogroups of STEC and EPEC isolated from <1-month-old diarrhoeic calves in Wallonia, Belgium.

Methods

(i) 233 enterohaemolysin-producing *E. coli* were isolated at ARSIA between November 2008 and February 2014 from diarrhoeic calves after growth on EHLY Medium®. They were tested with a triplex PCR targeting the *stx1*, *stx2* (Shiga toxins) and *eae* (AE lesion) genes.

(ii) triplex PCR-positive *E. coli* were assayed with two pentaplex PCR targeting the specific genes coding for the nine O serogroups listed above and for the O104 serogroup.

Conclusions

(i) 206 isolates tested positive with the triplex PCR. The most frequent pathotypes were *eae+stx1+* (102 isolates), *eae+* (78 isolates) and *eae+stx1+stx2+* (13 isolates).
(ii) the most frequent serogroups of EPEC and STEC were O26 (57 isolates) and O111 (36 isolates). A few additional isolates tested positive for the O103, O5, O145, O121 and O157 serogroups.

(iii) the future is to compare these EPEC and STEC with those isolated from healthy cattle and from humans, to identify host- and age-specific properties.
CHARACTERISTICS AND ANTIMICROBIAL SUSCEPTIBILITY OF TRUEPERELLA PYOGENES ISOLATED FROM BOVINE MASTITIS CASES IN CHINA

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Background

Trueperella (T) pyogenes is an opportunistic pathogen causes suppurative diseases in dairy cows. However, the pathogenesis and antibiotics resistance of T. pyogenes are still unclear.

Objectives

T. pyogenes isolates morphological characteristics, the presence of genes (plo, cbpA, fimA, fimC, fimE, fimG, nanH, nanP, tet/W/, erm/X/, erm/B/), biofilm formation, the cytopathological effects in intracellular assay and antimicrobial resistance were investigated.

Methods

T. pyogenes was isolated from 50 out of 275 clinical and subclinical bovine mastitis cases in China. Both pyolysin (plo) and collagen-binding protein (cbpA) virulence factor genes were detected by conventional PCR in all T. pyogenes isolates. The tissue culture plate method was used to assay the capacity of T. pyogenes for biofilm formation and showed that 90% of T. pyogenes isolates were able to form biofilms with different production amount. Minimum inhibitory concentrations (MICs) of 14 antimicrobial agents were determined and observed high susceptibility to rifampin (96%), while high resistance to trimethoprim–sulfamethoxazole (90%) and bacitracin (98%). Intracellular assay revealed that 4 different T. pyogenes isolates had different cytopathological effects on infected bovine mammary gland epithelial cells.

Conclusions

18% T. pyogenes isolates indicates that T. pyogenes is important contributors to bovine mastitis. Moreover, the within-host quantitative, spatial, high occurrence of multidrug-resistant, biofilm producing and temporal dynamics of T. pyogenes interactions are key factors to better understand how immunity acts on infections with bacteria and how they evade immune surveillance; thus, highlighting the need for prudent use of antimicrobial agents in veterinary medicine (This research was supported by projects No. 2012BAD12B03, No. 313054, No. 20120008110042, No. 2014M561102 and No. GDT20141100043).
FUNCTIONAL RESILIENCE OF MICROBIAL ECOSYSTEMS IN SOIL: HOW IMPORTANT IS A SPATIAL ANALYSIS?

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Background

Microbial life in soil is exposed to fluctuating environmental conditions influencing the performance of ecosystem services such as biodegradation.

Objectives

However, as this environment is typically very heterogeneous, spatial aspects can be expected to play a major role for the ability to recover from a stress event.

Methods

To determine key processes for functional resilience, scenarios with varying stress intensities were simulated within a microbial simulation model and the biodegradation rate in the recovery phase measured. Besides temporal monitoring, results were analyzed regarding spatial and mechanistic aspects. Parameters including microbial growth and dispersal rates were varied over a typical range to consider microorganisms with varying properties.

Conclusions

Results of the mechanistic and spatial view show that key factors for functional recovery depend on stress intensity and the location of the observed habitats. The limiting factors near unstressed areas are spatial processes (motility and substrate diffusion), with increasing distance microbial growth becomes more important. To confirm this, we repeated the simulations including a dispersal network representing fungi in soil. The system benefits from an increased spatial performance due to higher bacterial mobility.

With these simulations we show the importance of spatial aspects even at the mm-scale for recovering after a severe stress event in a highly heterogeneous environment such as soil. In consequence a spatial-mechanistic view is necessary for examining the functional resilience as the temporal view alone could not have led to these conclusions.

Further research should explore the importance of a spatial view for quantifying functional resilience also after complex stress regimes.
PHYSIOLOGIC AND GENOMIC CHARACTERIZATION OF A NOVEL NITROSPIRA SPECIES ENRICHED UNDER ANAEROBIC, DENITRIFYING CONDITIONS

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Background

Members of the genus Nitrospira are dominant in many natural habitats and of vital importance for wastewater treatment. They are chemolithoautotrophic organisms capable of growth with nitrite and CO2 as sole energy and carbon source. It further has been demonstrated that some Nitrospira can utilize simple organic carbon compounds and molecular hydrogen as alternative substrates and can switch to nitrate reduction under oxygen limitation. However, this metabolism has been assumed to be a survival strategy and in most systems studied so far Nitrospira was outcompeted by denitrifying organisms when anoxic conditions prevailed.

Objectives

Here, a community of nitrogen cycle bacteria was enriched from the anaerobic compartment of a biofilter connected to a recirculating aquaculture system.

Methods

The culture was fed with filtered water from the aquaculture system, supplemented with ammonium, nitrite and nitrate.

Conclusions

A stable enrichment culture was obtained which anaerobically converted ammonium, nitrite and nitrate into dinitrogengas. The culture was dominated by anaerobic ammonium-oxidizing Brocadia species, a denitrifier (Denitratisoma), and a Nitrospira species. Anammox bacteria and Nitrospira co-aggregated, while the denitrifiers formed separate clusters. This culture shows that anaerobic ammonium oxidizers and nitrite oxidizers, who were considered to be mutually exclusive, can be grown as a stable co-culture. Furthermore, this shows that some Nitrospira species can be competitive under nitrate-reducing conditions. This study demonstrates another unexpected lifestyle for an organism believed to be only competitive under aerobic lithoautotrophic conditions. It further indicates that Nitrospira in combination with anammox can be of interest for anoxic wastewater treatment systems.
BACTERIAL PROFILING OF SAHARAN DUST DEPOSITION IN THE ATLANTIC OCEAN USING SEDIMENT TRAP MOORINGS – YEAR ONE RESULTS

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Background

Large quantities of dust are transported from the Sahara Desert across the Atlantic Ocean towards the Caribbean each year, with a large portion of it deposited in the ocean. This dust brings an array of minerals, nutrients and organic matter, both living and dead. This input potentially fertilizes phytoplankton growth, with resulting knock-on effects throughout the food chain. The input of terrestrial microbial life may also have an impact on the marine microbial community.

Objectives

The aim is to understand the links between dust input and the bacterial community and how this relates to ocean productivity and the carbon cycle.

Methods

The current multi-year project consists of a transect of floating dust collectors and sub-surface sediment traps placed at 12°N across the Atlantic Ocean. Sediment traps are located 1200m and 3500m below the sea surface and all are synchronized to collect samples for a period of two weeks.

Conclusions

The first set of sediment trap samples were recovered using the RV Pelagia in November 2013 with promising results. Results from 7 sediment traps (three at 1200m and four at 3500m) were obtained. In general, the total mass flux decreased as distance from the source increased and the upper traps generally held more material than those at 3500m. Denaturing Gradient Gel Electrophoresis (DGGE) was used as a screening technique, revealing highly varied profiles, with the upper (1200m) traps generally showing more variation throughout the year. Several samples have been submitted for high throughput DNA sequencing which will identify the variations in these samples.
LIFE IN A MICROWAVE: EFFECTS OF MICROWAVE RADIATION ON THERMUS SP.
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Background
Biological systems are frequently exposed to microwave radiation. Many studies have investigated the influence of microwaves on these systems, but controversy over methods to distinguish between thermal and non-thermal microwave effects remains.

Objectives
To differentiate between non-thermal and thermal microwave effects on a physiology of a microorganism, a thermophilic bacterium was grown in a constant-temperature microwave or a convection oven. Comparing the growth properties of the thermophile in these conditions will reveal non-thermal microwave effects on cell growth and
physiology. Biophysical and biochemical analysis will demonstrate changes in morphology and chemical composition arising from microwave exposure.

Methods
Cell growth was analyzed by optical density (OD) measurements (supported by independent quantitative DNA analysis), and cell morphologies were characterized using electron microscopy imaging (SEM, TEM), dynamic light scattering (DLS), and atomic force microscopy (AFM). AFM was also used to probe the biophysical characteristics of the cells, in conjunction with nano-infrared spectroscopy (Nano-IR). Attenuated total reflectance infrared spectroscopy (ATR-IR) and fatty acid methyl ester (FAMEs) analysis were used to determine biochemical differences between cells grown in microwave and oven conditions.

Conclusions
Thermophilic bacteria were grown in a synthetic microwave such that thermal effects and microwave effects were distinguishable. These data demonstrate that there are physiological differences between cells cultured in a dielectric field and a convection oven, and that microwaves induce non-thermal changes to the structure, physiology, and chemical composition of the organism.
Background

Increasingly appearing resistances to antibiotics and the subsequent need for new bioactive substances is one of the fundamental challenges of infection research. Myxobacteria have a rich secondary metabolism, which places them among the best known natural product producers. To date more than 100 new metabolites and more than 500 derivatives were described from these soil dwelling, predatory microorganisms. In the past it turns out that in particular new families, genera and species of myxobacteria are reliable sources for new compounds.

Objectives

Promising sources for the isolation of new myxobacteria are uncommon habitats like, for example, peat bogs or fen, which are characterized by low pH values. This habitat has been neglected with regard to the isolation of myxobacteria in the past. In our study, the diversity of myxobacteria in moor-samples from the Harz-region was evaluated by standard cultivation methods as well as by cultivation independent 16S rRNA-clone bank analyses.

Methods

A total of 297 clones of myxobacterial origin were sequenced and compared to sequences of cultures, isolated from these samples, by phylogenetic analyses. It turned out that the majority of myxobacteria is only represented by clone sequences and could not be cultivated. Comparing these sequences to sequences of a public database (NCBI) revealed that most of these uncultivated myxobacteria are exclusively related to other uncultivated myxobacterial sequences from acidic soils.

Conclusions

This suggests that peat bogs and fen harbor a big diversity of new myxobacterial species, genera and even families which themselves harbor a great potential of urgently needed, new secondary metabolites.
THE MICROORGANISM THAT NEVER READ THE LITERATURE - FASTEST GROWING PHOTODAMAGE TOLERANT ALGA ISOLATED FROM DESERT CRUSTS, DEPENDS ON PIONEER FILAMENTOUS CYANOBACTERIA SPECIES TO SURVIVE DESICCATION

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Background
With the premise that unique capabilities can be found in microorganisms facing extremely harsh conditions, we focused our efforts on desert biological sand crusts (BSC), one of the harshest environments to support life. Recently, we have isolated a small green alga (named Chlorella ohadii), which does not conform to some of the basic fundamentals in microbial ecology and photosynthesis:

Objectives

1. Acquired ability to acclimate to extreme environments is usually accompanied by reduced performance under optimal conditions. We intend to show that C. ohadii does not obey this rule. In its natural habitat, C. ohadii is facing diurnal desiccation-hydration cycles, vast temperature amplitudes and extremely high illumination intensities, yet when grown under optimal laboratory conditions it exhibits the fastest growth rates ever reported for an alga.

2. After many years of research, some feel that we have elucidated the functioning of the photosynthetic machinery, and what sets the upper limit for algal growth. However, the unparalleled fast growth, extremely high photosynthetic rates and resistance to photodamage, suggest this may not be the case. Namely, C. ohadii has been shown to be completely resistant to photoinhibition, and its productivity was unaffected by irradiances as high as twice full sun light.

3. Survival of C. ohadii in its BSC habitat depends on close association with filamentous cyanobacteria; a unique and novel mode of interspecies interaction.

Methods
We wish to present detailed physiological analysis of the unique properties supporting this phenomenal growth and resistances, combined with insights from their genomic, transcriptomic and metabolomic characterization.

Conclusions
LEVANSUCRASE OF PSEUDOMONAS SYRINGAE AS PRODUCER OF NOVEL PREBIOTIC FRUCTANS: A STRUCTURE-FUNCTION STUDY OF THE ENZYME

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Background

Gut microbiota has numerous roles in human life and its significance has thus far been underestimated. Therefore, more detailed studies on diversity and metabolism of gut microbiota are needed.

Objectives

Gut microbiota can be shaped by prebiotics. Levan-type (β 2,6-linked) fructans can be considered as potential prebiotics [1]. We have synthesized levan and levan-type fructooligosaccharides by heterologously expressed levansucrase Lsc3 of Pseudomonas syringae pv. tomato. These substrates should affect gut microbiota as were perfectly fermented by a gut generalist B. thetaiotaomicron [2]. Here we focus on structure-function relationships of the Lsc3 protein and its biotechnological potential.

Methods

The His-tagged Lsc3 protein was site-directedly mutated, biochemical methods were used to characterize the mutant proteins and 3D structure modelling was applied to interprete the results.

Conclusions

Levansucrase Lsc3 has high polymerizing activity (up to 80%) and extremely high stability confirming its biotechnological potential for the synthesis of fructans. Several novel catalysis-related positions for levansucrases were revealed and their predicted location on a 3D model of Lsc3 will be presented.

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References:


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Poster Discussion: Food Microbiology and Biotechnology

MATCHING GENOME AND TRANSCRIPTOME OF LACTOCOCCUS LACTIS STRAINS WITH ROBUSTNESS TOWARDS INDUSTRIAL STRESSES


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Background

*Lactococcus lactis* is industrially employed to manufacture fermented dairy products. Spray drying is the cheapest preservation method for *L. lactis* starter cultures, but during this process cultures encounter heat and oxidative stress, resulting in low survival. Viability of starter cultures is essential for their role in dairy acidification, supporting the need to understand and improve their robustness.

Objectives

We have measured the ability of 39 *L. lactis* strains to survive industrially relevant stresses. This set of strains contained different *L. lactis* strains from dairy as well as plant origin. The observed 4-log variation in heat and oxidative stress survival was compared with genomic content, resulting in the identification of genes associated with robustness. Presence and activity of genes can play an important role in the observed diversity in robustness.

Methods

Therefore, four *L. lactis* strains with varying robustness phenotypes were fermented under twelve different conditions, varying in temperature, salt concentration, pH, and oxygen level. Cells were harvested at exponential phase of growth for transcriptome analysis and survival measurements.

Conclusions

The varying growth conditions resulted in up to 4-log differences in robustness towards heat and oxidative stress. Moreover, clear differences in gene expression profiles were observed. Correlation of robustness phenotypes and gene expression levels revealed transcriptome signatures for oxidative and heat stress survival. For strain MG1363 this included the *metC-cysK* operon, involved in methionine and cysteine metabolism, which triggered us to grow this strain in the absence of...
cysteine, resulting in elevated expression levels of the *metC-cysK* operon and concomitant enhanced robustness towards oxidative stress.
COOPERATION BETWEEN SPECIES IN THE YOGURT CONSORTIUM IS AFFECTED BY THE MODULATION OF INTRACELLULAR PH THAT IS DRIVEN BY STREPTOCOCCUS THERMOPHILUS UREASE ACTIVITY

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Background

The proto-cooperation between Streptococcus thermophilus and Lactobacillus delbrueckii subsp. bulgaricus in the yogurt consortium enhances the growth rate and size of each population. In contrast, the independent growth of the two species in milk leads to a slower growth rate and a smaller population size.

Objectives

In this study, we evaluated how urease activity of S. thermophilus might affect the intracellular pH of L. delbrueckii, modulating its bioenergetics during milk fermentation.

Methods

Urease-mediated intracellular alkalization of S. thermophilus and L. delbrueckii was evaluated by flow cytometry using cFSE as pH-dependent fluorescent probe. Lactose consumption and lactic acid production was followed in vivo by ¹³C-NMR analysis. D-L lactic acid production in milk was measured enzymatically. For the evaluation of the pH-dependent glycolysis efficiency, glucose and lactic acid were measured by HPLC.

Conclusions

We observed that intracellular alkalization caused by urea hydrolysis or the addition of ammonia to milk boosted lactic acid production in S. thermophilus and in L. delbrueckii when the species were grown separately or in combination. Therefore, we propose that urease activity acts as an altruistic cooperative trait, which is costly for urease-positive individuals but provides a local benefit because other individuals can take advantage of urease-modulated pH. Compared to the interactions that are known to occur between S. thermophilus and L. delbrueckii in the yogurt consortium, the modulation of the bioenergetic efficiency due to the intracellular pH alkalization represents a new type of cooperation that directly affects the kinetic parameters of enzymes involved in homolactic fermentation in both the species.
LISTERIA MONOCYTGENES OF SEQUENCE TYPE 121 HARBOR SPECIFIC ADAPTATIONS SUPPORTING PERSISTENCE IN FOOD PRODUCTION PLANTS.
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Background
The foodborne pathogen Listeria monocytogenes is able to survive for months and even years in food production environments. Among a great strain diversity particularly strains belonging to sequence type (ST)121 are persistent in food production plants.

Objectives
To elucidate the molecular determinants responsible for persistence of L. monocytogenes.

Methods
We analysed the genomes of two L. monocytogenes ST121 strains, which persisted for up to eight years in food production plants in Ireland and Austria. Additionally we characterized two ST121 strain-specific genetic features: Tn6188 and lin0464/lin0465.

Conclusions
All ST121 genomes are highly similar and show a tremendously high degree of conservation among prophages and particularly among their plasmids, which are usually variable parts of genetic information in bacteria. This remarkably high level of conservation suggests a strong selective pressure.

In addition, all ST121 strains share adaptations related to persistence in food production environments such as the presence of Tn6188, a transposon responsible for increased tolerance against quaternary ammonium compounds, and the presence of homologues of the L. innocua genes lin0464 and lin0465, a transcriptional regulator and a putative pfpI protease. Deletion of lin0465 resulted in reduced survival under oxidative and alkaline conditions suggesting a role in stress response. Furthermore all ST121 strains reveal a yet undescribed insertion harboring recombination hotspot (RHS) repeat proteins, which are most likely involved in
competition against other bacteria. In conclusion we show that *L. monocytogenes* ST121 strains are highly similar to each other harboring conserved regions which provide fitness adaptations to survival in food production environments.
Background

While inimical food preservation approaches often result in heavily damaged and stressed populations of foodborne pathogens and spoilage microorganisms, very little is known about the actual molecular and genetic events and dynamics that govern the (sub)lethal injury and subsequent resuscitation or death of these cells. Nevertheless, insights into these phenomena might be decisive for a proper understanding of the resulting behavior and evolvability of the surviving subpopulation.

Objectives

In this study, we set out to monitor and dissect (sub)lethal injury and resuscitation phenomena in populations and cells of *Escherichia coli* that are stressed by heat or high hydrostatic pressure treatment.

Methods

Directed evolution, genetics and time-lapse fluorescence microscopy.

Conclusions

Based on adaptive mutations, genetics and live cell biology, different subcellular processes and structures have been delineated and monitored that play a role in the heterogeneous injury and resuscitation dynamics of high temperature or pressure stressed *E. coli* cells and populations, thereby revealing a surprising impact on the spatial dynamics of nucleoids and protein aggregation inside stressed cells.
MAXIMIZING BIOTECHNOLOGICAL PRODUCTION OF 3-HYDROXYPROPIONALDEHYDE AS A NATURAL FOOD PRESERVATIVE WHILE MINIMIZING ITS CONVERSION TO TOXIC ACROLEIN
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Background

3-hydroxypropionaldehyde (3-HPA) has strong antimicrobial activity and application potential as a natural food preservative for disinfection of minimally processed vegetables. During glycerol metabolism, Lactobacillus reuteri DSM 20016T can accumulate 3-HPA, which undergoes reversible dimerization and hydration, and can be further converted enzymatically into inactive 1,3-propanediol (1,3-PD), restoring intracellular redox potential. Dehydration of 3-HPA resulting in acrolein, a toxic metabolite, is also possible, however, conditions promoting this transformation have not been fully understood due to the lack of analytical methods to simultaneously quantify 3-HPA as well as acrolein.

Objectives

The aim of this study was to maximize the capacity for biotechnological processing of glycerol to 3-HPA using L. reuteri, while accounting for the accumulation of acrolein after production and storage to evaluate toxicological risk.

Methods

Strategies to improve 3-HPA production included the addition of redox-active compounds during glycerol fermentation to prevent conversion to 1,3-PD, the usage of immobilized cells resulting in high densities of viable cells as well as the optimization of process parameters in batch and fed-batch processes. Special attention was paid to cultivation conditions of L. reuteri prior to glycerol conversion. Acrolein and 3-HPA were simultaneously quantified using a newly established analytical method employing ion-chromatography with pulsed amperometric detection.

Conclusions

Redox-active compounds, process conditions and pre-treatment of L. reuteri cultures influenced 3-HPA yield as well as ratio of 3-HPA to 1,3-PD. Temperature and pH
optimization prevented formation of acrolein during biotechnological production and storage, promoting the safety of 3-HPA when applied as natural food preservative.
Background
Candida species are responsible for recurrent human infections, mostly in immunocompromised patients, due to their high vulnerability. Candida glabrata has been showing to have a major role in these infections being the second most prevalent species involved in human fungemia. Amphotericin B (AmB), a common antifungal drug, is a hospital-environment exclusive polyene, normally being efficient when used to fight candidiasis.

Objectives
The main goal of this work was to infer about the influence of AmB in Candida glabrata biofilms formation and its effect on matrix composition and ERG genes expression.

Methods
Candida glabrata biofilms were formed in the presence of AmB and analyzed by dry weight. Moreover, ERG genes expression was evaluated by qRT-PCR and matrix was analyzed in terms of composition in carbohydrates, proteins, beta-glucans and a new finding: ergosterol.

Conclusions
In addition to an inefficient reduction of the C. glabrata biofilms, this work showed that ERG genes seem to be less involved than the matrix composition in C. glabrata biofilms response to AmB. Specifically, C. glabrata biofilms matrices respond with an increase of carbohydrates, particularly beta-1,3 glucans, and with a decrease of total proteins. The ergosterol values did not expressively changed in the presence of AmB.

The present work support the theory of multifaceted mechanisms developed by C. glabrata biofilms as response to the presence of AmB.
EFFECT OF SHEAR STRESS ON PSEUDOMONAS AERUGINOSA ISOLATED FROM THE CYSTIC FIBROSIS LUNG

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Background
Chronic Pseudomonas aeruginosa lung infections are the main cause of morbidity in cystic fibrosis (CF) patients. Previously, we have sequenced the genome of the transmissible Belgian epidemic strain CF_PA39. This CF-adapted strain has been present in the UZ Brussel hospital environment for at least ten years. Furthermore, earlier experiments showed that culturing P. aeruginosa PAO1 in a low fluid shear environment, obtained by means of the Rotating Wall Vessel (RWV) technology, leads to the formation of a biofilm phenotype comparable to that observed in the CF lung.

Objectives
To study the effect of fluid shear on biofilm formation by an adapted P. aeruginosa CF strain in artificial sputum medium (ASM) at the genotypic, transcriptomic, and phenotypic level.

Methods
In this study, an RWV experiment was performed that closely resembled the in vivo situation by inoculating this adapted CF isolate in ASM in the RWV either with or without two glass beads, simulating high and low fluid shear conditions, respectively. Scanning electron microscopy (SEM) was utilized to determine biofilm formation in both conditions, while RNAseq and qPCR was used to study gene expression in both conditions.

Conclusions
Increasing fluid shear in the RWV model disrupted biofilm formation of an adapted P. aeruginosa CF isolate.
In accordance with the biofilm disruption, several genes involved in denitrification, tryptophane synthesis, choline metabolism, and alginate biosynthesis were down-regulated in the high fluid shear condition, resembling the planktonic stage of growth. Furthermore, we identified small RNAs that are differentially expressed.
Background

Listeria monocytogenes is a ubiquitous opportunistic human pathogen detected in many habitats spanning from the farm environment to food industry and the gastrointestinal tract.

Cell-cell communication participates to the adaptation of bacteria to their environment. In the species Listeria monocytogenes, the Agr system is required for full virulence and biofilm formation but its actual role is still poorly understood.

Objectives

To investigate whether the ability to communicate provides a benefit to L. monocytogenes in soil, a complex environment combining biotic and abiotic characteristics.

Methods

Deletion of the gene coding the response regulator or the signal propeptide. Population dynamics of the mutant and/or parental strains in soil microcosms. Both unsterilised and sterilised soils were investigated.

Conclusions

Deletion of the gene coding the regulator or the signal did not affect population dynamics in sterilized soil but survival was altered in biotic soil suggesting that the Agr system was involved to face the complex soil biotic environment. This was confirmed by co-incubation experiments. The fitness of the response negative mutant was lower either in the presence or absence of the parental strain but the fitness of the signal negative mutant depended on the strain with which it was coincubated. Survival of the signal negative mutant was higher when cocultured with the parental strain than when cocultured with the response negative mutant. These results showed that the ability to respond to communication provided a benefit to listerial cells. These results might also indicate that in soil, the Agr system controls private goods rather than public goods.
REGULATORY RNA ARISES FROM 3´-UTR OF AHL SENSOR ENCODED RNA AFTER ITS PREPROCESSING IN PECTOBACTERIUM ATROSEPTICUM

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Background
In a broad group of proteobacteria two quorum sensing-related genes, which encode synthase of autoinducers and their sensor, respectively, have convergent topology and overlap by their 3´-ends. It was suggested that the expression of one gene may antagonize the transcription of another one because of the convergent arrangement of these genes.

Objectives
We proposed that the topology of quorum sensing-related genes of plant pathogenic bacterium Pectobacterium atrosepticum, expI and expR, determines the mechanism of regulation of their expression.

Methods
We noticed that expR gene lacks any obvious transcriptional stop signals. Using chain-specific RT-qPCR we found that at the stationary growth phase expR mRNAs having long 3´-untranslated region (UTR) were formed and the expression of oppositely oriented expI gene was simultaneously decreased. It is likely, that RNA products of the expR regulatory gene may inhibit the expression of the autoinducer synthase gene through the interactions with expI mRNA. Using 3´-RACE method we found that at the stationary growth phase the extended expR transcripts underwent preprocessing resulting in their break down into two fragments. The first fragment encompassed the entire length of open reading frame (ORF) of expR gene and the second one included only a long 3´-UTR having a predicted regulatory function.

Conclusions
We found that cis-encoded RNA may participate in the regulation of expression of two convergent functionally related genes. The length of one of two transcripts varies respective to the cell growth phase affecting the expression of oppositely oriented gene. Moreover, cis-encoded regulatory RNAs arises from protein encoded RNAs that undergo preprocessing in their 3´-UTR.
METABOLIC TRANSFORMATIONS ENHANCES PSEUDOMONAS AERUGINOSA GROWTH ACTIVITY DURING INTERACTION WITH STAPHYLOCOCCUS AUREUS

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Background
Interactions among members of polymicrobial infections can result in altered pathogen behaviours such as enhanced virulence, biofilm formation or antibiotic tolerance, which may influence the disease phenotype and clinical outcome of the infection. Pseudomonas aeruginosa and Staphylococcus aureus are important opportunistic human pathogens and are both part of the polymicrobial infection communities in human hosts. Cell-cell communication can play a major role in the interaction behaviors between these bacteria, which subsequently can affect the physiology of the individual strains. However, the extent to which evolutionary processes may remodel interspecies interactions and bacterial physiology during the course of infection and therapy is currently not understood.

Objectives
Therefore, the aim of this study was to analyze in vitro interactions between S. aureus and a human host adapted P. aeruginosa strain from a dominant lineage, DK2, that have evolved through decades of growth in chronically infected patients.

Methods
By using a combination of in vitro agar assays, molecular genetics and MALDI-TOF imaging mass spectrometry we demonstrate that DK2 – in contrast to other P. aeruginosa strains - show a commensal-like interaction with S. aureus during in vitro co-culturing, where the growth activity of P. aeruginosa was increased in the presence of S. aureus. The commensal interaction was observed both under aerobic and anaerobic conditions as well as during growth on artificial CF sputum medium, however was abolished with S. aureus strains mutated in the agr quorum sensing system.

Conclusions
Our results suggest that metabolic transformations during interaction with S. aureus enhance the growth activity of P. aeruginosa.
Background

Many of the world’s most precious artworks are made of stone. Their irreversible deterioration due to biological attack is a worldwide concern. Microorganisms colonize outdoor lithic surfaces and develop into biofilms at the interface solid/air (subaerial biofilms, SABs), which, in turn might cause aesthetic, chemical and physical decay. Although it has been estimated that at least 99% of the world’s microbial biomass exists in biofilms, the role and behavior of microorganisms within the biofilm matrix and their complex interactions with the external environment is still unknown.

Objectives

This work provides a pioneering and multidisciplinary research to investigate the behavior of microorganisms within the biofilm matrix for sorting out time-spatial relationships and to elucidate microorganism-EPS, inter-organism, biofilm-atmosphere and biofilm-stone interactions.

Methods

This work spans sophisticated molecular, chemical, physical and data modeling techniques and it is approached from two complementary angles:
1- Lab-scale study to delineate specific transcriptional responses of mono- and multi-species biofilms as well as the biofilm-stone interactions under controlled environmental conditions.
2- Real heritage case studies to investigate the shifts in the microbial community structure and function under different environmental conditions. Through comparing phylogenetic and functional diversity under different environmental scenarios, we
provide evidence that any intuition gained from the lab-scale experiments is relevant to true environmental biofilms.

Conclusions

The findings obtained so far will contribute to better understand the complexity of all the interactions encountered within SAB communities, and how these interactions may influence the biofilm outcome and the biodeterioration of the stone materials under different environmental conditions.
Background

The reduction of nitrite to NO is considered the defining step of denitrification and a point of no return due to the toxicity of NO. Nevertheless, some accumulation of NO is necessary since it is central to the induction of its reductase and frequently also its enzymatic origin. A positive feedback loop perpetuated by a toxic substance is intuitively a play with fire. However, phenotypic studies of organisms known, or assumed to hold a product induced nitrite reductase have with few exceptions revealed stringent NO control.

Objectives

We studied the gas kinetics of \textit{P. denitrificans} during the oxic-anoxic transition in order to reveal new aspects of NO as a regulator of denitrification.

Methods

The core instrumentation was a semi-automatic incubation system similar to that described by Molstad \textit{et al.} (2007), where relevant headspace gases are monitored in respiring batch cultures.

Conclusions

In cultures incubated at 20°C, [NO] reached a semi-steady state at \textasciitilde15nM. This homeostasis could almost entirely be explained by an estimated \textit{in vivo} \(K_m\) of 50nM for NO reductase. Although assumedly eliminating toxic effects, upholding such low [NO] appears to have some drawbacks. By combining experimental data with modelling approaches, we found strong evidence that the gross majority of cells were essentially entrapped in anoxia due to delayed induction of nitrite reductase. In contrast, when incubated at 30°C, [NO] was elevated (\textasciitilde75nM) and the entire population apparently succeeded in switching to N-oxide reduction. This illustrates some of the challenges of wielding a substance with such a double-edged nature: toxic intermediate and essential regulator in one.
METHANOL AS A SUBSTRATE FOR METHANOGENESIS OF THE PRINCIPAL HUMAN GUT ARCHAEOAL COMMENSAL METHANOBREVIBACTER SMITHII

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Background
Methanobrevibacter smithii (Mb. smithii) is the predominant methanogen of the human gut microbiota using CO₂ and H₂ (or formate) for methanogenesis. Other gut methanogens utilize methanol (Methanosphaera stadtmanae) and methylamines (members of newly identified order Methanomassiliicoccales) by H₂-dependent methanogenesis.

Objectives
Genomic analysis revealed that genes encoding the methanol-utilizing methanogenesis pathway (mtaABC) are present in most of the human gut strains of Mb. smithii, encompassing the completely sequenced DSMZ 861 strain. This arises the possibility that this methanobacteriale is also able to use methanol as substrate in the gut. This hypothesis was therefore tested.

Methods
The reference strain DSMZ 861 (ATCC 35061) was obtained from DSMZ (Braunschweig, Germany) and was cultured in DSM141 media under different conditions regards to H₂, CO₂ and methanol. Gene expression of mtaB and mcrA genes (involved respectively in the methanol use and in the final common step of the methanogenesis pathways) were quantitatively evaluated by qPCR.

Conclusions
The DSMZ 861 strain originating from the human gut is able to perform methanogenesis with methanol and H₂. A lower methane production rate was recorded when compared to normal hydrogenotrophic pathway. Methanol induces the expression of the mtaB gene while mcrA relative expression was lowered. This capability to use methanol coupled with its standard hydrogenotrophic methanogenesis provides a clue to explain the predominance of Mb. smithii over other methanogens in the human gut.
SIGMA54-DEPENDENT REGULATION OF P-ETHYLPHENOL AND P-HYDROXYACETOPHENONE DEGRADATION IN “AROMATOLEUM AROMATICUM” EBN1

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Background
Anaerobic $p$-ethylphenol and $p$-hydroxyacetophenone degradation in the denitrifying bacterium “Aromatoleum aromaticum” EbN1 is predicted to be transcriptionally controlled by the sigma54-dependent regulator EbA324. The latter is encoded in between the two operons comprising genes of $p$-ethylphenol/$p$-hydroxyacetophenone degradation and detoxification.

Objectives
Generation of an unmarked, in-frame deletion mutant (strain EbN1 ΔebA324) to validate EbA324 as mediator of substrate-specific transcriptional activation.

Methods
The established genetic system for strain EbN1 was applied to generate deletion mutant strain ΔebA324 and the corresponding complementation strain ΔebA324::ebA324. Both mutant strains and the wild type were physiologically analyzed applying $p$-ethylphenol, $p$-hydroxyacetophenone, benzoate or a mixture of benzoate-$p$-hydroxyacetophenone and subsequently studied on the transcript and protein level.

Conclusions
The ΔebA324 mutant was unable to grow with both $p$-ethylphenol and $p$-hydroxyacetophenone. Accordingly, no transcripts of the $p$-ethylphenol operons were detected in the ΔebA324 mutant using benzoate as growth promoting co-substrate. In contrast, transcripts and proteins of both operons were detected in the wild type and complemented mutant strain, supporting the proposed function of EbA324.

Furthermore, transcriptional activation by EbA324 depends on the presence of either $p$-ethylphenol or $p$-hydroxy-acetophenone: high level constitutive formation of EbA324 in the complemented mutant only yields operon expression in the presence of these compounds.
ADAPTATION OF LISTERIA WEIHENSTEPHANENSIS TO ANAEROBIOSIS

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Background

Listeria weihenstephanensis is a recently new described Listeria specie. It was isolated from the water plant Lemna trisulca from a fresh water pond in Bavaria, Germany. The isolate is non-haemolytic and has not been associated with animal or human diseases. It has an optimal growth at pH 7-8 and temperature at 28-34 °C. It requires a relative low demand of nutrients and is able to grow under both aerobic and anaerobic conditions.

Objectives

In the present study, the focus was the investigation of the adaptation of L. weihenstephanensis to oxygen availability, with respect to the temperature-dependent influence.

Methods

Growth analyses for L. weihenstephanensis were performed aerobically and anaerobically in BHI medium at 18 °C and 34 °C. Global transcriptional analyses, via next generation RNA sequencing, were then performed for cells grown aerobically or anaerobically to an OD _600_ = 0.85-0.90. The oxygen dependent gene expression of selected genes was further validated via qPCR.

Conclusions

At 18 °C, 49 genes were found to be stronger transcribed aerobically while 40 genes were found to be stronger transcribed anaerobically. At 34 °C, 52 genes exhibited aerobically a stronger transcription while transcription of 27 genes was induced anaerobically. Among the differently regulated genes, many encode for metabolic enzymes indicating broad metabolic adaptations with respect to oxygen availability. Moreover, the transcriptional results clarified that temperature does not play a key role in this oxygen-dependent regulation for L. weihenstephanensis.
C4-DICARBOXYLATE TRANSPORT AND GLOBAL TRANSCRIPTOME ANALYSIS OF C4-DICARBOXYLATE METABOLISM IN ACTINOBACILLUS SUCCINOGENES

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Background

Actinobacillus succinogenes that is known to produce large amounts of succinate during hexose fermentation was able to grow on C4-dicarboxylates like fumarate under aerobic and anaerobic condition. Succinate was the major product from anaerobic growth on fumarate, similar to growth on glucose. It indicates the involvement of fumarate respiration and the presence of proteins for C4-dicarboxylate transport.

Objectives

This study aims to characterize transport properties related to C4-dicarboxylate metabolism of A. succinogenes.

Methods

Transport of 14C-fumarate and 14C-succinate were studied with bacterial cells of A. succinogenes grown on fumarate under anaerobic condition. Transport properties were determined concentration-, time-, pH-dependently or in the presence of ionophores.

Global transcriptome was analyzed by total-RNA-sequencing using MiSeq platform. The total RNAs were isolated from cells grown on fumarate or glucose under aerobic or anaerobic condition.

Conclusions

Transport assay revealed that A. succinogenes has several transport systems representing specific uptake activities dependent on concentration range of for C4-dicarboxylates. Some of them required proton potential, but the others Na⁺ gradient.
The genome of *A. succinogenes* possesses candidate genes encoding 18 different transport systems. Global transcriptome analysis showed that six transport systems are expressed during anaerobic growth on fumarate, which includes transporters of the Dcu, DcuC, DASS, and TRAP family.

Overall, *A. succinogenes* could utilize six different protein systems for anaerobic C₄-dicarboxylates transport, according to availability of driving force and substrate concentration.
AN UNUSUAL METHANOGENIC PATHWAY FOUND IN Methanomassiliicoccus luminyensis
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Background

Methanomassiliicoccus (Mmc.) luminyensis is a methanogenic archaeon and was first isolated from human feces. The organism belongs to the recently discovered order of the Methanomassiliicoccales and uses methylated compounds together with H₂ as substrates, which is distinct from other methanogenic pathways.

Objectives

Mmc. luminyensis shows unique metabolic features since it combines different methanogenic strategies in the course of heterodisulfide reduction possessing both, a soluble heterodisulfide reductase (HdrABC) and HdrD as compound of the membrane bound heterodisulfide reductase found in members of the order Methanosarcinales. Therefore the heterodisulfide reduction system of Mmc. luminyensis is of great scientific interest.

Methods

In a novel approach using quantitative reverse transcriptase PCR we identified key enzymes of the methanogenic pathway of Mmc. luminyensis. To further investigate the role of membrane bound enzymes during methanogenesis membranes of Mmc. luminyensis were isolated and tested for the oxidation or reduction of artificial electron mediators such as viologen dyes. Additionally enzymes from Mmc. luminyensis will be produced heterologously and characterized in order to elucidate the methanogenic metabolism of the organism in detail. A special focus will be placed on the headless F₄₂₀H₂ dehydrogenase (Fpo) complex lacking subunit FpoF, which is essential for F₄₂₀H₂ oxidation, and HdrD since the HdrE subunit of the usually membrane bound heterodisulfide reductase is missing.

Conclusions

If those enzymes are active despite their deficiency of certain key subunits Mmc. luminyensis could employ a new methanogenic pathway combining classical enzymes known from other methanogens in a so far not described manner.
REDUCTIVE DECHLORINATION OF EITHER TETRACHLOROETHENE OR 1,2-DICHLOROETHANE BY DEHALOGENASES WITH SIMILAR PROPERTIES BUT DIFFERENT REACTION MECHANISMS

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Background
Organohalide respiring bacteria are able to use halogenated organic compounds as terminal electron acceptors for growth. The key enzyme for the conversion of these substrates is the reductive dehalogenase (RDase), a cobamide-containing iron-sulfur protein. Biochemical and structural data showed the cobamide cofactor as the reactive species in the RDase's active site [1]. Numerous RDase protein sequences are listed in databases [2], but only a few enzymes were biochemically characterized. The RDase of Sulfurospirillum multivorans, PceA, converts tetrachloroethene via trichloroethene to cis-1,2-dichloroethene in a hydrogenolysis reaction [3]. In contrast, the RDase of Desulfitobacterium dichloroeliminans, DcaA, catalyzes the formation of ethene from 1,2-dichloroethane by a dihaloelimination mechanism [4].

Objectives
The structural determinants in RDases will be identified, which are responsible for the catalysis of either a hydrogenolysis or a dihaloelimination reaction.

Methods
In silico structural analysis of DcaA based on the crystal structure of PceA is conducted. The substrate spectrum of recombinant DcaA is determined via enzyme activity measurements. Site-directed mutagenesis is applied to unravel the role of amino acids forming the DcaA’s active site.

Conclusions
From the results of this study, a better understanding of the diversity in RDase substrate spectra and reaction mechanisms and an improved classification of unknown RDases based on their protein sequence is expected.

THE ANAEROBIC LINALOOL METABOLISM OF THAUERA LINALOOLENTIS 47LOL
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Background
Thauera linaloolentis 47Lol\textsuperscript{T} grows on linalool (3,7-dimethylocta-1,6-dien-3-ol), a tertiary monoterpene alcohol, as sole carbon and energy source under denitrifying conditions. Linalool mineralization is initiated by isomerization into the primary alcohol geraniol by a novel enzyme, a linalool isomerase (Foss and Harder, 1997, Foss and Harder, 1998). Geraniol degradation may proceed in analogy to the acyclic terpene utilization (ATU) pathway, as described for various Pseudomonas strains (Förster-Fromme and Jendrossek, 2010). This pathway involves an oxidation of geraniol to geranic acid. After activation by coenzyme A thioester formation, the monoterpene structure is shortened by beta-oxidation, and rearrangement to 3-methylcrotonyl-CoA, ultimately entering the leucine degradation pathway (LIU).

Objectives
The objective of this study was to elucidate the anaerobic linalool metabolism in Thauera linaloolentis 47Lol.

Methods
Recently, draft genomes of T. linaloolentis 47Lol became available and we detected candidate genes of the ATU and LIU pathways in the genome. Expressed proteins present in linalool-grown cells included many corresponding proteins.

Conclusions
So far, the ATU pathway has only been described in various Pseudomonas strains growing aerobically. Our findings are the first observation for the utilization of the ATU pathway under anaerobic conditions.
BONCAT TAGGING FOR DIRECT QUANTIFICATION OF RESPIRING AND NON-RESPIRING SUBPOPULATIONS IN PARACOCCUS DENITRIFICANS DURING TRANSITIONS FROM AEROBIC RESPIRATION TO DENITRIFICATION

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Background
Ecophysiological studies have revealed that the regulation of denitrification pathways is highly variable among bacterial strains. One aspect of the regulatory phenotype is the maintenance and distribution of the denitrification proteome within clonal populations. Gas kinetics studies of batch cultures during the transition from aerobic respiration to denitrification have indicated that in many organisms there is a uniform onset of nitric oxide reduction amongst cells in the population. Mathematical modelling suggests that the model organism for denitrification studies, Paracoccus denitrificans, has a different strategy reminiscent of bet-hedging where only a subset of cells (Fden) start to denitrify, and thus continue to grow, upon oxygen depletion [1].

Objectives
We aim to estimate the proportion of cells successfully switching to denitrification by quantifying directly the cells that are showing translational activity after the onset of denitrifying conditions.

Methods
The bioorthogonal noncanonical amino acid tagging (BONCAT) approach takes advantage of chemical groups that react with each other at near-physiological conditions, but not with other cell components. These are appended to the bacterial proteome during translation using supplementation of a noncanonical amino acid, in this case azidohomoalanine (AHA). A fluorescent dye displaying a complementary moiety is then used to visualize cells displaying the modified proteins.

Conclusions
The BONCAT method was shown to be successful in differentiating between fully respiring cells and those that were inactive, only showing low background fluorescence. Studies are currently undertaken to determine Fden for P. denitrificans cultures undergoing the transition from aerobic to denitrifying growth.

Anaerobic physiology

A NOVEL ANAEROBIC TOLUENE-DEGRADING BACTERIAL STRAIN OF CLOSTRIDIACEAE FROM AN OIL-CONTAMINATED SOIL

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Background

Anaerobic degradation of aromatic hydrocarbons is important for bioremediation of contaminated subsurface environments. Information on the microbial diversity and metabolic pathways involved in anaerobic degradation of hydrocarbons is crucial for successful cleanup of hydrocarbons in subsurface. Although members of the family Clostridiaceae have been frequently observed in the analysis of various hydrocarbon-contaminated microbial communities, there are limited studies on the biodegradation potential of Clostridiaceae.

Objectives

The aims of this study are to isolate hydrocarbon degrading strains of Clostridiaceae and elucidate the biochemical and genomic properties of the strain regarding anaerobic biodegradation.

Methods

Enrichment cultures inoculated with various soils from oil spill sites were incubated for several years with petroleum amendment. After enrichment, several colonies were obtained using the roll tube method. Among the isolates, one strain, designated IRF9, was found to degrade toluene and thus selected for further biochemical and genomic analysis.

Conclusions

An alkalipilic and heterotrophic aromatic hydrocarbon-degrading bacterial strain, IRF9, was isolated from the enrichment cultures. Cells of strain IRF9 were straight or curved rod-shaped, motile, and Gram-negative. Optimal growth of strain IRF9 was observed at pH 9.0–9.5 and 40 °C. The strain was found to grow within pH and temperature ranges of 6.5–10.0 and 25–45 °C, respectively. Ferric iron was used as an electron acceptor but not sulfate, nitrate. The biochemical and genomic studies of the strain IRF9 will shed light on the potential of Clostridiaceae for the anaerobic biodegradation of hydrocarbons in subsurface environments.
Background
The photosynthetic purple sulphur bacterium, *Thiocapsa roseopersicina* harbours four functional [NiFe] hydrogenases. Two of them are attached to the periplasmic membrane (HynSL, HupSL) and the other two are localized in the cytoplasm (Hox1, Hox2). It prefers to utilize reduced sulphur compounds for anaerobic phototrophic growth.

Objectives
HupSL is a hydrogen uptake hydrogenase which can recycle hydrogen in the environment and provides energy for the cells. The gene expression and enzyme activity correlate with the applied thiosulphate concentration, HupSL shows increased hydrogen uptake activity under low sodium-thiosulphate concentration. The hydrogenase small and large subunits are encoded by the *hupS* and *hupL* genes in the *hupSLCDHIOR* operon. The function of *hupO* gene and its product was completely unknown. In order to investigate the physiological role of the HupO, the *hupO* gene was inactivated by in-frame deletion mutagenesis. The *hupO* gene was deleted in the ∆hynSL, ∆hox1H, ∆hox2H (GB113141) mutant (resulting strain HOD134). The effects of the mutation were analyzed by activity and expression measurements.

Methods
The *in vivo* activity of the HupSL hydrogenase in the HOD134 strain substantially increased under low sodium thiosulphate concentration compared to the wild-type and complemented strains. These activity measurements were supported by Western analyses and gene expression studies.

Conclusions
This study adds to the clarification of the so far unresolved HupSL regulation. The HupO plays a crucial role in this regulation at least under low thiosulphate conditions, where the masking effect of thiosulphate is avoided.
EFFECT OF CARBON DIOXIDE ON EXOPOLYSACCHARIDE PRODUCTION AND SURVIVAL OF BIFIDOBACTERIUM BIFIDUM AFTER FREEZE-DRYING
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Background

*Bifidobacteria* are particularly sensitive to dissolved oxygen potentially present in bioreactor during the production process [1]. However, oxygen stripping by another gas (i.e., nitrogen and hydrogen) decreases the oxido-reductive potential and improved survival of *B. bifidum* during storage [2]. It is noted that carbon dioxide can not only maintain the anaerobic growth conditions of *Bifidobacterium longum* JBL05 but also enhance the cell concentrations and the exopolysaccharide (EPS) secretion [1]. Moreover, EPS was known to be an important factor for the protection of *B. bifidum* during freeze-drying [3].

Objectives

Optimization of oxygen stripping by using different gases (i.e., CO₂ and N₂) to reduce the redox potential in *B. bifidum* THT0101 culture aimed at (i) improving the survival of cells during freeze-drying and (ii) increasing the EPS secretion during the fermentation.

Methods

*B. bifidum* THT0101 was cultured following different degassing procedures. After culturing and freeze-drying, the concentration of exopolysaccharide and the viability of *B. bifidum* THT0101 were evaluated.

The results pointed out a significant increase of viability (5.06-15.6 folds by comparison with controls) in relation with the duration of degassing by CO₂. Besides, the CO₂ treatments led to a decrease of the redox potential. The concentration of exopolysaccharide increased related to the flow rate of CO₂ and *B. bifidum* THT 0101 survival.

Conclusions

The flow rate of CO₂ in culture led to the increase of exopolysaccharide concentration and *B. bifidum* THT0101 survival.
Background
Terpenes are natural hydrocarbons mainly produced by plants. Monoterpenes are released to the environment in teragram amounts every year. Castellaniella defragrans 65Phen, a betaproteobacterium able to grow under denitrifying conditions, uses several aliphatic and cyclic monoterpenes as sole carbon and energy sources (1). This metabolic capability is currently studied intensively and the genome has recently been described (2).

Objectives
Identify genes and proteins relevant to the metabolism of the bicyclic monoterpenes 3-carene, sabinene and alpha-pinene.

Methods
Random transposon mutagenesis and differential proteomics were used.

Conclusions
Firstly, in our study the bicyclic monoterpenes 3-carene, sabinene and alpha-pinene seemed to be isomerized to monocyclic monoterpenes and then further degraded by proteins encoded in a 70 kb genetic island (2). The nucleotide composition of the island is indicative for horizontal gene transfer from several other hydrocarbon degraders to C. defragrans 65Phen (2). Beta-oxidation of perillic acid and the leucine/isovalerate (liu) pathway were also involved in the metabolic steps of all monoterpenes under investigation. Additionally, several permeases and stress-response proteins had a relevant role in the survival and growth of C. defragrans 65Phen on monoterpenes, most likely due to the monoterpenes toxicity.

Background
Generally, sulfate-reducing bacteria (SRB) grow optimally at neutral pH but sulfate reduction at low pH has been reported in both natural and engineered environments. Despite evidence of their activity in situ, acidophilic SRB have been difficult to isolate and cultivate in vitro, possibly due to the inhibitory effect at low pH of metabolites such as sulfide or acetic acid. Thus far, only three species have been formally described: Desulfosporosinus acidiphilus, D. acididurans and Thermodesulfobium narugense.

Objectives
Our objective was to isolate novel acidophilic SRB to study their ecophysiology.

Methods
Enrichments were set up using sediments from an acid rock drainage environment (Tinto River) and several isolation methods were applied (enrichment, serial dilution and pasteurization).

Conclusions
The isolate, strain I, represents a novel lineage of SRB within the Firmicutes phylum, proposed as Desulfobacillus spp. (93% similarity with the genera Desulfosporosinus and Desulfitobacterium). The genome-guided characterization shows a very versatile metabolism; it is able to use multiple electron donors and electron acceptors for respiratory growth, and to perform fermentation and disproportionation. Strain I shows a unique metabolism compared to other acidophilic SRB by being able to oxidize acetic acid. As a consequence, strain I is the most acidophilic strain described so far, able to grow while reducing sulfate at a pH as low as 3.8 (optimum at pH 5).

Its high tolerance to low pH and its acetate detoxifying metabolism together with the alkalinity generation coupled to growth of SRB suggest its important role in promoting microbial niches with milder conditions in acidic environments.
CONVERSION OF CIS-2-CARBOXYCYCLOHEXYLACETIC ACID IN THE DOWN-STREAM PATHWAY OF ANAEROBIC NAPHTHALENE DEGRADATION

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Background
The ability of anaerobic naphthalene degradation has been demonstrated for the deltaproteobacterial strains N47 and NaphS2 under sulfate-reducing conditions (Galushko et al., 1999, Meckenstock et al., 2000). It was shown recently that after carboxylation to 2-naphthoic acid (Mouttaki et al., 2012) and formation of the corresponding CoA-ester, the latter is stepwise reduced to a hexahydro-2-naphthoyl-CoA with unknown conformation of the remaining double bonds (Eberlein et al., 2013). Furthermore, cis-2-carboxycyclohexylacetic acid could be identified as a metabolite of the down-stream pathway (Annweiler et al., 2002).

Objectives
Our aim for this study was the elucidation of the next metabolites of the pathway following cis-2-carboxycyclohexylacetyl-CoA and the identification of the enzyme reactions finally leading to the second ring-cleavage.

Methods
The metabolite cis-2-carboxycyclohexylacetic acid was chemically synthesised and converted to the corresponding coenzyme A thioester. The latter was tested for conversion in cell-free-extracts of N47 and NaphS2 in the presence of different potential co-factors and electron acceptors. Samples were analysed by HPLC and emerging metabolites were further characterised via GC-MS.

Conclusions
Conversion of cis-2-carboxycyclohexylacetyl-CoA was only observed in the presence of ferrocenium hexafluorophosphate, which can oxidise enzyme-bound co-factors like FADH₂. The first emerging metabolite had a double-bond introduced, indicating the activity of a FAD-dependent acyl-CoA dehydrogenase. Also the product of the next enzyme, probably a hydratase, could be detected. Combining these data, we can for the first time make a detailed proposal for the steps leading to the second ring-cleavage in the down-stream pathway of anaerobic naphthalene degradation.
EXPLORING A RICE STRAW DEGRADING MICROBIAL COMMUNITY ENRICHED FROM SWINE MANURE DIGESTER MATERIAL

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Background
Rice straw is one of the most abundant agricultural waste residues in the world. Currently it is burned in the fields, which causes pollution and greenhouse gas emission, whereas it has potential as feed for biogas production.

Objectives
To understand and stimulate biogas production from rice straw, we aim to isolate novel microbes with metabolic capacities that are key in the degradation of rice straw. Ingham et al. (2007) developed the MicroDish Culture Chip (MDCC) for cultivation of micro-colonies.

Methods
We set up a method for using the MDCC in anaerobic conditions which allows for screening key capacities and studying microbial interactions.

Conclusions
After five weeks incubation of anaerobic liquid cultures at 37°C, 75-80% of the rice straw dry weight was degraded and 250 L*kg VS⁻¹ methane was formed, and acetate, propionate, benzoate and 3-phenyl propionate were measured as intermediates. The rice straw-degrading community was recently transferred to specific substrates. After 25 days of incubation methane was formed from cellulose and xylan, but not from lignin.

Reference
Background
Plasmid gene delivery vectors offer several advantages over viral vectors, including safety, lower toxicity and easier preparation. Selective marker genes coding for antibiotic resistance have been traditionally an essential part of these plasmids. However, health authorities recommended the use of plasmids without resistance markers in order to prevent the spread of resistance genes.

Objectives
The aim of our study was to determine the possibility, frequency and efficiency of therapeutic plasmid pORF-hIL12 AmpR transformation into different bacterial strains.

Methods
The strains used were selected from 69 isolates belonging to 17 genera, which were isolated from dogs skin before application of electrogene therapy (EGT) (1) and from 89 isolates belonging to 17 genera retrieved after EGT. Several enterobacterial strains were also included into the study. Transformation of electrocompetent bacteria from genera Staphylococcus, Bacillus, Acinetobacter, Kocuria, Micrococcus, Rhodococcus, Streptomyces, Mycobacterium and Arthrobacter yielded no transformants with pORF-hIL12. The transformation frequency (efficiency) of Escherichia coli, E. hermanii, Buttiauxella sp., Salmonella eneterica, Proteus vulgaris, Erwinia chrysanthemii, Yersinia sp., Providencia sp., Enterobacter sp., Shigella boydii, Herbaspirillum frisingense and Stingmonas sp. ranged between $10^{-4}$ and $10^{-10}$ ($10^1$ and $10^5$).

Conclusions
Our study revealed that pORF-hIL12 could be introduced into several enterobacterial species, if they are present on the dogs skin and naturally competent. Thus a potential risk for the spread of the ampicillin resistance gene from pORF-hIL12 by horizontal gene transfer can not be excluded.

Reference
IN VITRO SUSCEPTIBILITY ACTIVITIES OF EIGHT ANTIFUNGAL DRUGS AGAINST CLINICAL AND ENVIRONMENTAL ISOLATES OF PHAEOACREMONIUM SPECIES

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Background

Phaeoacremonium species are chiefly found in the environment and can either cause a plant disease which is an uncommon cause of human disease. Most cases of Phaeoacremonium infection in humans involve traumatic inoculation and present as subcutaneous abscesses, cysts, or osteoarthritis. The spectrum of disease caused by Phaeoacremonium species especially *P. parasiticum* is variable and ranges from subcutaneous infections to eumycetoma, arthritis, osteomyelitis, endocarditis, fungemia and disseminated disease.

Objectives

Despite worrying clinical pictures associated with *Phaeoacremonium*, there is little information regarding the antifungal susceptibility patterns against currently available antifungal agents.

Methods

Thus, the *in vitro* activities of eight antifungal agents against clinical and environmental strains of *Phaeoacremonium* were obtained. The Clinical and Laboratory Standards Institute (CLSI) provides no specific guidelines for testing the *in vitro* antifungal susceptibility of this fungus. No other testing protocol has also been validated for testing the susceptibility of *Phaeoacremonium*.

Conclusions

In the present study, the resulting MIC₉₀ₐₛ for all strains were as follows, in increasing order: POS (0.125 μg/ml); VOR (0.5 μg/ml); AmB (0.5 μg/ml); ISA (1 μg/ml); CAS (8 μg/ml); ANID (8 μg/ml); ITC (16 μg/ml); and FLU (64 μg/ml). The difference in the MIC₉₀ₐₛ between the two groups of isolates did not differ by more than one
Therefore, the present study based on *in vitro* activity showed that posaconazole and voriconazole might have a potent activity with a best choice of alternative to amphotericin B. In
IMPACT OF WASTEWATER DISCHARGES IN SOILS: DEFINITION OF LIMITS OF DETECTION AND QUANTIFICATION OF ANTIBIOTIC RESISTANCE GENES

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Background
Wastewater can contribute to the spread of antibiotic resistant bacteria and resistance genes (ARB&G) in the environment. The discharge of treated wastewater in soils may lead to the accumulation of different contaminants, including ARB&G. However, due to technical limitations and low abundance of ARB&G in water, it is possible that the effects remain unnoticed before a long term exposure has occurred.

Objectives
To test this hypothesis, we designed an experiment to assess the limits of detection of ARG in soil.

Methods
An agricultural soil was soaked with a synthetic effluent containing two ARB to reach bacterial densities from $10^2$ up to $10^9$ CFU/g. ARB were *Escherichia coli* strain A2FCC14, which harbours the genes *bla*TEM, *bla*CTX-M-15, *bla*OXA-1, aac(6')-Ib-cr and integrase IntI-1, and *Enterococcus faecalis* H1EV10, with the *vanA* gene. ARB&G detection limits were assessed based on culture-dependent and culture-independent methods (PCR and qPCR).

Conclusions
The limit of detection by conventional PCR was gene-dependent and corresponded to bacterial densities between $10^4$ (in the case of aac(6')-Ib-cr) and $10^7$ CFU/g soil (in the case of *bla*CTX-M-15 and *bla*OXA-1). Using qPCR, the quantification limit corresponded to bacterial densities of $10^3$ CFU/g soil, although detection was possible for lower genes abundances. However, these values can be considered high for certain up to now uncommon ARG, which can spread in soils remaining unnoticed. The relatively high detection limits observed will be discussed, highlighting the meaning of this shortcoming for an adequate risk assessment of the implications of wastewater discharges in soil.
Background

Evolution of bacterial antibiotic resistance represents one of the most threatening health care problems worldwide. This problem urgently requires a better understanding of important reservoirs and paths of dissemination, such as those represented by wastewater treatment and reuse.

Objectives

While coordinated European Surveillance programs exist for clinically-associated antibiotic resistance, equivalent programs addressing the (aquatic) environments are missing. The aim of this study was to obtain a first insight into the abundance of antibiotic resistance genes in the treated effluent discharged by urban wastewater treatment plants (UWTP) from distinct geographic areas. This is the first pan-European synopsis on the levels of antibiotic resistance genes in treated wastewater.
Methods

Within the 2014 WG5 Norman campaign we screened 20 (UWTP) in 13 different nations in Europe to screen for five environmentally and medically relevant antibiotic resistance and related genes (ARG, blaTEM; vanA; qnrS; sul1; ctx-m 32 and intI1). For three consecutive days, 24h-composite samples were collected and processed immediately for DNA extraction.

Conclusions

All analysed ARG were detected in samples from all countries, with differences between different UWTP effluents, suggesting that the investigated determinants are adequate indicators. A north to south gradient within Europe could be observed, with the northern UWTPs showing the lower abundance of ARGs in the effluent than southern countries. This study sets the base for further studies investigating correlations between resistances and treatment processes, and linking environmental and clinical occurrence.
VIROME-ASSOCIATED ANTIBIOTIC RESISTANCE GENES IN FRESHWATER AQUACULTURE

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Background

Due to their structural characteristics in extracellular phase, bacteriophages persist quite successfully in the environment and are resistant to natural or man generated stressors. Aquatic compartments may therefore play a significant role in driving antibiotic resistance genes (ARGs) transfer.

Objectives

The project is aimed to highlight the impact of viral populations in the spreading of ARGs in aquaculture.

Methods
Upstream and downstream water samples from an experimental salmonid aquaculture were treated with a tangential flow filtration system to separate and concentrate viral-like particles and microbial cells. Then viromes were investigated through shotgun NGS while DNA from microbial communities underwent NGS 16S rRNA gene profiling. Bioinformatic analysis was performed using QIIME and Meta-Vir tools, CARD, NCBI, and COG database. The AR colonies frequency was evaluated by standard plating.

Conclusions

Upstream and downstream samples were characterized by limited changes in viral community composition at family level, while a large variation in bacterial population was observed. The relative abundance of AR drug classes did not significantly change whereas the taxonomic distribution of ARGs and bacterial metabolic genes in the viromes were remarkably different between upstream and downstream samples thereby reflecting the taxonomic composition and diversity of the bacterial communities. Interestingly, an inverse relationship between the relative abundance of genes belonging to a bacterial family in the virome and the same bacterial family abundance in the microbioma was observed. Furthermore, the frequency of AR resistant colonies showed an inverse relationship with respect to the abundance of specific virome-associated ARGs.
CHARACTERIZATION OF YBJG, A PYROPHOSPHATE PHOSPHATASE FROM E. COLI INVOLVED IN THE LIPID CARRIER UNDECAPRENYL PHOSPHATE METABOLISM

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Background

Undecaprenyl phosphate (C₁₅-P) is an essential lipid carrier involved in the biosynthesis of cell surface carbohydrate polymers such as the peptidoglycan. C₅₅-P is the result of the dephosphorylation of the undecaprenyl pyrophosphate (C₅₅-PP) by specific phosphatases. In Escherichia coli this dephosphorylation can be performed by four integral membrane proteins, BacA, and three members of the type 2 phosphatidic acid phosphatase family (PAP2), PgpB, YbjG, and LpxT.

Objectives

The aim of this project is to characterize YbjG and contributes to the understanding of the physiological role and mechanism of action of this enzyme in the C₅₅-P metabolism. The C₅₅-PP phosphatases could become an interesting target in the search for new molecules with antibacterial activity.

Methods

In parallel the stability of YbjG and its activity against C₁₅-PP were assessed in 94 different detergents. Moreover the enzymatic activity of YbjG was studied: substrate specificity, optimum pH and temperature, effect of detergent concentration.

Conclusions

For the first time, YbjG has been purified and we show its ability to dephosphorylate C₁₅-PP, DGPP and C₅₅-PP in vitro with respectively decreasing efficiency. No activity has been detected on five other potential substrates (PPI, PA, C₅-PP, G6P & PNPP).

The phosphatase activity on C₁₅-PP is maximum at pH 6.5 and 25 °C. Moreover Cymal6, LMNG, & ωUDM are good detergent both for the stability and the C₁₅-PP
phosphatase activity of YbjG, but approximately half of the 94 tested detergents show C_{15}-PP phosphatase activity on the qualitative enzymatic test.
Antibiotic resistance and environment

CONSTITUTIVE PRESENCE OF ANTIBIOTIC RESISTANCE GENES WITHIN THE BACTERIAL COMMUNITY OF A LARGE SUBALPINE LAKE

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Background

The fate of antibiotic resistance genes (ARGs) in natural bacterial communities is of primary concern as prodromal of their potential spread within pathogenic bacteria. Although of diverse origin, the persistence of ARGs in aquatic environments is highly influenced by anthropic activities, allowing potential control actions in well-studied environments. However, knowledge of their abundance and dynamics in ecosystems is still scarce.

Objectives

We investigated the presence and the abundance of twelve ARGs in large, subalpine Lake Maggiore (Italy/Switzerland), using qPCR, at different stations, depths and seasons. In order to compare seasonal dynamics to short-term changes we sampled one station daily over the course of a week. We then evaluated correlations between ARGs and the bacterial community composition assessed with Illumina sequencing of the 16S rDNA and a number of environmental parameters, including bacterial density, temperature, chlorophyll, oxygen and water transparency.

Methods

Conclusions

Our results suggest the constitutive presence of at least three to four ARGs within the natural bacterial community, with a high proportion of bacteria potentially resistant to tetracyclines and sulphonamides. However, in some cases we also observed short-lived peaks of ARGs (one day) which also co-occurred with a significant change of the bacterial diversity in the system. These observations, on the one hand propose the lake as an ideal reservoir for the persistence of certain ARGs, and on the other hand it suffers from point pollution with ARGs. This highlights the need for an appropriate management of the antibiotic consumption in the catchment basin and of the potential sources of contamination.
ANTIBIOTIC RESISTANCE OF BACTERIAL STRAINS ISOLATED FROM THE AIR OF.Sorting Facilities

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Background
The spread of resistant strains of bacteria in the community is recognized as a recent trend, and this fact adds new risk factors usually not taken into consideration in particular for professional activities where the exposure to biological agents is larger than usual, as is the case of waste sorting facilities.

Objectives
The study focuses on the characterization of the antibiotic resistance profile of Staphylococcus spp. and presumptive Enterobacteriaceae strains isolated from the air of sorting facilities.

Methods
A total of 49 Staphylococcus spp. strains isolated from air samples were used for the detection of mecA and pvl genes by RT-PCR. Phenotypic resistance has been determined by disk diffusion and agar dilution methods. From an initial screening of 133 air samples using McConkey agar containing meropenem (2mg/L), 37 Gram (-) strains were phenotypically characterized. Strains resistant to meropenem were characterized in terms of resistance to other 15 antibiotics of 7 different classes using the disk diffusion method.

Conclusions
None of the Staphylococcus spp. strains was pvl (+). The presence of the gene mecA was observed in 8% of the Staphylococcus spp. strains tested and 25 of the strains were able to grow in the presence of oxacillin concentrations between 0.5 mg/L and 4 mg/L. The presumptive Enterobacteriaceae strains isolated in the initial screening showed high prevalence of resistance to at least one antibiotic (95%) and 43% of the strains were resistant to 6 different antibiotics. Meropenem resistance was confirmed in 51% of the strains.
Background
River water is one of the most important reservoir and vehicle of bacterial resistance dissemination, in nature. The overexposure of aquatic environments to human and animal intestinal commensals and pathogens, and the selective pressure exerted by factors as antibiotic residues, are important sources of resistance. Our previous work showed multi-resistant ESBL producing E.coli, in river water in Portugal, leading us to improve our research about this public health concern.

Objectives
Our aim was the detection of ESBL and carbapenemase producing Enterobacteriaceae in recreational river waters in the North of Portugal.

Methods
Water samples of 5 different rivers with recreational activities, were analysed by membrane filtration on MacConkey agar and MacConkey Agar with cefotaxime, ciprofloxacin and meropenem. Representatives of different colony morphotypes were tested for antimicrobial susceptibility and carbapenemase screening, according to CLSI and EUCAST. ESBL producers were confirmed by the double-disk-synergy-test and PCR was performed for detection of blaTEM, blaOXA, blaSHV, blaCTX-M-group-1 and aac(6’)-lb-cr genes. Isolates were identified by ID-32-GN.

Conclusions
Our study showed the dissemination of blaCTX-M-group-1 producing Enterobacteriaceae in this aquatic environment and the presence of Gram-negative-non-fermentative bacteria with reduced susceptibility to carbapenems. From 56 isolates analyzed, 33 showed multi-resistance, 17 were ESBL producers. Isolates of Escherichia coli, Klebsiella pneumoniae and Enterobacter spp. showed positive for blaCTX-M-group-1 (13), blaOXA (12), blaTEM (1), blaSHV (6) and aac(6’)-lb-cr (1), genes. Results indicate that recreational river waters of the North of Portugal are reservoirs of ESBL producing Enterobacteriaceae and non-fermentative-Gram-negatives with reduced susceptibility to carbapenems, suggesting a public health concern.
CATTLE AS RESERVOIR OF CIPROFLOXACIN RESISTANT COLIFORMS
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Background

Human intestinal colonization by antimicrobial resistant E.coli, suggests environmental reservoirs as water, food or direct contact with animals. Descriptions of fecal carriage in food-producing animals, might suggest cross-transmission between humans and animals or a common environmental source. Meat can be a source of resistant bacteria that can spread through the food chain.

Objectives

The aim of our study was the detection of ciprofloxacin resistant (CR) coliforms in cattle fecal flora of two different type facilities in the North of Portugal.

Methods

Faecal samples were suspended in TSB and incubated at 37°C overnight. Isolates were selected on MacConkey agar with and without ciprofloxacin. Colonies of lactose fermenters were randomly selected and susceptibility to antimicrobial agents was determined by the agar diffusion test using CLSI guidelines. Strains were identified by ID 32 GN and API 20 E. Genetic information was investigated for qnr and aac(6')-Ib-cr genes, by PCR.

Conclusions

Our study showed the emergence of qnrA and aac(6')-Ib-cr mediated ciprofloxacin resistance in coliforms from Portuguese cattle. CR E.coli isolates were detected in all samples of intensive production. ESBL producers were detected among the CR isolates, and resistance to tetracycline, chloramphenicol, trimethoprim-sulfamethoxazole, β-lactams and β-lactam/inhibitor association, was frequent.

Intestinal flora of cattle might act as reservoir of CR coliforms. Fluoroquinolones used in veterinary suggest a potential for selection of CR coliforms in the faecal flora of
production animals. Bovines can be a source of resistant bacteria colonization through the food chain, by unexpected contamination of carcasses by CR fecal isolates.
Background The spread of antibiotic resistance among bacteria has become one of the most important health problems nowadays. The problem relies on the dissemination of antibiotic resistance genes by conjugative plasmids. Increasing antibiotic concentrations in clinical and natural environments enhances this problem. Many antibacterial compounds have their roots in natural environments, such as soil. As a consequence, bacteria have coevolved and shared resistance mechanisms and genes along evolution.

Objectives This work is focused on the search of new conjugative plasmids from environmental unique soils in which high concentrations of antibiotics are present as a result of animal manure amendments. Additionally, we aim to study the effect of these plasmids on the fitness and phenotype of recipient bacteria.

Methods Exogenous Plasmid Isolation technique was used to collect conjugative plasmids from the bacterial community present in different soils. We obtained a total of 17 E. coli transconjugants from 10 different soils selected with ampicillin, erythromycin, chloramphenicol or streptomycin. To study the effect that the acquisition of these plasmids has on the fitness and phenotype of E. coli transconjugants, we determined growth curves, antibiotic MIC values, and community-level physiological profiles (BIOLOGTM) which provide information on carbon substrate utilization patterns.

Conclusions We conclude that acquired conjugative plasmids affect both fitness and phenotype of recipient bacteria.
EXTRAINTESTINAL PATHOGENIC ESCHERICHIA COLI SEQUENCE TYPE 131 IN HEALTHY BROILER CHICKEN MEAT IN ITALY: A COMBINATION OF ANTIBIOTIC RESISTANCE WITH VIRULENCE

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Background
Extrainestinal pathogenic Escherichia coli sequence type 131 (ExPEC-ST131) has emerged internationally as a multidrug-resistant pathogen. It has been shown that the transmission can occur from animals to humans. Based on existing evidence, poultry is the food animal source most closely linked to human E. coli.

Objectives
The objective was to characterize ExPEC from raw chicken meats purchased from Palermo, Italy.

Methods
In a total of 187 raw chicken meat from retail markets in Palermo, Italy during March-November 2014, selective cultures and disk-diffusion assays for the isolation and characterization of antimicrobial-resistant E. coli were performed. Polymerase chain reaction-based assays were done to detect Extended-Spectrum ß-Lactamases (ESBLs), plasmid-mediated AmpC genes, plasmid-mediated quinolone resistance (PMQR) genes and aac(6')-Ib-cr genes, phylogenetic group typing and to define ExPEC-associated traits. A single nucleotide polymorphism (SNP) PCR was done to detect E. coli ST131.

Conclusions
In total, 179 E.coli Isolates were isolated from 187 chicken meat samples. Fifteen (8.4%) isolates were belong to phylogenetic group B2, at least one of the ESBLs genes and/or PMQR determinants were detected in all of these isolates. Three isolates of group B2 contained aac(6')-Ib-cr. Based on the molecular definition of ExPEC, all of the B2 isolates had the status of ExEPC. The most frequent virulence factors were kpsMT II and iutA. SNP-PCR results showed that eight isolates were ST131 which is the first report from Italy and poses a potential zoonotic risk.
Investigations into the ability of our poultry ExPEC to cause human infections are warranted.
Background
A major contributor to water pollution is the discharge of domestic and industrial waste into the environment. Although developed countries treat wastewater and sludge before releasing it into the environment, many contaminants are not completely removed by the wastewater treatment plants (WWTP) and subsequently end up in environmental waters. There are concerns that these contaminants contribute to increased virulence and/or antibiotic resistance in microbes in the environment and thus pose a health risk.

Objectives
To design and apply a multi-gene expression system for *Pseudomonas aeruginosa* to predict the potential impact of soluble components released from sewage sludge on bacteria.

Methods
Selected genes associated with stress response, virulence and antibiotic resistance were analysed by quantitative PCR in *P. aeruginosa* subjected to tobramycin or heat shock treatments for validation, or sewage sludge leachates from 3 Swedish WWTPs (Eskilstuna, Örebro and Västerås). *P. aeruginosa* was selected because it is normally in the environment and is also an opportunistic pathogen. The molecular profiles of the sludge leachate treatments were subjected to principle component analysis.

Conclusions
The sludge leachate from Eskilstuna significantly affected more *P. aeruginosa* genes, responsible for general stress response and virulence, while sludge leachate from Västerås affected fewer genes and clustered more closely to the control. Örebro sludge leachate produced a different gene expression profile from the other two WWTPs. This suggests that the presented strategy of *P. aeruginosa* toxicogenomics has potential applications in evaluating the effects of soluble contaminants on bacteria in environmental waters for considerations in risk assessment.
Background:

Expanded spectrum cephalosporin resistance continues to be a global threat with regard to infection control management and therapeutic options. *Escherichia coli*, being versatile in host tropism, acts as a major contributor in horizontal and vertical expansion of antibiotic resistance even without external antibiotic pressure. Food remains an important growth media for both pathogenic and non-pathogenic bacteria and a source for genetic exchange between them leading to transfer of antibiotic resistance. Thus, knowledge of these resistance determinants in non-clinical host is critical in tracking down their dissemination in trans specific level.

Objectives:

The current investigation was undertaken to assess the occurrence of class C beta lactamase mediated cephalosporin resistance and their transmission dynamics among non-clinical isolates of *E. coli* in a high altitude city of Northeastern India.

Methods:

A total of 56 *E. coli* were recovered from 61 ready to eat food samples. Phenotypic and molecular characterization of AmpC genes was performed and MIC against oxy-iminocephalosporins was also determined. Isolates positive for *bla*\textsubscript{CMY} was checked with CMY specific primers and amplicons were sequenced. Horizontal transferability was tested by conjugation and transformation. Plasmids were further characterized by PCR based replicon typing.

Conclusions:

A total of 17 isolates were found harbouring *bla*\textsubscript{CMY-96} type. The gene was conjugatively transferable and maintained within W inc type plasmid of approximately 23 kb in size. Carriage of this rare type resistant determinant is of clinical and epidemiological interest as non-pathogenic organism may act as vehicle for their dissemination in community and within hospital environment.
Background

Aspergillus flavus is the second leading cause of invasive and non-invasive aspergillosis, also common cause of fungal sinus and cutaneous infections. Since resistance to these drugs has been seen in patients, susceptibility testing can helpful in defining the activity spectrum of an antifungal and determining the appropriate drug for treatment.

Objectives

The aim of this study was to analyse the in vitro activities of amphotericin B, itraconazole, voriconazole, posaconazole, and caspofungine against clinical isolates of A. flavus recovered from clinical and environmental specimens in Iran.

Methods

199 isolates of A. flavus (171 clinical and 28 environmental) were included in the study. All isolates were identified by typical colony and microscopic characteristics, also characterized by molecular methods. MICs and MECs were determined according to recommendations stated in the Clinical and Laboratory Standards Institute (CLSI) M38-A2 document.

Conclusions

The range of MIC/MEC and MIC/MEC90 values were as follows: amphotericin B, 0.031-8 and 1 µg/ml; voriconazole, 0.031-8 and 1 µg/ml; itraconazole, 0.031-2 and 1 µg/ml; posaconazole, 0.008-0.5 and 0.341 µg/ml; and caspofungin, 0.008-0.25 and 0.063 µg/ml, respectively. Caspofungin was the most active drug, followed by posaconazole, itraconazole, Voriconazole, and amphotericin B. Caspofungin exhibited highly potent activity (100% of strains had MEC50 and MEC90 values of ≥0.5 µg/ml) against A. flavus strains. Among the azoles, posaconazole (MICs ≥1 µg/ml) was the most potent followed by itraconazole and voriconazole. We were not able to find significant differences in antifungal susceptibilities between strains recovered from clinical and environmental specimens (P >0.01).
HEAVY METAL TOLERANCE OF STAPHYLOCOCCUS AUREUS ISOLATES FROM PATIENTS TREATED WITH SILVER COATED WOUND DRESSINGS

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Background

Silver coated wound dressings are a new useful tool to fight bacterial infections. In vitro studies with heavy metal surfaces showed good results but only by direct contact with the patch and with a dry metal surface. But the efficacy of the coatings seems to be less effective over a longer application time.

Objectives

The aim of the study was to test heavy metal tolerance of Staphylococcus aureus wound isolates that had contact to silver coated wound dressing in comparison to isolates that had not.

Methods

120 Staphylococcus aureus patients isolates (MRSA and MSSA, with or without metal contact). Strains were analysed in a Bioscreen devise over a period of 72 hours, with copper concentrations up to 20 mM copper chloride. Additionally the strains were screened for genes thought to be responsible for heavy metal tolerance (copA, copB).

Conclusions

Some isolates of both groups showed high tolerance levels for copper chloride. Surprisingly MRSA (28% positive for copB) came up with an overall lower metal tolerance (around 8mM) in contrast to MSSA with 9mM. Prior contact to silver dressing did not show any difference in their metal tolerance. MRSA were positive in 28% for copB whereas MSSA were positive in 10%. Interestingly some of the high tolerance strains showed a prolonged lag-phase over days. Therapy with silver dressings does not favor strains with higher metal tolerance. In contrast to other studies the Austrian MRSA showed lower copper tolerance than MSSA.
CHARACTERIZATION OF LIPID COMPOSITION OF VANCOMYCIN-RESISTANT STAPHYLOCCUS EPIDERMIDIS 33 GISK

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Background

Numerous studies have revealed that the emergence of vancomycin-resistant staphylococcal phenotype is a result of multiple mutations that markedly change biological properties of these bacteria.

Objectives

The goal of these investigations was the analysis of adaptation changes of S. epidermidis 33 GISK (Moscow) lipid composition and S. epidermidis 33 GISK Van⁻ (its derivative strain) obtained via selection on media with increasing vancomycin concentrations.

Methods

Extraction of bacterial lipids was realized following the modified Bligh and Dyer method. Silica-gel plates Sorbfil (Russia) were used in thin-layer chromatography of lipid extracts in chloroform-methanol-water system (65:25:4, v/v). Identification of individual lipids was implemented via their functional groups’ staining with specific dyes. Analysis of fatty acid methyl esters was made using RTX-5MS column (Restek) on gas chromatograph 6890N with MS-detector 5973N (Agilent Technologies). Chromatography was carried out in autonomous temperature mode (180 to 280°C) at a rate of 5°/min. Peak substances were identified with NIST98 database software library.

Conclusions

Increase of bacterial resistance to vancomycin is accompanied by augmentation of cardiolipin and phosphatidylglycerol content in cell membrane lipid components, and the occurrence of lysyl phosphatidylglycerol and phospholipid lysoforms therein, but decline in glycolipid content.

According to data obtained the main point of lipid composition alteration in Van⁻ strain was the occurrence of unidentified phospholipid containing the alanine residue. Gas-chromatographic assay demonstrated that saturated and mono-unsaturated fatty acids C₁₆ and C₁₈:₁ were the predominant components of Van⁻-lipids, whereas fatty acids C₁₅ and C₁₇ were typical for the initial strain.

The work was supported by RFBR (14-04-00687_a).
Background

*E. coli* exhibits an increasing resistance to broad spectrum antimicrobial agents as well as the subsequent generations of these drugs. Previous genetic diversity studies indicated transmission of food-borne *E. coli* strains to human patients. Therefore it is important to study the genetic diversity and antibiotic-resistance patterns both in clinical and environmental *E. coli* to predict potential of transmission of organisms and genes antibiotic resistance.

Objectives

The aim of this study was to compare the genetic diversity and the antibiotic resistance patterns of uropathogenic *E. coli* (UPEC), to strains from freshwater, seawater and iguanas.

Methods

*E. coli* isolates were identified using API20E and ATCC 25922, Rep-(GTG)$_5$ PCR and BOX-PCR extragenic DNA fingerprinting, and FigTree™. The antibiotic resistance was assessed using the Kirby-Bauer disc diffusion against the eleven most commonly prescribed antibiotics in Grenada. Excel t-test and Statistica™ were utilized for comparison of patterns of drug resistance among four
Conclusions

Both DNA fingerprinting methods targeted non-protein coding or extragenic DNA and demonstrated enormous diversity within the population of the studied bacteria. UPEC expressed resistance patterns similar to the marine for 50%, while both to iguana and to fresh water isolates for 72.7% of drugs tested. There were only three drugs which showed significant difference between the UPEC and the freshwater isolates, namely ampicillin, amoxicillin/clavulanate and nitrofurantoin. We identified resistance to β-lactam antibiotics and lack of resistance to ciprofloxacin and gentamycin both in clinical strains of *E. coli*. The fractions of drug resistance strains and resistance patterns of the UPECs were similar to *E.coli* isolated from natural sources in Grenada.
Background
Due to escalating resistance to a range of antimicrobial agents, carbapenem resistant Acinetobacter baumannii (CRAB) has become a worldwide concern. Rapidly increasing prevalence of CRAB in many parts of the world in the past few years has questioned the reliability of the carbapenems.

Objectives
This study aimed to correctly identify A. baumannii species, antimicrobial susceptibility, and clonal relationships of CRAB clinical isolates.

Methods
Clinical isolates of A. baumannii collected from Clinical Bacteriology Laboratory between Sept 2013 and June 2014 were used in this study. Genospecies identification was verified by MALDI-TOF, and using polymerase chain reaction for OXA-51 & gyrB genes. MIC for meropenem, imipenem & doripenem was determined using agar dilution method. Antibiotic susceptibility was performed using Kirby-Bauer disc diffusion method. Multi-locus sequence typing (MLST) was performed to study clonal relationships.

Conclusions
Our results demonstrate clonal spread of CRAB in our hospitals. Further MLST studies on a nationwide basis would provide a better insight into the transmission and dissemination of CRAB clones.
IDENTIFICATION OF A HEAT STABLE BETA-LACTAMASE, WHICH IS PRODUCE BY BACILLUS CEREUS NBRC13494, INACTIVATES THE MEROPENEM

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Background
Antibiotics are used for the medical care, the domestic animals growth, and the control of health administration. Some antibiotics exposed in the environment are stable and keep the activities, which potentially causes the potential ecological and human health issue. The penem antibiotics, which are hardly to be degraded, are frequently used in Japan.

Objectives
In order to reduce the antibiotics pollution, we aimed to isolate soil bacteria that produce meropenem digestion enzymes.

Methods
Genome sequence analysis reveals that the some Bacillus species potentially produce antibiotic degradation enzymes. We screened Bacillus strains that can digest the meropenem from Biological Resource center in Japan and the soils.

Conclusions
We found B. cereus strain NBRC13494 and three natural isolates, SUBC1002, SUBC1008, and SUBC1010. The MICs of meropenem were 6.7, 2.5, 3.3, 3.4 mg/l, respectively. B. cereus strain NBRC13494 possessed the highest beta-lactamase activity among them. We found that the supernatant of B. cereus NBRC13494 after the heat treatment at 80 °C was also effective to degrade the meropenem. We resolved the proteins included in the supernatant of B. cereus NBRC13494 by SDS-PAGE and found major two bands with the molecular mass of 28-kDa and 33-kDa bands. We determined their amino acid sequences by LC-MS/MS. The 28-kDa protein corresponded to the B. cereus 5/B/6 beta-lactamase II. The 33-kDa protein corresponded to the B. cereus 5/B beta-lactamase I. These beta-lactamases belong to the different protein family and highly conserved among B. cereus group. These results suggested that some beta-lactamases produced by B. cereus strains potentially digested the meropenem.
IDENTIFICATION AND PREVALENCE OF ANTIBIOTIC RESISTANCE GENES IN FRESHWATER URBAN WATERSHEDS

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Background

Antibiotic resistance genes (ARGs) of environmental origin have been identified as a potential source of the rise in antibiotic resistance in communities. To understand the risk of ARGs in source waters, identification and quantification of ARGs in watersheds is necessary. Here, two urban watersheds with different land-use profiles were studied: a river situated in an urbanized environment and a lake situated in a reserved environment.

Objectives

To identify and quantify ARGs using GeoChip microarray and quantitative PCR (qPCR).

Methods

To screen for ARGs, GeoChip version 5, which target 17 families of ARGs, was used to screen DNA extracted from three sampling sites at each watershed. Primers for qPCR were designed based on ARGs yielding the highest GeoChip intensities.

Conclusions

Based on GeoChip analysis, ABC transporters, multidrug and toxic efflux, multidrug resistance efflux pumps, multi-facilitator superfamily (MFS), small multidrug resistance (SMR), β-lactamases (A-D), tetracycline resistance (tet), vancomycin, virginiamycin resistance gene B (vgb), fosfomycin resistance (fosA, fosB, fosX) and quinolone resistance (qnrC) were detected in both watersheds. There was significantly higher intensity of β-lactamase A (p<0.05), MFS (p<0.01), β-lactamase C (p<0.01), SMR (p<0.01), and vgb genes (p<0.01) in the urban environment than in the reserve; contrarily, fosA had a higher intensity in the reserve.

Quantification of tet and MFS revealed presence of these genes in low relative abundance (<2.5%) to 16S rRNA genes in both watersheds. Taken together this data
suggests that there are more ARGs in the urban than the reserve watershed and that only a sub-population of the bacterial community harbours these genes.
Antibiotic resistance and environment

CARRIAGE TRANSMISSION AND SELECTION OF ANTIBIOTIC-RESISTANT ESCHERICHIA COLI IN AN EXPERIMENTAL BIRD MODEL


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Background

Very low concentrations of antibiotics, more than a hundred times lower than the minimal inhibitory concentration (sub MIC), can select for antibiotic resistant bacteria in vitro.

Objectives

To investigate if sub MIC concentrations of antibiotics in the environment can select for antibiotic resistant bacteria present in the intestine of mallards and to study if and how rapidly different ESBL E. coli strains transmit within a flock of mallards.

Methods

In the selection study birds were infected with an equal amount of two isogenic ESBL producing E. coli strains where one is resistant to ciprofloxacin. Mallards were exposed to different concentrations of ciprofloxacin in the drinking water. Faecal samples were analysed and the ratios of ciprofloxacin resistant over ciprofloxacin susceptible bacteria were calculated.

In the carriage and transmission study four different ESBL E. coli strains with different resistance patterns were used to inoculate a group of mallards. Faeces samples from each bird were collected up to 29 days post infection and colonies were patched onto selective plates to follow each of the inoculated strains.

Conclusions

Low levels of antibiotics (<20 ng/ml) in the water source of birds enrich for antibiotic resistant bacteria. Resistant bacteria can be carried in a flock of birds for at least a month without any antibiotic selective pressure and antibiotic resistant bacteria rapidly spread in the flock of birds.
EXTRACELLULAR VESICLES AND RESISTANCE OF MYCOPLASMAS TO FLUOROQUINOLONES: RESISTOME OF ACHOLEPLASMA LAIDLAWII

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Background
Control of mycoplasma infections presents a serious problem which solution is connected with elucidating molecular-genetic mechanisms of mycoplasma adaptation to environments. Sequencing of bacterial genomes provides an opportunity for application of postgenomic technologies to study the above-mentioned processes. In our studies it was shown for the first time that adaptation to environments of Acholeplasma laidlawii – the “ubiquitous” mycoplasma which is the major contaminant of cell cultures and the causative agent of phytomycoplasmoses, was associated with not only specific modulation of transcriptome and proteome profiles of the bacterium but secretion of extracellular vesicles.

Objectives
Comparative analysis of genome and proteome profiles and vesicular content of Acholeplasma laidlawii strains with different susceptibility to ciprofloxacin was the object of our work.

Methods
The TEM, SEM, LC-ESI-MS/MS and DNA sequencing were used in our study.

Conclusions
As a result of our studies it was found that A. laidlawii strains with different susceptibility to ciprofloxacin differed significantly on complete nucleotide sequences of genomes, vesiculation level, cellular and vesicular proteomes. Furthermore, it was revealed that the extracellular vesicles of A. laidlawii mediated export of ciprofloxacin as well as mutant nucleotide sequences of the genes that encoded the antibiotic-targeted proteins. The obtained data provided a necessity for correction of current notion about the mechanisms that facilitate the rapid development of resistance to antibacterial agents in mycoplasmas and approaches to control of mycoplasma infections.

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Background
Hospital and municipal effluents receive high doses of antibiotic resistant bacteria and their genes (ARB&ARG), some of which can persist in the environment. Little is known about the factors influencing the fate of ARG when they are introduced in complex microbial communities.

Objectives
This study aimed at assessing the fate of some indicator genes hosted by different bacterial species when inoculated in wastewater.

Methods
Assays were performed in microcosms of hospital effluent, raw wastewater, treated wastewater and sterile ultrapure water. Replicas of these microcosms were inoculated with Acinetobacter johnsonii (intI1), Escherichia coli (blaTEM, intI1) and Enterococcus faecium (vanA) or with Acinetobacter johnsonii (intI1), Klebsiella pneumoniae (blaTEM, intI1) and Enterococcus faecalis (vanA). Inoculated and non-inoculated microcosms incubated for 15 days at 18 or 30 ºC, the genes were quantified by qPCR and bacterial community composition was assessed based on 16S rRNA gene-denaturing gradient gel electrophoresis.

Conclusions
In general, the bacterial community richness, diversity and evenness indices did not vary over the incubation period and similar results were observed at 18ºC or 30 ºC. Both vanA and blaTEM were observed to decay over time, with higher reduction than the genes 16S rRNA and intI1. The gene vanA decreased sharply after the incubation period irrespective of the gene host or type of water used in the microcosm. In contrast, the decay of the gene blaTEM varied with the host and type of water. Comparatively, K.pneumoniae presented a higher fitness in water than E.coli, suggesting the importance of the host in the spread of some ARG.
Background
Aquatic environments are important habitats for bacteria and water is a privileged mode of dissemination of microorganisms. Freshwater is inhabited by different bacterial populations, mostly strictly environmental, in pristine environments, and also of human/animal origin, in human-impacted areas. The rivers that flow in urban areas are often inhabited by both environmental and animal/human derived microorganisms.

Objectives
The study of the diversity and antibiotic resistance patterns of bacteria colonizing freshwater habitats brings interesting insights into the role of the natural and human-derived microbiota in the evolution of antibiotic resistance. This study explored the taxonomic diversity and antibiotic resistance patterns of bacteria isolated from freshwater.

Methods
Water and/or sediment samples were collected from rivers Ferreira (43 km) and Douro (897 km), in Northern Portugal. Heterotrophic bacteria were enumerated and isolated on plate count agar (PCA) or on PCA supplemented with antibiotics (PCA+A). The isolates (n=173) were characterized for their antibiotic resistance phenotypes and multidrug resistant (MDR) bacteria (to antibiotics of ≥4 classes) were identified based on 16S rRNA gene sequence analysis.

Conclusions
In both rivers, PCA counts ranged $10^5$ CFU/g of the sediments being about 1-3 log less on PCA+A (ciprofloxacin, meropenem, gentamycin). In both Ferreira and Douro, the most prevalent antibiotic resistance phenotypes were for beta-lactams and sulfonamides and the least prevalent was for tetracycline. Ceftazidime, meropenem and ciprofloxacin resistance was more prevalent in Douro than in Ferreira isolates. About 40% of the isolates from PCA+A were MDR and the most common genera were Pseudomonas, Chryseobacterium and Stenotrophomonas, in Ferreira and Chitinophaga in Douro.
Background

The bacterial SOS response is a global stress response involved in DNA-damage repair. In *Escherichia coli* it is induced after DNA damage, for example after treatment with antibiotics that target DNA. The SOS response is also involved in genome plasticity and in the acquisition of resistance to antibiotics. Unexpectedly, the SOS response is induced in *Vibrio cholerae* by exposure to sub-inhibitory concentrations (sub-MIC) of different families of antibiotics that do not target DNA, such as aminoglycosides, chloramphenicol and tetracycline. The use of a genetic screen led to the isolation of mutants in which induction of the SOS response by sub-MICs of aminoglycosides is lost. One of these mutants is inactivated for the vc1636 gene, which encodes a putative DNA/RNA helicase.

Objectives

The purpose of this project is to characterize the protein encoded by the *V. cholerae* vc1636 gene and to understand the induction of the SOS response by antibiotics that do not target DNA in *V. cholerae*.

Methods

Towards this goal we use several bacterial genetics assays.

Conclusions

The first results suggest that VC1636 plays an important role in DNA-damage repair.
ANTIMICROBIAL AND VIRULENCE FACTORS OF V. PARAHAEOMOlyticus ISOLATED FROM THE FINAL EFFLUENTS OF WASTEwATER TREATMENT PLANTS

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Background
Enteric pathogens, including Vibrio species, increasingly survive wastewater treatment processes, thus constituting threat to humans as final effluents are discharged into receiving water bodies used primarily as source of domestic or recreational waters. V. cholerae has been reported from various water sources however, other pathogenic species have been under reported in wastewater hence the focus is on V. parahaemolyticus for this study.

Objectives
The objectives of the study were to: Isolate and confirm presumptive Vibrio species while determining their incidence; Confirm V. parahaemolyticus using PCR; Determine the antibiogram characteristics of V. parahaemolytics and determine the virulence factors specifically the tlh and trh gene of V. parahaemolyticus.

Methods
Wastewater final effluents were collected once, monthly, for a period of twelve months, membrane filtered and analysed in accordance with standard technique. Presumptive Vibrio isolates were confirmed via 16SrRNA gene and V. parahaemolyticus identified with toxR gene. Antimicrobial susceptibility test was done using 10 different antibiotics and virulence factors namely the thermostable direct haemolysin related haemolysins (trh gene) and thermolabile haemolysin (tlh gene) were assessed using PCR

Conclusions
Incidence of Vibrio species varied with months and treatment plants. Resistance was shown against certain antibiotics while in other antibiotics resistance varied. The incidence of trh genes was nil, while the tlh gene was 5.7 %. Although, the incidence of virulence determinants in V. parahaemolyticus were few, their presence portends a public health risk to the receiving water bodies used by humans and animals especially with the increasing resistance of the pathogens to antibiotics.
PHENOTYPIC AND MOLECULAR CHARACTERIZATION OF KLEBSIELLA SP FROM URINE SAMPLES OF IN- AND OUT-PATIENTS IN SOME HEALTH INSTITUTIONS IN NIGERIA

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Background

Klebsiella species are important pathogens in nosocomial infections, most especially the multiple antibiotic resistant strains.

Objectives

The aim of this study was to isolate and identify Klebsiella sp from urine samples of in- and out-patients in some health institutions in Ile-Ife, Nigeria; determine the resistance pattern; detect plasmid DNA, resistance and virulence genes in multiple antibiotic resistant Klebsiella isolates.

Methods

Isolation of klebsiella sp was carried out on MacConkey agar plate at 37°C. Identification was done using standard techniques. Antibiotic sensitivity was determined using Kirby-Bauer’s disc diffusion method. Plasmid DNA was extracted by alkaline lysis, resistance (blaCTX, tetM) and virulence (cps and fimH) genes were detected by PCR.

Conclusions

A total of 32 (7.02%) Klebsiella sp were identified, out of which 29 (90.63%) were multiple antibiotic resistant (MAR). Resistance to antibiotics varied greatly. The isolates showed high resistance to augmentin (93.75%) and tetracycline (71.88%). There was diversity in the MAR pattern with 21 different antibiotypes which ranged from two to ten antibiotics. Thirteen out of the 18 randomly selected MAR isolates haboured plasmids ranging from 900bp to 9416bp. Three (16.67%) of the isolates had blaCTX (543 bp) and cps (418 bp) genes while one (5.56%) harboured fimH (1250 bp) gene. However, tetM (974 bp) gene was not detected in any of the isolates. The incidence of MAR Klebsiella sp in the study is high and calls for great concern in health care delivery with great economic and health consequences.
Antibiotic resistance and environment

AEROMONAS SPP. – B-LACTAMASES RESISTANCE IN WASTEWATER TREATMENT PLANT STRAINS

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Background

Bacteria of the Aeromonas genus are fish and human pathogens commonly occurring in various aquatic ecosystems (Janda and Abott, 2010). Literature data show a wide spread of drug resistance genes among these microorganisms with a high proportion of chromosomal and plasmid-mediated genes (Piotrowska and Popowska, 2014).

Objectives

The aim of this project was determination of antibiotic resistance genes in the β-lactam unsusceptible Aeromonas strains isolated from wastewater with different levels of purification. In addition, in drug-resistant strains, their plasmid profile and the conjugation potential was defined.

Methods

The double-disk synergy test was performed to determine the mechanisms of β-lactamases resistance (AmpC, ESBL, carbapenemases). In addition, disk-diffusion were performed to determine the bacterial antibiogram. Identification of resistance genes was performed by PCR (Multiplex) and confirmed by sequencing. Plasmids were isolated by alkaline lysis and commercial kits. Diversity of plasmids were evaluated by plasmid restriction profiles by comparison of their patterns obtained after digestion with different restriction enzymes. The conjugative transfer ability and host range of selected plasmids were performed according to standard methods.

Conclusions

Aeromonas strains existing in wastewater are β-lactam resistant and possess a variety of β-lactamases. We found and confirmed the genes encoding β-lactamase from variety of families e.g. OXA, TEM, SHV, GES, VEB, FOX and characteristic to Aeromonas spp. chromosomal metallo-β-lactamase CphA. Some of the resistant strains possess plasmids which were characterized by a different profiles.
IMPACT OF URBAN CONTAMINATION OF LA PAZ RIVER ON THE ABUNDANCE OF FECAL COLIFORMS, AND ANTIBIOTIC RESISTANT GRAM-NEGATIVE BACTERIA IN RIVER WATER, SOIL AND VEGETABLES

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Background
Worldwide, contamination of fresh water resources affects food security and ecosystem sustainability. This situation is aggravated in developing countries and markedly in dry regions such as the Bolivian Highlands. In this region along with a shortage of water, urban wastewater is directly discharged into fresh water bodies contributing to aquatic ecosystem degradation, as observed in La Paz River, which receives urban discharges that further contaminate agricultural lowlands, where vegetables and soil are directly irrigated by river water

Objectives
To evaluate the microbial quality of river surface water at unimpacted and impacted sites, and tested for antibiotic resistance of gram-negative bacteria isolated from river water, soil and vegetable samples.

Methods
Culture dependent methods.

Conclusions
High levels of fecal coliform density were found along impacted sites and green-leafy vegetable samples, exceeding at least in two and one order of magnitude respectively, the allowable concentration. Antibiotic resistance among gram- negative bacterial isolates was common at all sites. A spatial distribution of antibiotic resistance was found, where resistance frequency to not commonly used antibiotics was higher at unimpacted in contrast to impacted sites and vice versa, this distribution was associated with the density of fecal coliforms. Multiple Antibiotic Resistance (MAR) index values ranged from 0.176 to 0.310, being higher in soil samples. Overall, this study suggest that surface water, soil and vegetables can be reservoirs of antibiotic resistant gram- negative bacteria, highlighting health risk associated with vegetables production and consumption in the region.
Background

Resistance to quinolones has been little studied in lactobacilli and the breakpoint value established to discriminate resistant from sensitive strains, the same for all Lactobacillus species, has not been updated since 2003 (1).

High level of resistance is, in most cases, due to mutations in the quinolone resistance determining regions (QRDR) of the genes encoding DNA gyrase (gyrA) or topoisomerase IV (parC). Reduced sensitivity to quinolones is conferred by genes, among which the most widespread are the qnr genes for pentapeptide repeat family proteins that protect gyrase from quinolone inhibition, encoded by plasmids or integrons in Gram-negative bacteria and chromosomally in some Gram-positive bacteria (2).

Objectives

This study was aimed at understanding if the species of probiotic interest Lactobacillus casei/paracasei/rhamnosus may have a role in the spread of quinolone resistance.

Methods

Strains resistant to ciprofloxacin were selected among 184 L. casei/paracasei/rhamnosus bacteria by using the breakpoint concentration for lactobacilli (4 mg/l) in LSM medium. These were tested for the presence of mutations in the gyrA and parC QRDRs and of genes conferring quinolone resistance.

Conclusions

Among the 31 strains resistant to the breakpoint value for ciprofloxacin only one L. rhamnosus strain presented a mutation in the gyrA QRDR, not corresponding to those commonly associated to resistance. Results showed that quinolone resistance is intrinsic in some strains of L. casei/paracasei/rhamnosus.

SCREENING BACTERIA ISOLATED FROM AQUATIC ENVIRONMENTS FOR THE POTENTIAL TO DEGRADE SULFAMETOXAZOLE
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Background
In contrast to beta-lactam antibiotics, the biodegradation of sulfonamides seems to be uncommon in nature, in spite of the extensive use and environmental contamination associated with these drugs.

Objectives
The aim of this study was to investigate the potential of sulfonamide resistant bacteria isolated from wastewater, surface water, bottled mineral water, and different sites within a drinking water treatment plant, including distribution system and tap water, to degrade sulfamethoxazole (SMX).

Methods
From a total of 133 isolates, were selected 51 representative strains affiliated to 19 genera - Pseudomonas, Sphingomonas, Blastomonas, Acinetobacter, Aeromonas, Variovorax, Microbacterium, Duganella, Bosea, Afipia, Brevundimonas, Acidovorax, Rhizobium, Klebsiella, Citrobacter, Raoultella, Enterobacter, Kluyvera, and Stenotrophomonas to test the ability to degrade SMX. The biodegradation experiments were carried out in mineral and complex culture media supplemented with 50 mg/L SMX.

Conclusions
Five isolates belonging to the genus Pseudomonas and one to the genus Stenotrophomonas were able to degrade SMX in complex culture medium. Pseudomonas mandelli isolated from bottled mineral water and Stenotrophomonas maltophilia isolated from river water were the most efficient SMX-degrading bacteria, removing 67% and 38% of the initial SMX content after 15 days of incubation, respectively.

Based on the potential application of these microorganisms or catalysts thereof, further research on the SMX degradation kinetics and metabolites are under study and will be discussed with what is described for other Proteobacteria and Actinobacteria.
Background
Extended-spectrum beta-lactamase (ESBL)-producing Enterobacteriaceae is an increasing public health problem. Extensive collection of these strains is needed to analyse the issue more thoroughly. Human Microbiota Biobank (HUMB) at University of Tartu (www.eemb.ut.ee/eng/, WFCC-MIRCEN no. 977) is a scientific collection that contains beneficial and opportunistic microbial strains of human origin.

Objectives
To form the subcollection of ESBL positive Enterobacteriaceae strains in HUMB for further pheno- and genotypic characterisation.

Methods
More than 1200 strains of ESBL positive Escherichia coli and Klebsiella pneumoniae were collected from 20 hospitals in Estonia (n=627), Latvia (n=182), Lithuania (n=134) and St. Petersburg region in Russia (n=322) in 2011-2012. The strains were identified by MALDI-TOF MS and deposited at the HUMB collection (stored at -80°C in multiple parallels in multiple freezers). Bacterial isolates were compared according to clinical material, geographical origin, phylogenetic group, antibiotic susceptibility and virulence factor genes.

First results of the subcollection of ESBL producing strains have been published (Pavelkovich 2014; Lillo 2014). The regional differences of virulence factor genes and carbapenemase resistance remained statistically significant after taking into account the phylogenetic distribution in the countries. MALDI-TOF MS-based assay appeared to be suitable, quick and cost-effective method for the initial confirmation of carbapenemase production.

Conclusions
Subcollection of ESBL producing strains stored in HUMB have significant scientific value that has to be further studied for epidemiological purposes and for creating evidence-based suggestions for clinicians.
Wastewater treatment plants are one source of microbial contamination. In the light of recent studies there are reports of increased levels of multi-resistant bacteria and antibiotic resistance genes (ARGs) in wastewater treatment plants (WWTPs) and downstream habitats. Conventional WWTPs represent a reservoir for the dissemination of ARGs in the bacterial communities. Horizontal gene transfer (HGT) of antibiotic resistance genes contributes to increased abundances of hygienically relevant, antibiotic resistant bacteria in aquatic systems. Advanced oxidation processes (AOPs) as additional wastewater treatment techniques are investigated to decrease the bacterial load and the dissemination of antibiotic resistances in adjacent downstream aquatic habitats.

To investigate the efficiency of ozonation to reduce the bacterial load, the project includes detection and quantification of facultative pathogens and medically relevant antibiotic resistance genes in microbial communities by qPCR analysis. We investigate the impact of ozonation by molecular quantification of the methicillin resistance gene mecA from staphylococci, the ampicillin resistance gene ampC from Enterobacteriaceae, the transposon mediated vancomycin resistance gene vanA from enterococci, the imipenem resistance gene blaVIM from Pseudomonas aeruginosa, and the erythromycin resistance gene ermB from streptococci.

As expected, most of the hygienically relevant bacteria were reduced during ozonation, but it became obvious that bacteria carrying antibiotic resistance genes showed a considerable robustness and fitness against ozonation. This indicates a selection-like process favoring already antibiotic resistant bacteria during oxidative wastewater treatment. Further investigations regarding drug-dependent predispositions and different bacterial stress response mechanisms are currently analyzed to characterize the bacterial adaptation.
RESISTANCE TO AZTREONAM IN ENVIRONMENTAL ISOLATES OF PSEUDOMONAS AERUGINOSA RESULTS FROM HYPEREXPRESSION OF MEXAB-OPRM INDUCED BY MUTATION IN NALC GENE.

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Background

Bacteria belonging to the genus \textit{Pseudomonas} have a great metabolic versatility and acquire resistance to a wide variety of antimicrobials. Resistance to the monobactam aztreonam occurs primarily by changing in the efflux pump MexAB-Opm system.

Three transcriptional regulators modulate the expression of this operon: MexR, from MarR family, and NalD and NalC, TetR family members. NalC is an anti-repressor that allosterically inhibits the MexR dimer formation and, then, MexR is released from the mexAB-oprm promoter, leading to an upregulation of this operon.

Objectives

This study aimed to determine the resistance profile to aztreonam in 50 \textit{P. aeruginosa} isolates and to determine if mutations in the regulatory genes affected the expression of MexAB-OprM efflux pump.

Methods

The soil sample was collected and processed according to the method describe by Mukherjee et al. (2010), and the resistance profile was determined by the Minimum Inhibitory Concentration (MIC) method. The transcriptional regulators \textit{mexR}, \textit{nalC} and \textit{nalD} were amplified by PCR, and the expression of MexAB-OprM was measured by quantitative real-time reverse transcriptase-PCR (qRT-PCR).

Conclusions

All isolates analyzed had some amino acid substitution in NalC, only one isolate showed a substitution at MexR and no single mutation was detected into NalD protein. Just the isolate that presented mutation at position 186 had MIC more than 250 µg mL\textsuperscript{-1} and showed a greater increase in the level of \textit{mexA} expression, about 92.6 fold more than that found in the wild-type PAO1 strain.
Background

P. aeruginosa is an opportunistic microorganism widely spread in different terrestrial and aquatic environments, but it is also known multiresistant pathogen in hospital setting.

Objectives

We aimed to characterize the population structure of P. aeruginosa isolated from human, animal and environmental samples; to identify the potential transfer routes between various environments and to look for the associations between antibiotic resistance and potential resistance genes.

Methods

Ninety strains were collected between 2004 and 2014: 41 clinical isolates from patients with infection; 4 colonising strains from nasal and faecal samples of 287 healthy volunteers; 29 isolates from sick animals and 16 from environmental samples.

Study has been approved by Research Ethics Committee of the University of Tartu.

All samples underwent whole genome sequencing. Draft genomes were assembled followed by in silico identification of markers for multilocus sequence typing (MLST).
analysis. Genetic relatedness was presented as minimum spanning tree.

Minimum spanning tree of *P. aeruginosa* from various sources. Each circle corresponds to a unique sequence type from MLST database (pubmlst.org/paeruginosa). Circle size is proportional to sequence type frequency.
Coding regions for beta-lactamases were predicted from draft genomes using custom HMM profiles for Ambler class A, D, C and B beta-lactamases.

MICs to 10 antipseudomonal agents (ceftazidime, cefepime, meropenem, imipenem, piperacillin/tazobactam, amikacin, gentamycin, tobramycin, ciprofloxacin, colistin) were detected using epsilometer test. EUCAST breakpoints and multiresistance definition were used for interpretation.

Phenotypic resistance was compared to the betalactamase genes predicted from the genomic sequences.

Conclusions

Our data indicates that *P. aeruginosa* strains are transferred between different hosts and environments. The high variety of phenotypic resistance and resistance genes in bacteria of the same sequence type suggests rapid horizontal transfer of resistance.
Background

Developing dosage guidelines for antibiotics with mitigation of adverse risks is a goal of the rational use of medications. The use of checkerboard tests enables evaluation “in vitro” of the association of antibiotics in different concentrations and optimization of gentamicin and ampicillin concentrations in terms of therapeutic effectiveness, contributing to more effective therapeutic treatment guidelines.

Objectives

The objective of the study is to evaluate the synergism of action of gentamicin associated to ampicillin through “in vitro” studies in enterococci strains.

Methods

A total of 50 Enterococcus sp strains were used and the susceptibility profile to ampicillin and gentamicin evaluated separately and in combination. For each enterococci strain, a total of 36 tests were performed in 1 mL tubes with different concentrations of ampicillin and gentamicin. Figure 1.

Conclusions

A total of 1,800 tests were conducted with different concentrations of antibiotics. After 24 hours and 48 hours of incubation, the turbidity, representing the action of antimicrobial on the enterococci strains, was observed.

The synergism of action was observed in 10% of samples, with 54% indifference and 32% revealing therapeutic treatment failure. In other words, mere association of the antibiotics does not result in successful therapeutic treatment. Evaluation “in vitro” of different concentrations of antibiotics can help guide proper treatment of infections through appropriate doses that permit synergy of action and therapeutic success.

Figure 1: Checkerboard – Evaluation of the synergism of action between different concentrations of ampicillin and gentamicin on enterococci strains, $1.5 \times 10^8$ UFC/mL.
The initial dose of antibiotics was 0.4mg/mL, corresponding to 10μg discs with progressive dilution of the antibiotics.

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FEMS-2912
Antibiotic resistance and environment

CORRELATING METAL CONTENT IN (EXTREMELY) POLLUTED SOIL WITH BACTERIAL RESISTANCE TO ANTIBIOTICS AND METALS
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Background

Microbes require some (trace) metals for growth, while high metal concentrations can be toxic. Metal-resistant microbes have evolved in metal-polluted environments, some of which may also have (co-)developed antibiotics resistance.

Objectives
This study aims to determine the environmental risk of (spread of) antibiotic resistance to humans from historically metal-contaminated urban areas.

Methods
This study analysed the spatial variation of the concentrations of 16 metals in 375 topsoil samples from the Swansea/Neath/Port Talbot urban area, which contain some of the highest metal-polluted soils of the UK. Pairwise correlations showed mostly significant positive associations between 16 of the studied elements, e.g. copper and arsenic. Factor analysis showed that 68% of the observed variation in concentrations could be explained by four principal components. The archived soils were then subsampled over a range of metal concentrations, yielding 65 subsamples for further biomolecular analysis.

Conclusions
Culture-dependent studies on dilute nutrient rich and humic acid agar media yielded many isolates with medium to high tolerance to several metals and antibiotics. Culture-independent studies are performed to determine the bacterial diversity and abundance of resistance genes over the wide range of observed (mixed) metal concentrations.
Background

The human gut harbors a densely populated microbial ecosystem, termed the gut microbiota. The gut microbiota can serve as a reservoir for antibiotic resistance genes ('the resistome') and drug-resistant opportunistic pathogens.

Objectives

In this study, we determined the dynamics of the microbiota and the resistome in thirteen patients that were hospitalized in Intensive Care Units. Because gut-associated bacteria and their resistance genes may be shed into the environment through sewage, we also assayed the levels of antibiotic resistance genes in hospital sewage and in subsequent stages of wastewater treatment.

Methods

16S rRNA profiling, functional metagenomics, metagenomic sequencing and nanolitre-scale quantitative PCRs were used in this study.

Conclusions

The gut microbiota of hospitalized patients contained genes that are predicted to confer resistance to fourteen classes of antimicrobials. Levels of antibiotic resistance genes increased during ICU hospitalization and were associated with changing patterns of the composition of the microbiota. Sequencing of vector inserts in drug-resistant fosmid clones indicated that resistance genes were mainly carried by anaerobic gut commensals (Clostridium and Bacteroides). Resistance genes were associated with mobile genetic elements, which could facilitate horizontal gene transfer of resistance genes from commensal bacteria to opportunistic pathogens. Antibiotic resistance genes were abundant in hospital sewage but the influence of hospital sewage on the abundance of resistance genes in the wastewater treatment
plant was limited. Levels of resistance genes and human-associated bacteria further decreased during wastewater treatment, limiting the dispersal of antibiotic resistance genes from hospitals into the environment.
Background

Molecular epidemiology-based studies using methods such as multilocus sequence typing (MLST) are useful to evaluate the dissemination of antibiotic resistant bacteria and their genes (ARB&ARG) in the environment. If the ARB bacteria tracked are typical human commensals like *Escherichia coli*, it is also possible to infer about the humans-environment relationship.

Objectives

This study aimed at assessing the genetic relatedness of quinolone unsusceptible *E. coli* isolated from different types of water, gulls, birds of prey and hospitalized patients, in order to infer about possible modes of transmission of ARB in the environment.

Methods

A group of 143 quinolone unsusceptible *E. coli* from clinical, wild birds and water isolates was compared based on MLST and resistance phenotype. Water isolates were further analyzed regarding the presence of chromosomal mutations in the genes *gyrA* and *parC*; of plasmid-mediated quinolone resistance (PMQR, *qnrA, B, C, D, S, VC; qepA; oqxAB; aac(6’)-ib-cr* and beta-lactam resistance (*bla*_{OXA-A,B,TEM,CTX,SHV,CMY}))-encoding genes; and of Inc group plasmid replicons. Water isolates carrying PMQR-encoding genes were tested for their ability to transfer resistance by conjugation.

Conclusions

Isolates from different sources clustered together, suggesting the ability to persist and spread across different environments. The most common sequence types were ST131 (n=26) and ST10 (n=9). 77% of the isolates were resistant to three or more classes of antibiotics. Eight water isolates harbored PMQR-encoding genes, combined with *bla*_{TEM} (n=6) and *bla*_{CTX-M-15} (n=2), from which seven could transfer
resistance through conjugation. These results confirm the role of multidrug-resistant \textit{E.coli} in the dissemination of ARB\&ARG across different environments.
Background

Over the last years, members of the genus *Aeromonas* have been recognized as one of the principal reservoirs of antibiotic resistance in aquatic environments. While several studies have documented the levels of antibiotic resistance associated with these bacteria in urban waterways, the contribution of hospital effluents remains unclear.

Objectives

The objective of this study was to elucidate the role of this genus in the dissemination of antibiotic resistance from hospital effluent into the environment.

Methods

A set of quinolone unsusceptible *Aeromonas* was isolated from untreated hospital effluent and from the inflow and treated effluent of the receiving urban wastewater treatment plant (UWTP). Isolates were identified and genotyped based on the 16S rRNA, *cpn60* and *gyrB* gene sequences. Further characterization involved determination of the antibiotic resistance phenotypes and screening for antibiotic resistance and mobile genetic elements (*qnrA,B,C,D,S,VC; qepA; oqxAB; aac(6')-ib-cr; blaOXA; incU*).

Conclusions

The most prevalent taxonomic groups were *A. hydrophila* and *A. caviae*, found in all types of water, while *A. salmonicida* was found exclusively in the hospital effluent and raw UWTP inflow, and *A. allosaccharophyla* and *A. veronii* in the UWTP. The genetic determinants *qnrS2+incU* were a signature of the UWTP and almost absent in the hospital effluent. The genes *aac(6')-ib-cr* and *blaOXA* were distributed in isolates from all sources. The dynamics of resistance genes and mobile genetic elements was evidenced by the fact that closely related isolates harbored different combinations of those genetic elements. This observation highlights the role of *Aeromonas* as important carriers of antibiotic resistance genes in aquatic environments.
CULTIVABLE GRAM-NEGATIVE BACTERIA IN DRINKING WATER: UBIQUITY, PERSISTENCE AND ANTIMICROBIAL RESISTANCE

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Background
To achieve required standards of quality and safety, in most world regions, drinking water must undergo purification and disinfection processes prior to its distribution. However, tap water is colonized by a myriad of bacteria comprising those that survive disinfection or that enter at any stage of the distribution network. Despite the multiple factors affecting water microbiota, some bacterial groups are known to prevail in drinking water\textsuperscript{1}. Good examples are given by genera of Alpha-, Beta- and Gamma-Proteobacteria, some of which display high levels of antibiotic resistance.

Objectives
The objective of this study was to characterize the diversity of cultivable Gram-negative bacteria isolated from tap water, collected over nine months. The aim was to assess which were the most prevalent, ubiquitous and persistent groups in drinking water and to evaluate their antimicrobial resistance potential.

Methods
More than 3000 isolates recovered from the water source, the water treatment and distribution system, and from 12 household taps, were screened for Gram-negative staining (51%) and absence of repetitions (same plate and identical genotype). From this selection resulted approximately 1100 isolates that were, most of which, identified by 16S rRNA or rpoB gene sequence analysis. The most representative groups occurring in tap water were characterized for their antimicrobial resistance phenotypes.

Conclusions
Among most ubiquitous groups were the genera Blastomonas, Brevundimonas, Methylobacterium, Sphingobium, Sphingomonas, Acidovorax, Ralstonia, Acinetobacter and Pseudomonas. Some of these groups comprise opportunistic pathogens and present profiles of multidrug resistance, making their presence in treated drinking water an issue of concern.

Reference:\textsuperscript{1}Vaz-Moreira et al., FEMS Microbiol Rev 38(4), 761-778.
FEMS-1377
Antibiotic resistance and environment

MULTIDRUG RESISTANT GRAM-NEGATIVE BACTERIA FROM HOSPITAL EFFLUENT

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Background
Hospital effluents are frequently discharged untreated in municipal collectors. However, this type of wastewater deserves a special attention given the high prevalence, and potential emerging character, of antibiotic resistance found in clinical environments.

Objectives
This study aimed at identifying quinolone resistant Gram-negative bacteria with the highest multidrug resistance index (MDR) thriving in hospital effluent or in the receiving municipal wastewater treatment plant. It was aimed to assess if these bacteria may represent critical vectors for the spread of antibiotic resistance from the hospital to the environment.

Methods
The criteria for inclusion in this study was to be Gram-negative, other than Escherichia coli and present reduced susceptibility to quinolones, a MDR phenotype (to ≥6 classes of antibiotics) or of resistance to meropenem or ceftazidime (n=25). These potential vectors of resistance were identified and characterized for antibiotic resistance phenotype, plasmid mediated quinolone resistance (PMQR) genes, presence of integron gene cassettes and conjugative capacity.

Conclusions
MDR isolates comprised bacteria of the genera Aeromonas, Acinetobacter, Citrobacter, Enterobacter, Klebsiella, Pseudomonas, Chryseobacterium and Myroides. About half of these isolates had at least one PMQR gene (aac(6')-Ib-cr, qnrB, qnrS or oqxAB) and a class 1 integron gene cassette encoding aminoglycoside, sulfonamide or carbapeneme resistance. Some isolates, mainly Enterobacteriaceae, could transfer through conjugation some resistance traits. These results showed that ubiquitous bacteria, other than the groups whose monitoring in water quality controls is recommended, are important vectors of antibiotic resistance. This information is useful to support management options aiming the control of this public health problem.
Antibiotic resistance and environment

COUNTERACTING ANTIBIOTIC RESISTANCE IN AGRICULTURE: BIOTECHNOLOGICAL BARRIERS MIGHT DO THE TRICK

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Background

The application of untreated manure and dung to agricultural areas provides essential fertilization for crops and pasture. However, it enables the entry of veterinary pharmacological compounds like antibiotics and heavy metals into the environment, thus promoting the distribution of resistant and potentially pathogenic microorganisms in soil, as well as ground and surface waters.

Objectives

The aim is to assess the risk of untreated agricultural wastes containing pathogens and antibiotic resistance genes on environment and human health while ascertaining the beneficial effects of manure treatment through fermentation.

Methods

Within a BMBF joint research project, we were able to monitor input and output samples of 15 German biogas plants, compiling extensive data on the chemical and microbiological status. Key organisms (e.g. Enterobactericeae, Enterococcaceae, Clostridiaceae) of every sample were genetically identified and archived in a strain collection to be tested for resistance and screened for tetracycline and sulphonamide resistance genes. Additionally, 4 resistance genes were quantified in all total DNA extracts and monitored in laboratory bioreactors under the influence of antibiotic supplementing over the course of 2 months.

Conclusions

Micro- and molecular biological approaches indicate a reduction of aerobic cultivable bacteria through fermentation including pathogens of the Enterobactericeae and Enterococcaceae without promotion of anaerobic and spore forming pathogens e.g. Clostridium spp. Although antibiotic resistance genes were found ubiquitously in both manure and digestate, a decline in resistance activity and resistance gene quantity after fermentation could be detected. Ultimately, biogas fermentation presents a
promising strategy to limit environmental pollution and risks for human health caused by agricultural wastes.
Background
Staphylococci are opportunistic pathogens that have emerged as a major cause of nosocomial infections. Methicillin resistance is associated with the mecA gene, which is located on a mobile genetic element called staphylococcal cassette chromosome mec (SCCmec). A new mecA homolog designated as mecA.LGA251 has been described recently.

Objectives
The purpose of this study was to evaluate multidrug resistant staphylococci recovered from hands of the general public.

Methods
267 staphylococci isolates were isolated from the general public in London, UK. All isolates were cultured, identified using MALDI-TOF-MS and tested for susceptibility to 13 antibiotics using disk diffusion method. In addition, MICs to oxacillin were also determined. mecA and mecC genes and SCCmec types were determined using PCR.

Conclusions
267 isolates belonging to 10 different staphylococcal species were recovered in this study. 223 (83.5%) staphylococcal isolates were resistant to more than two antibiotics. MICs to oxacillin varied from 0.06 mg/L to 256 mg/L. Seven out of 223 staphylococcal isolates were mecA positive and 3 mecC positive. SCCmec types were assigned for 4 out of 10 mecA and mecC positive strains. 2 SCCmec types V and 1 type VI were mecA positive and 1 SCCmec type IV was mecC positive.

We report on the widespread distribution of antibiotic resistant strains on the hands of the general public. We also report for the first time the isolation of mecA homologue mecC gene from randomly selected individuals. This emphasises the importance of good hygiene within the community.
EVALUATION OF CLASS 1-3 INTEGRONS AMONG PSEUDOMONAS AERUGINOSA CLINICAL ISOLATES IN MILAD HOSPITAL, TEHRAN, IRAN

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Background
Pseudomonas aeruginosa is a Gram-negative opportunistic pathogen that is found in soil, water and moist environments. P. aeruginosa infections are major cause of hospital morbidity and mortality.

Objectives
The current study aimed to demonstrate the frequency of class 1, 2 and 3 integrons in P. aeruginosa clinical isolates collected from urinary tract infections in Milad Hospital, Tehran, Iran.

Methods
Sixty P. aeruginosa strains were isolated from urinary tract infections during period December to March 2012. Antibiotic susceptibility testing was performed by disk diffusion method according to CLSI. Then, DNA of isolates were extracted and the presence of class 1, 2 and 3 integrons were examined by PCR.

Conclusions

Results: The antibiotic susceptibility pattern of P. aeruginosa clinical isolates were as follows: resistance to cefepime, ceftriaxone, Aztreonam, gentamicin, amikacin, ciprofloxacin, ofloxacin, ceftazidime, piperacillin tazobactam, and imipenem were 73.3%, 40%, 51.6%, 31.7%, 26.6%, 43.3%, 45%, 35%, 23.3%, and 15%, respectively. Based on our analysis, 73.3% and 1.6% of P. aeruginosa clinical isolates were positive int-1 and int-2, respectively. The negative results obtained for int-3.

Conclusion: Our findings demonstrated high frequency of class I integron in P. aeruginosa clinical isolates and therefore, the class I integron might play a special role in transfer of antibiotic resistant genes
Antimicrobial resistance

PHENOTYPIC DETECTION OF EFFLUX PUMPS IN MULTI-DRUG RESISTANT PSEUDOMONAS AERUGINOSA ISOLATED FROM SUEZ CANAL UNIVERSITY HOSPITAL

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Background

Infections caused by Pseudomonas aeruginosa are often severe, life threatening and difficult to treat because of the limited susceptibility to antimicrobials, the high frequency of emergence of antibiotic resistance during therapy and the emergence of multi-drug resistant (MDR) strains that develop resistance by various mechanisms including efflux pumps. Efflux pumps contribute to multidrug resistance by expelling antibiotics out of cells.

Objectives

Detection of the role of efflux pumps in MDR Pseudomonas aeruginosa isolates from Suez Canal university hospital in Ismailia.

Methods

The study included 49 Pseudomonas aeruginosa strains isolated from 307 nosocomially infected patients. Antibiotic susceptibility tests were done by disc diffusion method. For detection of association of efflux pumps, strains proved to be MDR were further tested for minimal inhibitory concentration (MIC), using the agar dilution susceptibility method, of four antibiotics of different groups (ciprofloxacin, meropenem, cefotaxime and gentamicin) before and after addition of the efflux pump inhibitor carbonyl cyanide m-chlorophenyl hydrazone (CCCP).

Conclusions

Using disc diffusion, 28 strains (57.1%) were proved to be MDR. The efflux pumps-associated resistance was proved through reduction of MIC in 18 (64.3%) , but not in 10 (35.7), of the MDR strains for ciprofloxacin and meropenem: 7 strains for ciprofloxacin, 7 for meropenem and 4 strains for both drugs together. This approximately twice association suggests a role of efflux pumps in MDR Pseudomonas aeruginosa. Understanding the mechanisms by which these pumps
act and how to overcome its activity opens the door for restoring the antibiotic activity and constitute a promising target for novel antibacterial agents.
FEMS-2214
Antimicrobial resistance

FIRST DESCRIPTION IN EUROPE OF A CIPROFLOXACIN RESISTANT SALMONELLA TYPHIMURIUM HARBOURING AAC6'-IB-CR AND OQXAB IN AN INCHI2 PLASMID
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Background
Fluoroquinolones are critical antibiotics (AB) for the treatment of Salmonella non-typhoid infections. Plasmid-mediated quinolone resistance (PMQR) genes have been often associated with decreased susceptibility to these AB worldwide, but concomitant presence of oqxAB and aac6'-Ib-cr genes, increasingly described in Asia, is still scarce in Europe.

Objectives
To characterize the genetic background (clonality, AB resistance genes, their transferability and genetic environment) of ciprofloxacin resistant S.Typhimurium clinical isolates.

Methods
Two clinical isolates of S.Typhimurium resistant to ciprofloxacin [young children (feaces)/one hospital/2012/Portugal] were studied. Susceptibility to AB and detection of ESBL were assessed by disk diffusion and/or Etest methods (CLSI/EUCAST). Characterization of AB resistance genes, including topoisomerases (gyrA/gyrB/parC/parE) and PMQR [qnrA/qnrB/qnrC/qnrD/qnrS/qepA/aac(6')-Ib-cr/oqxAB] genes, and metals (copper/silver/mercury/arsenic/tellurite) tolerance genes was performed (PCR/sequencing). Clonality (XbaI-PFGE/MLST), class 1 integron, plasmid characterization (PCR/sequencing), conjugation/transformation assays and genomic location (I-Ceu/I-S1-PFGE hybridization) were also performed.

Conclusions
Presence of multidrug-resistant IncHI2~180Kb plasmid encoding oqxAB and aac-6'-Ib-cr in a ciprofloxacin resistant S.Typhimurium/ST34, also carrying gyrA mutation (Asp87Asn), is here firstly described in Europe. Co-resistance to nalidixic acid,
chloramphenicol-cmlA/floR/catB3, gentamicin-aac(3)-IV, kanamycin-aphA1, streptomycin-aadA1/strA-strB, sulfamethoxazole-sul1/sul2/sul3, tetracycline-tetB, trimethoprim-dfrA12, ampicillin-blaTEM/blaOXA-1 and genes coding for metals tolerance (merA/silA/pcoD/terF) were found. Transformation of PMQR carrying plasmid was associated with resistance acquisition to several antibiotics (cmlA/floR/catB3/aac(3)-IV/aphA1/aadA1/sul1/sul2/sul3/dfrA12/blaOXA-1) and an eight-fold increase on ciprofloxacin MIC. Interestingly, the IncHI2 plasmid carried oqxAB genes flanked by IS26 and an incomplete class 1 integron containing aac-6’-Ib-cr-blaOXA-1-catB3-arr3-qacEΔ1-sul1, atypical sul3-integron and terF. A similar PMQR genetic environment in Salmonella with IncHI2 circulating in Asia was observed, highlighting the role of these plasmids in oqxAB and aac-6’-Ib-cr global spread.
FEMS-0512
Antimicrobial resistance

ASSESSMENT OF MULTIPLE ANTIBIOTIC RESISTANT ENTEROCOCCI IN COMMERCIAL AND COMMUNAL CATTLE FAECES AND THEIR WATER SOURCES IN MAFIKENG, NORTH-WEST PROVINCE, SOUTH AFRICA
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Background
Enterococci are part of the intestinal microflora of most mammals.

Objectives
To isolate and determine the antibiotic resistance profiles of Enterococcus species from cattle farms in South Africa.

Methods
A total of 49 faecal and 16 water samples from 2 commercial farms (Molelwane and Rooigrond) and a communal farm (Gelukspan) were analysed for Enterococcus species. 138 potential isolates were obtained and 129 enterococci were positively identified. All the 129 isolates were subjected to antibiotic susceptibility tests. The antibiotic inhibition zone diameter data of the isolates were subjected to cluster analysis and the results expressed as dendrograms. Vancomycin and tetracycline specific PCR was also performed.

Conclusions
Enterococcus avium (61%) and Enterococcus faecium (32%) were the dominant species isolated from both diarrhoeal and healthy cattle compared to Enterococcus durans (5%) and S. bovis (2%). Only Enterococcus avium (13.9%) and Enterococcus faecium (10.1%) were isolated from water samples. Large proportion (30%) of the isolates were resistant to tetracycline compared to vancomycin (17%). The MAR phenotype GM-SMX-NAL-NIT-C-K-S was dominant (38%) among E. faecium isolated from communal and commercial cattle while GM-SMX-NAL-NIT-K-S was dominant among E. faecium (40%), E. durans (60%) and E. avium (42%) isolated healthy and diarrhoeal cattle in both the communal and commercial farms. None of the enterococci isolates were positive for the vanA and vanB genes but the vanC gene was detected in 20% and 1.6% of E. durans and E. avium, respectively. Cluster analysis could not distinguish between isolates from commercial and communal farms indicating that these isolates had similar antibiotic resistant profiles.
FEMS-1443
Antimicrobial resistance

MOROCCAN BIOACTIVE ACTINOBACTERIA ISOLATES PRODUCING EFFLUX PUMPS INHIBITORS (EPI) OF RESISTANT BACTERIA
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Background

Traditional antimicrobials antibiotics are increasingly suffering from the emergence of multidrug resistance among pathogenic microorganisms. Among the antibiotic resistance mechanisms, efflux pumps have recently received a particular attention. Hence, there is an acute need for new active agents.

Objectives

Reducing the rate of emergence of antibiotic-resistant is our objective by studying the ability of Moroccan actinobacteria isolates to produce natural efflux pumps inhibitors (EPI) of medical interest.

Methods

210 actinobacteria isolates were screened for their ability to produce efflux pumps inhibitors using agar diffusion method. As test strains we have used in this study the wild type strain E. coli AG100 and its mutant AG100A with non-functional pump efflux system and the two clinical strains of S. aureus: the sensitive one SA-1199 and its mutant SA1199B. Phe-Arg-\(\beta\)-naphtylamide (PA\(\beta\)N) which inhibits the efflux system of many bacteria was used as control. Finally the biological and chemical characterizations of active compounds were conducted from four liters fermentation culture of selected bioactive.

Conclusions

Our screening program showed that among the 210 screened actinobacteria isolates, 9% were able to produce EPI. From the most promising ones we have purified five different bioactive compounds and their structure elucidation were under investigation. Screening only a subsection of our natural product library led to purify five molecules as EPI capable of sensitizing Gram-negative and Gram positive bacteria to antibiotics to which they are ordinarily intrinsically resistant. This result demonstrates the great potential of this approach in expanding antibiotic effectiveness in the face of the growing challenge of resistance.
IN VITRO ACTIVITIES OF SEVEN ANTIFUNGAL DRUGS AGAINST CANDIDA AFRICANA, AN ATYPICAL AGENT OF VULVOVAGINAL CANDIDIASIS

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Background

Vulvovaginal candidiasis (VVC) is a common vaginal fungal infection, affecting 70–75% of young women at least once during their lifetime. Candida africana is an opportunistic yeast pathogen that was proposed as a new species within the C. albicans species complex in 2001. It has been reported from 12 countries predominantly as an agent of VVC. Although, recently a large variety of antifungal drugs are used for treatment of VVC, treatment is a big challenge.

Objectives

Therefore in the present investigation, we evaluated the in vitro antifungal susceptibility testing of 7 agents against C. africana isolated from vulvovaginitis patients.

Methods

5 strains out of 350 vaginal swabs were identified as C. africana based molecular methods. The MIC of AmB, FLU, ITC, VOR, POS, ISA, MICA, ANID were determined based on CLSI document M27-A3.

Conclusions

The isolates were susceptible to Amphotericin B (MIC= 0.25 μg/ml), fluconazole (MIC= 1 μg/ml),itraconazole (MIC range, 0.5 - <0.016), voriconazole (MIC range, 0.5 - <0.016), posaconazole (MIC range, 0.5 - <0.016), isavuconazole (MIC range, 0.5 - <0.016), micafungin (MIC range, 0.016 - <0.008), anidulafungin (MIC range, 0.031 - <0.008).

C. africana has been largely associated with superficial infections of the genital tract, despite it has been associated with deep-seated infections, suggesting its invasive potential. The global patterns of resistance/susceptibility to most antifungal drugs remain largely unknown in C. africana; However high MIC of voriconazole, flucytosine and terbinafine against C. africana has been demonstrated. Antifungal susceptibility testing should be determined before therapy to avoid treatment failures.
PROPIONIBACTERIUM ACNES IN BIOMATERIALS-ASSOCIATED INFECTION:
FREQUENCY AND ANTIMICROBIAL RESISTANCE

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Background
Propionibacterium acnes can cause invasive deep seated infections, usually in the setting of implantable devices and its antimicrobial resistance is beginning to emerge.

Objectives
P. acnes can act as an opportunistic pathogen causing invasive and chronic biomaterials-associated infection through a biofilm mode of growth. In this study we aimed to review the frequency and the antimicrobial resistance of P. acnes in infections involving implants.

Methods
From March 2011 to April 2014, cerebrospinal fluid specimens (CFS) of 46 patients with internal shunt infection (ISI), corneal scrapings (CS) of 55 patients with delayed postoperative endophthalmitis (DPE) developed after lens implantation, periprosthetic tissues (PT) of 11 patients with prosthetic shoulder infection (PSI) and 17 patients with prosthetic joints infection (PJI) were studied. Samples were inoculated on supplemented Schaedler agars and fluid thioglycolate medium and were incubated during 15 days in anaerobic jars (Oxoid) on anaerobic conditions. Anaerobic bacteria were identified by API 20 A (BioMerieux). Susceptibility testing for penicillin, ampicillin+sulbactam, cefoxitin, clindamycin, tetracycline, rifampicin was performed with E-test (BioMerieux).

Conclusions
P. acnes was isolated from 2(4.3%) CFS, 16(29%) CS, 1(9 %) PT sample of patients with PSI and from 2(11.7%)PT samples of patients with PJI. These 21 P. acnes strains were susceptible to penicillin, ampicillin+sulbactam, cefoxitin and rifampicin whereas 38% (8/21) were resistant to clindamycin (MIC 90:32 mg/L), 14.2% (3/21) were resistant to tetracycline (MIC90: 4 mg/L). P. acnes is most frequently found in CS of patients with DPE developed after lens implantation. Its increasing clindamycin and tetracycline resistance should be taken into consideration during the treatment of biomaterials-associated infection.
BACKGROUND

Gram negative non-fermenting bacteria related infections are among the major problems encountered in health care facilities. *Pseudomonas aeruginosa* and *Acinetobacter baumanii* are common and widely distributed non-fermenter pathogens which mostly cause opportunistic and nosocomial infections in patients at intensive care units as well as general wards.

OBJECTIVES

In this retrospective study, we aimed to present antimicrobial susceptibility trends of common non-fermenters (*P. aeruginosa* and *A. baumanii*) in a tertiary referral hospital in Istanbul, Turkey. All chosen antimicrobial agents are consistently tested during study period, which covers eight years from 2006 to 2013.

METHODS

All *P. aeruginosa* (n:1312) and *A. baumanii* (n:1809) isolates from various specimens of inpatients from various clinics including ICUs were tested for antibiotic susceptibility for certain agents, between January 2006 and December 2013. Isolates were identified by conventional methods and/or commercial identification systems - API 20 NE (2006-2007) and VITEK 2 (2007-20013). Susceptibility testing of non-fermenters (n:3121) is performed by Kirby-Bauer disc diffusion method and results are interpreted according to CLSI guidelines.

CONCLUSIONS

Multidrug resistance in *P. aeruginosa* and *A. baumanii* is a worldwide problem. The key point of our study is the demonstration of antibiotic susceptibilities of *P. aeruginosa* and *A. baumanii* isolates with more than 3000 isolates within a long period of eight years within our tertiary training hospital. Increasing carbapenem resistance is a major problem in our setting especially for *A. baumanii* strains. Among
tested aminoglycosides, tobramycin is found more effective than amikacin for
\textit{A.baumanii}. Quinolones were found quite effective against \textit{P.aeruginosa in vitro} and
even more effective than carbapenems for some years. But resistance rates were
found very high against \textit{A.baumanii}. 
FEMS-2143
Antimicrobial resistance

FUSIDIC ACID AND MUPIROCIN RESISTANCE RATES AMONG STAPHYLOCOCCUS AUREUS CLINICAL ISOLATES

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Background

*Staphylococcus aureus* is one of the major pathogens causing infections both in community and in health care settings. Both mupirocin and fucidic acid are topical antimicrobial agents that can be used for the treatment of staphylococcal skin infections and/or elimination of its nasal carriage.

Objectives

We aim to determine high level (HL) mupirocin resistance and *in vitro* activity of fusidic acid in clinical *S. aureus* strains isolated between 2011 and 2013.

Methods

A total of 208 *S. aureus* strains, 86 methicillin resistant *S. aureus* (MRSA) and 122 methicillin susceptible *S. aureus* (MSSA), were tested for HL mupirocin with a 200 µg disc according to the CLSI guidelines. Fusidic acid resistance was investigated by 10 µg disc according to the criteria of EUCAST. HL mupirocin resistance was also confirmed by PCR for *ileS2* gene by using “*ileS2-M1 5’ GTTTATCTTCTGATGCTGAG 3’ and *ileS2-M2 5’ CCCCAGTTACACCGATATAA 3’*” primers. Resistance to mupirocin and fusidic acid were found as 7.2% (15/208) and 17.8% (37/208). Resistance rates of mupirocin for MRSA and MSSA were detected...
as 15.1% (13/86) and 1.6% (2/122). Resistance rates of fusidic acid for MRSA and MSSA were determined as 25.6% (22/86) and 12.3% (15/122).

**Conclusions**

We detected lower resistance rates for mupirocin than fusidic acid among clinical *S. aureus* strains in our hospital. Today increased topical preparations and inappropriate clinical use of antibiotics seem major factors causing to resistance and clonal selection. Prescription of topical mupirocin or fusidic acid especially for prolonged treatment of skin infections and for decolonization of *S. aureus* from nares should be reconsidered.
Antimicrobial resistance

QUEST FOR NEW ANTIMICROBIAL DRUG TARGETS: GENOME-WIDE SCREENING OF ESSENTIAL GENES OF STREPTOCOCCUS SUIS AND VALIDATION OF YYCF/G TWO-COMPONENT SYSTEM AS ANTIMICROBIAL DRUG TARGET

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Background

Streptococcus suis is a major bacterial pathogen of swine, causing high mortality and economic losses worldwide. S. suis is also emerging as an important cause of meningitis in humans in South East and East Asia. Antibiotic resistance of S. suis clinical isolates has been reported worldwide.

Objectives

Identification of novel antimicrobial drug targets of S. suis and validation of some of these targets.

Methods

Essential genes of S. suis S10 have been identified in a high-throughput genome-wide screen using a mariner transposon insertion approach (Tn-Seq). We constructed a list of S. suis S10 essential genes based on Tn-Seq results and published lists of orthologous essential genes of three Gram-positive bacteria. Based on this we selected several putative drug targets. We generated conditionally knock-out of yyCF/G to elucidate its regulatory network and to validate its essentiality.

Conclusions

1) Genome-wide screening of a transposon insertion library of S. suis S10 revealed 281 essential genes of which 54 genes were shared with three other Gram-positive bacteria. 2) yyCF/G was the only essential Two-component System in four selected Gram-positive bacteria. 3) A conditional knock-out strain of yyCF/G in S. suis S10 validated its essentiality and potential as a drug target for novel antimicrobials.
Background
Yeast species identified in a previous study in the North West Province are associated with superficial mucosal infections and life-threatening diseases. Treatment is by available antifungal drugs. Due to continuous exposure of aquatic yeast species to sub-therapeutic levels of antifungal agents, these species have developed resistance.

Objectives
The aim of this study was to determine the diversity and antifungal susceptibility patterns of yeasts in the Harts River of the North West Province, South Africa.

Methods
Yeast colonies were enumerated using yeast-malt-extract (YM) agar (containing 100 ppm chloramphenicol). Purified isolates from 37°C incubation were identified by biochemical tests and 26S rRNA gene sequencing. Disc diffusion antifungal susceptibility tests were conducted on the yeast species.

Conclusions
Yeasts levels ranged between 320-4200 cfu/L and 27-2573 cfu/L for room temperature and 37°C. All the yeast colonies isolated were non-pigmented. DBB tests determined the yeasts isolates as ascomycetes. Yeasts isolates were identified and belonged to the genera Arxiozyma, Candida, Clavispora, Cyberlindnera, Leucythophora, Pichia, Saccharomyces, and Wickerhamomyces. Saccharomyces cerevisiae and Candida glabrata were most frequently isolated species. A large number of isolates were resistant to azoles, especially fluconazole as well as other antifungal classes. Most of the Candida species were resistant to almost all the antifungals. Several of the isolated yeast species are opportunistic pathogens. These could cause infections in sensitive individuals during occasional direct contact. A major health concern is that these infections may not respond to available antifungal agents.
Antimicrobial resistance

C. PARAPSILOSIS AZOLE RESISTANCE CONFERRED BY A GAIN-OF-FUNCTION MUTATION IN MRR1 TRANSCRIPTION FACTOR
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Background
The human fungal pathogen, Candida parapsilosis, is mostly isolated from patients with bloodstream infections and its incidence in some countries is equal or exceeds that of C. albicans. For the treatment of these infections, azoles are widely used as prophylactic and therapeutic drugs. However, C. parapsilosis clinical azole resistance has been increasing in the last decade and little is known about the antifungal resistance mechanisms involved in the development of such resistance. Recently, voriconazole resistance mechanisms were associated with MDR1 (encoding for a multidrug efflux pump) and MRR1 (transcription factor) genes overexpression. Furthermore, a mutation in the MRR1 gene (A2619C) that resulted in an alteration of lysine to asparagine, K873N, in polypeptide chain was firstly reported.

Objectives
Knowing that gain-of-function mutations found in transcription factors genes play a crucial role in azole resistance acquisition, we intend to elucidate the importance of A2619C mutation in MRR1 gene in C. parapsilosis azole resistance.

Methods
For that, MRR1_RVRC mutated gene was integrated in its native locus in a double deleted strain, Δmrr1/Δmrr1. Azole susceptibility and gene expression profile of the set of strains used in this study were characterized.

Conclusions
The MRR1_RVRC complemented strain changed azole susceptibility profile, from susceptible to resistant, and also displayed an up regulation of MRR1 and MDR1 genes. Thus, missense mutation A2619C described in MRR1 transcription factor is a
gain of function mutation, decisive for azole resistance. This study was supported by FEDER, COMPETE and FCT (PTDC/DTP-EPI/1660/2012).
Background

In Portugal, *C. parapsilosis* is the second most common yeast species isolated from patients with bloodstream infections. However, *C. parapsilosis* increasing prevalence was been found in other European countries and in Latin America and Asia. Azoles are widely used as prophylactic and therapeutic drug; but *C. parapsilosis* acquires azole resistance in a rapid and stable manner.

Objectives

This study aims to characterize antifungal susceptible profile of *C. parapsilosis* sensu strico etiologic agent of human infections from two major Portuguese hospitals, Centro Hospitalar de Coimbra and Centro Hospitalar São João and uncover the molecular mechanisms underlying antifungal resistance.

Methods

Clinical isolates (n=93) collected from distinct body sites/products were depicted regarding antifungal susceptibility profile to azoles (fluconazole, voriconazole, posaconazole) and echinocandins (caspofungin, micafungin, anidulafungin). The minimal inhibitory concentration (MIC) of each antifungal drug was determined according to the M27-A3 protocol of the Clinical and Laboratory Standard Institute. Gene expression profile of the resistant isolates was carried out by RT-PCR.

Conclusions

Among *C. parapsilosis* isolates, 10% were resistant to fluconazole; 6% were resistant to voriconazole and 2% were resistant to posaconazole. The incidence of susceptible–dose-dependent found was 6% for FLC and 10% for VRC. No resistance was found for echinocandins. Azole resistance development was mainly associated with *MDR1* (encoding multidrug efflux pump) and *MRR1* (transcription factor that
regulates \textit{MDR1} expression) genes overexpression. Our study showed that prevalence of azole \textit{C. parapsilosis} resistant isolates has increased dramatically since the last survey carried out in 2009. This study was supported by FEDER, COMPETE and FCT (PTDC/DTP-EPI/1660/2012).
Background

C. jejuni has shown increasing resistance to quinolone. The resistance is mediated by a point mutation in the quinolone resistance-determining region (QRDR) of gyrA with the high frequency at position 86, followed by position 90. However, no direct evidence of the association between these mutations and the quinolone resistance has been shown to date.

Objectives

To clarify the impact of an amino acid substitution at positions 86 or 90 in C. jejuni GyrA on DNA gyrase activity in the presence or absence of quinolones

Methods

The recombinant DNA gyrase with amino acid substitutions at position 86, and 90 in GyrA were expressed in Escherichia coli and purified using Ni-NTA agarose column chromatography. Wild-type and mutant DNA gyrase were reconstituted in vitro by mixing recombinant GyrA and GyrB. Correlation between the amino acid substitution and quinolone resistance was assessed by quinolone-inhibited supercoiling assay.

Conclusions

The quinolone-inhibited supercoiling assay demonstrated the important role of these amino acid substitutions in reduced sensitivity to quinolone with mark influence by amino acid substitution, especially Thr86Ile. This study confirmed the association of these mutations with the quinolone resistance in C. jejuni and demonstrated that sitafloxacin might be a candidate for the treatment of campylobacteriosis caused by ciprofloxacin-resistant C. jejuni.
Background
The bacterial SOS response to DNA damages is involved in antibiotic resistance acquisition and dissemination. It can regulate expression of resistance genes (eg \textit{qnrB}), or acquisition of resistance genes (eg expression of the integrase of integrons, \textit{IntI}). Integrons are genetic elements that play a major role in the dissemination of antibiotic resistance in Gram-negative bacteria. \textit{IntI} catalyses insertion and excision of resistance genes embedded within gene cassettes. Dissemination of resistance genes might also be correlated to the SOS response as shown for bacterial conjugation. Various stresses including antibiotics can induce the SOS response.

Objectives
Our aim was to estimate the level of the SOS response and of the \textit{IntI} expression and activity \textit{in vivo} in the mouse digestive tract.

Methods
Germ-free mice were inoculated intragastrically with a strain of \textit{Escherichia coli} carrying two plasmids: one harbouring the \textit{intI1} gene under the control of its own promoter, the second containing a gene cassette array. Efficiency of bacterial colonisation, plasmid stability and \textit{IntI1} activity were estimated from mice feces. \textit{intI1} and \textit{sfiA} (SOS regulon) genes expression level was estimated by qPCR.

Conclusions
We observed a higher expression of the SOS response and of \textit{IntI1} \textit{in vivo} than \textit{in vitro} and a 1.5 log increase of the \textit{IntI1} excision frequency.

Our results indicate that the digestive tract is an environment where the SOS response is induced. Moreover, the higher expression and activity of integrase highlight the role of the gut in disseminating antibiotic resistance.
FEMS-0889
Antimicrobial resistance

A DIFFERENT ATP-BINDING CASSETTE (ABC) SYSTEM LIKELY ASSOCIATED WITH LINEZOLID RESISTANCE IN STAPHYLOCOCCUS HAEMOLYTICUS

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Background
Linezolid is an oxazolidinone antimicrobial agent for treating infections caused by multi-drug resistant Gram-positive cocci. Mutations in domain V of 23S rRNA, an adenosine modification catalyzed by Cfr and mutations in the ribosomal proteins L3, L4 and L22 are mechanisms that have emerged in linezolid-resistant isolates. We identified a clinical Staphylococcus haemolyticus strain (ShRL1) exhibiting a high level of minimum inhibitory concentrations (MIC, 64µg/mL) for linezolid, which did not show any of these mechanisms. Moreover, the ShRL1 strain shows genes encoding proteins of the ATP-binding cassette (ABC) superfamily which were not identified in its susceptible derivative ShSL2 (MIC, 1µg/mL) after 7 subcultures under no selective antibiotic pressure.

Objectives
To compare the whole genome sequencing of the ShRL1 and ShSL2 strains in order to determine a cause of linezolid resistance.

Methods

Genome sequencing was performed using the Ion Torrent PGM platform (Life Technologies) kit400bp. The sequences were de novo assembled using Mira v4.0.2 and spades v3.1.1. Scaffold was obtained using CLC Genomics Workbench v7.5 and Geneious 8.0.5 with the genome reference S, haemolyticus JCSC1435. Lastly comparative analyses were performed using Mauve v3.2.

Conclusions
Rare studies have associated efflux-pump systems with linezolid decreased susceptibility. We propose a resistance mechanism other than target site modifications that may be responsible for elevated MIC for linezolid in *S. haemolyticus*. Further studies are necessary to support it. However, our data strongly suggest that the ATP-binding cassette (ABC) system, which share about 70% sequence homology to efflux proteins of *S. aureus*, has an important role in linezolid resistance in *S. haemolyticus*. 
THE ANTIBACTERIAL EFFICACY OF DIFFERENT MOUTHＲINＳＥＳ ON PERIODONTAL PATHOＧENＳ

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Background

The widespread use of systemic antibiotics in medicine and dentistry becomes an increasingly serious problem by eliciting drug resistance and undesirable side effects. In dentistry, there are several antiseptic mouthrinses for topical use available, showing different mechanisms of action to inhibit dental pathogens, for instance those associated with periodontitis and peri-implantitis.

Objectives

Aim of the study was to investigate the antimicrobial efficacy of eight antiseptic mouthrinses containing six different types of active ingredients for the use on planktonic periodontitis-associated pathogens.

Methods

Six representative periodontitis/peri-implantitis associated pathogens were grown anaerobically for 24h. After suspension, the six-strain-ｃocktail was exposed to the following agents for 30 seconds:

1. NaCl (saline as negative control), 2. CHX (0.2%) (chlorhexidine), 3. CHX (0.06%),
4. CHX+CPC (CHX 0.05% + cetylpyridinium chloride 0.05%), 5. PH (polyhexanide 0.04%), 6. PH 0.1%+betaine, 7. Oct (octenidine dihydrochloride + phenoxyethanol),
8. FF (amine fluoride/stannous fluoride), and 9. EO (mouthwash containing essential oils). Microbial viability was monitored by both colony growth (CFU/ml) and fluorescence-based vitality. Six independent series of experiments were conducted. Statistical analysis of data was performed using mean-based 95% confidence intervals, ANOVA analysis and the Tukey test.

Conclusions

The strongest antibacterial CFU reduction of 8.75 log10 units could be shown for the agents 2, 4 and 7. Furthermore, strong colony growth reduction of 7.25-6.36 was
caused by the agents 6, 8 and 9. Moderate to low antibacterial activity was induced by the exposure to agents 3 and 5 (log\textsubscript{10} CFU reduction of 3.80-2.25). Bacterial vitality was strongly reduced by all antiseptic agents.
CHARACTERIZATION OF ESBL PRODUCING ESCHERICHIA COLI IN NORTHERN THAILAND

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Background

The prevalence of extended spectrum beta-lactamase (ESBL) producing Escherichia coli has reached to 41-44 % of total E. coli infections in a tertiary center Maharaj Nakorn Chiang Mai Hospital (MNCMH) during past six years. To date, there is no report on distribution of ESBL genes and levels of resistance to third generation cephalosporins among E. coli clinical isolates from any hospital in Northern Thailand.

Objectives

To study the correlation between distribution of ESBL genes and levels of resistance to cefotaxime and ceftazidime.

Methods

435 E. coli strains collected from MNCMH during April to May, 2010 were screened for production of ESBL and AmpC BL. MICs of cefotaxime and ceftazidime were determined in ESBL producers. ESBL and AmpC BL genes were detected using multiplex PCR and nucleotide sequencing.

Conclusions

Of 182 ESBL producers, 25 strains also produced AmpC BL. The MIC₅₀/MIC₉₀ of CTX and CAZ were 128/512 and 16/128 μg/ml, respectively. ESBL genes detected were blaTEM (82.9%), blaCTX-M (80.7%; blaCTX-M-group 1 and blaCTX-M-group 9), and blaSHV (11%). blaCTX-M-15 and blaCTX-M-65 predominated among blaCTX-M-group 1 whereas blaCMY2 among AmpC BL producers. Most strains carried multiple beta-lactamase genes. The predominant genotypes were blaTEM + blaCTX-M-group 1 and blaTEM + blaCTX-M-group 9. Levels of resistance to cefotaxime and ceftazidime depended on type of beta-lactamase gene, not the accumulating number of genes. The presence of blaCTX-M-group 1 was associated with resistance to both cefotaxime and ceftazidime. However, the
presence of \textit{bla}CTX-M-group 9 was associated with more resistance to cefotaxime and more sensitivity to ceftazidime.
FITNESS STUDIES OF PZA RESISTANT MYCOBACTERIUM TUBERCULOSIS STRAINS

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Background

Pyrazinamide (PZA) is the only first line tuberculosis (TB) drugs to have significant activity against dormant TB. PZA resistance in Mycobacterium tuberculosis is most frequently conferred by mutations in pyrazinamidase (PZase), the enzyme required to activate PZA. These resistance mutations are sometimes stated to have no fitness cost, but in fact laboratory evidence seems absent for either a fitness cost or lack thereof.

PZase is non-essential during active growth, but its transcription is upregulated during oxygen and nutrient limitation. The role of PZase in NAD salvaging supports requirement of PZase during dormancy, and thus PZA resistant strains without functional PZase may have a measurably decreased fitness. Epidemiological data showing only limited spread of PZA resistant strains also suggests a lower fitness.

Objectives

Objective of this study is to determine the fitness of PZA resistant TB strains in a dormancy model.

Methods

Isogenic PZA susceptible and resistant strains of M. tuberculosis are cultured in the Wayne hypoxic dormancy model in non-selective media, and viability investigated over time.

Conclusions

In the first 2 months of dormancy model culture, proportions of both the PZA resistant and susceptible bacteria remained viable as determined by live/dead staining and culture. Auramine/NileRed stainings demonstrate decreasing proportions of acid-fast bacilli and increasing proportions of lipid inclusion over time in susceptible and resistant strains.
No measurable difference in fitness was observed between the strains during the first two months, viability will be further assessed for additional time points to assess the long term survival of these dormant cultures.
Antimicrobial resistance

ANTIBACTERIAL ACTIVITY AND MECHANISM OF ACTION OF EUGENOL AGAINST MULTI-DRUG RESISTANT PATHOGENIC BACTERIA

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Background

Antibiotic resistance in clinical isolates is an increasing life-threatening global threat. Newer approaches and drugs are required to control these organisms in the current scenario. The in vitro antibacterial activity and mechanism of eugenol, a natural phenylpropene was studied against multi-drug resistant clinical isolates.

Objectives

The present study was conducted to evaluate the antibacterial property of eugenol against multi-drug resistant clinical bacterial pathogens. Further, minimum inhibitory concentration and bactericidal concentration of eugenol along with mechanistic action was determined.

Methods

Antimicrobial minimum inhibitory concentration and bactericidal concentration experiments with eugenol was measured against a range of Gram positive and Gram negative bacteria isolated from clinical specimens using Resazurin micro-titre assay. Analysis of killing kinetics of eugenol was performed against two strains (Escherichia coli and Staphylococcus aureus). The effect of divalent salts on the activity of eugenol was determined. Bioluminescence method was used to determine the effect of eugenol on the intracellular pool of bacterial ATP. Membrane depolarization studies were conducted to study the cell wall permeabilization. The morphological changes in bacterial membrane due to the activity of eugenol were investigated using scanning and transmission electron microscopy studies.

Conclusions

The results provide additional experimental data that are consistent with the hypothesis that eugenol represents a new class of antibacterial agents for treating multi-drug resistant pathogenic bacteria and acts via different mechanism from conventional antibiotics. The results suggest that antibacterial activity is due to
interaction of eugenol on bacterial cell membrane. The results also revealed that cell wall and membrane of treated bacteria were significantly damaged.
PREVALENCE AND CHARACTERIZATION OF BLA CTX-M, BLA TEM AND BLA SHV FROM ESBL PRODUCING ORGANISMS FROM HOSPITALS IN LAGOS, NIGERIA.

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Background

The increasingly prevalent of Extended-Spectrum β-Lactamase (ESBL) among clinical isolates are becoming widespread globally. Most reports on ESBL from Nigeria have always rested on their phenotypic statuses with just a few with information on their molecular types.

Objectives

We therefore set out to characterize the Beta-lactamases in multidrug resistant clinical isolates of Enterobacteriaceae by molecular techniques.

Methods

One hundred and sixteen non-duplicate clinical isolates from 3 hospitals including a Teaching hospital were screened for ESBL production, 42 of them were phenotypically confirmed as ESBL producers. Molecular analysis by PCR and Pulsed Gel Electrophoresis used in the detection of TEM, SHV and CTX-M ESBL genes revealed 100% prevalence for CTX-M, 57.1% TEM and 64.3% SHV. Klebsiella pneumoniae and E. coli respectively had the most frequency of ESBL genes with Serratia rubidae with the lowest frequency of occurrence. All ESBL producers showed resistance to at least 2 or more classes of antibiotics and varied resistance to meropenem (carbepanem).

Conclusions

Despite several researches and reports on ESBL in Nigeria, there is still no known awareness for its routine detection in the private and public hospital laboratories. It
seems as though the fact is simply ignored. This study has brought to light the extent of the spread of these organisms in Lagos and call for stringent policies to be put in place to reduce antibiotic misuse and abuse.
RETROSPECTIVE DETECTION OF THE HIDDEN SPREAD OF OXA-48-PRODUCING KLEBSIELLA PNEUMONIAE IN A FRENCH TEACHING HOSPITAL

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Background

In our hospital, an outbreak of OXA-48-producing \textit{Klebsiella pneumoniae} (OXA48KP) was identified in the end of 2013. For these strains, the Vitek2 System (V2S) categorised the ertapenem intermediate or resistant (I/R), whereas it could have been categorised susceptible by E-test (E-test is systematically performed if ertapenem is categorised I/R with the V2S). Before the outbreak, no screening for carbapenemases was performed for isolates identified as susceptible to ertapenem by E-tests.

Objectives

The objective of this study was to determine the presence of OXA48KP before the moment of the outbreak detection.

Methods

From January 2010 to the end of 2013, all \textit{K. pneumoniae} isolates for which ertapenem was tested by E-test were selected. These isolates were then tested for the presence of carbapenemase with the KPC, MBL and OXA-48 confirm Kit (ROSCO Diagnostica, Taastrup, Denmark). The isolates with an inhibition diameter around temocillin $<12$ mm were then tested for OXA-48 by PCR and compared by PFGE.

Conclusions

Among the 21 isolates included, 3 were identified as OXA48KP. PFGE analysis demonstrated that two of them were closely-related to the outbreak isolates and one was not related. They had been initially considered susceptible by E-test (MIC from 0.25 to 0.5 mg/L) and had been identified from samples collected within the 6 months preceding outbreak detection. Therefore, there was probably a hidden spread of the outbreak before triggering the alert at the end of 2013. In addition, these results show the high sensitivity of the V2S for carbapenemase detection.
CHARACTERIZATION OF THE RESISTANCE MECHANISM TO DITYROMYCIN OF THE STREPTOMYCES PRODUCER STRAIN.
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Background

Dityromycin is a peptide antibiotic produced by *Streptomycyes sp.* strain AM-2504 with the structure highly related to that of GE82832. The crystal structure of Dityromycin - 70S ribosome complex revealed that the binding involves conserved amino acid residues of ribosomal protein S12. Dityromycin inhibits EF-G activity by disrupting critical contacts between EF-G and S12 that are required to stabilize the post-translocational conformation of EF-G, thereby preventing the ribosome-EF-G complex from entering a productive arrangement for translocation.

Objectives

The aim of this work was to identify the mechanism of resistance to Dityromycin of the producer strain *Streptomycyes sp.* AM-2504.

Methods

Cell-free *in vitro* translation tests, genome walking and direct sequencing of the *rpsL* gene were used to characterize the mechanism of resistance of this antibiotic.

Conclusions

The results obtained in this work indicate that the resistance of the producer strain to Dityromycin is due to a modification of its ribosomes. In particular, three amino acids changing mutations were found in an highly conserved region of the protein S12 corresponding to the binding site of the antibiotic. These mutations cause a substantial loss of affinity of Dityromycin for the 30S ribosomal subunit thus conferring resistance to the producer strain. Results of this study contribute to the knowledge of the molecular determinants developed and successfully implemented by the antibiotic production/resistance machinery.

References

HAS THE USE OF FLUOROQUINOLONES CONTRIBUTED TO THE WIDESPREAD DISSEMINATION OF METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS AND EXTENDED-SPECTRUM B-LACTAMASE-PRODUCING KLEBSIELLA PNEUMONIAE IN THE HEALTHCARE SETTING?

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Background

We recently demonstrated that fitness cost associated with resistance to fluoroquinolones is diverse across clones of both methicillin-resistant Staphylococcus aureus (MRSA) and extended-spectrum β-lactamase (ESBL)-producing Klebsiella pneumoniae in adult hospital wards. We observed that major clone MRSA and ESBL-producing K. pneumoniae strains with high-level resistance to fluoroquinolones retain their fitness while the vitality of other S. aureus and K. pneumoniae isolates become more compromised during the development of resistance to these antibiotics. Our observation with MRSA was subsequently confirmed by British scientists. In addition our findings may account for the replacement of SHV type ESBL genes by CTX-M-15 in K. pneumoniae reported during the last decade.

Objectives

Investigation of whether or not this mechanism - the selection of the major clones of MRSA and ESBL-producing K. pneumoniae by fluoroquinolones - could have contributed to the widespread dissemination of both pathogens.

Methods

Review of related literature.

Conclusions

An overwhelming part of the related literature supports the assumption.

Our findings suggest that a more judicious use of fluoroquinolones could significantly decrease the rates for both MRSA and ESBL-producing K. pneumoniae in the adult healthcare setting.
WHICH ANTIBIOTIC SHOULD BE PRESCRIBED FOR ERADICATION OF KLEBSIELLA PNEUMONIAE NOSOCOMIAL INFECTIONS IN ILAM HOSPITALS? AMOXICILLIN OR AMIKACIN?
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Background
The relax prescription of antibiotics in Iran made an evolution in bacterial populations. Antibiotic resistance strains specially noscomial strains cause too issues for treatment of bacteria responsible for nosocomial infections. Klebsiella pneumoniae is also known as a hard to treat bacterium in Ilam hospitals and the rate of resistance is different from antibiotic to antibiotic.

Objectives
Therefore, the current study aims to evaluate the prevalence of different antibiotics for treatment of K. pneumoniae clinical isolates and open new insight for antibiotic suspension or prescription.

Methods
48 K. pneumoniae clinical isolates were identified as standard procedures during March 2013 to March 2014 in Ilam hospitals. The isolates were collected from in and out patients with urinary tract infections. The isolates were then subjected to antibiotic susceptibility testing via disk diffusion method as CLSI guideline. The antibiotic resistance pattern and significant difference among K. pneumonia and different antibiotics were calculated by spss 16 via chi-square.

Conclusions
The antibiotic susceptibility pattern demonstrated 39.6%, 27.1%, 27.1%, 16.7%, 39.6%, 37.5%, 10.4%, 54.2%, 8.3%, 75%, 27.1%, 50%, and 97.9% resistance to gentamicin, ciprofloxacin, cefoxitin, pipracilin, aztreonam, choloramphenicol, nitrofurantein, cephalotin, amikacin, ceforuxim, tetrycycline, cepazolin, and amoxicillin, respectively. The findings demonstrated high resistance in K. pneumoniae clinical strains. This study recommended the new policy with prescription of amikacin and non-prescription of amoxicillin as a dominant susceptible and resistance antibiotic in K. pneumoniae clinical isolates.
FEMS-2127
Antimicrobial resistance

ADES MUTATIONS IN MDR-ACINETOBACTER BAUMANNII CLINICAL ISOLATES WITH TIGECYCLINE REDUCED SUSCEPTIBILITY FROM CAIRO, EGYPT.
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Background
Infections caused by multidrug resistant (MDR) Acinetobacter baumannii (A. baumannii) especially in intensive care units have limited therapeutic options, one of them is Tigecycline. Overexpression of the adeABC efflux pump may be caused either by the ISAba-1 insertion or by specific point mutations in adeR and adeS, therefore plays a major role in conferring MDR- A. baumannii.

Objectives
We aimed in our study to monitor the TGC susceptibility and to study the role of ISAba-1 and the adeS regulator within the AdeABC efflux pump among MDR- A. baumannii clinical isolates.

Methods
Over 18 months from April 2012 to September 2013, 63 MDR-A. baumannii identified by detection of OXA-51 like gene were isolated from ICU patients. TGC MIC was determined by E test and confirmed by broth microdilution. PCR analysis of adeR, adeS, adeB, and ISAba1 genes were done with further sequencing of adeS gene.

Conclusions
Reduced susceptibility to TGC (MIC: 3-4ml/L) was noticed in 6/63 (9.5 %) MDR- A. baumannii isolates
*baumannii* isolates, insertion sequence IS*Aba*-1 was detected in 3 isolates, two of them showed amino acid substitutions in the ade*S* operon by sequencing. IS*Aba*-1 may play a role in conferring reduced TGC susceptibility in MDR- *A. baumannii* in association with amino acids mutations in the ade*S* gene.
MOLECULAR CHARACTERIZATION AND EPIDEMIOLOGY OF CEFXITIN RESISTANCE AMONG ENTEROBACTERIACEAE LACKING INDUCIBLE CHROMOSOMAL AMPC GENES FROM HOSPITALIZED AND NON-HOSPITALIZED PATIENTS IN ALGERIA

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Background

Plasmid-mediated AmpC β-lactamases have emerged and are being reported worldwide with varying prevalence rates. In Algeria, only few reports on plasmid-encoded AmpC in Enterobacteriaceae strains recovered from hospital settings were published.

Objectives

The aim of this study was to investigate the prevalence and molecular epidemiology of cefoxitin resistance among Enterobacteriaceae isolates recovered from hospitalized and non-hospitalized patients in Bejaia locality (Algeria). The association of pAmpC with ESBL and plasmid-mediated quinolone resistance determinant was also studied.

Methods

In this study, 922 consecutive non-duplicate clinical isolates of Enterobacteriaceae obtained from hospitalized and non-hospitalized patients at Bejaia, Algeria, were analyzed for AmpC type β-lactamases production. The ampC genes and their genetic environment were characterized using PCR and sequencing. Plasmid incompatibility groups were determined using PCR-based replicon typing. Phylogenetic grouping and Multilocus sequence typing (MLST) were determined for molecular typing of the plasmid encoded AmpC (pAmpC)-producing isolates.

Conclusions

Of the isolates, 15 were identified as AmpC producers including 14 CMY-4-producing isolates and one DHA-1-producing K. pneumoniae. All AmpC-producing isolates co-expressed the broad-spectrum TEM-1 β-lactamase and three of them co-produced CTX-M and/or SHV-12 ESBL. Phylogenetic grouping and virulence genotyping of the
*E. coli* isolates revealed that most of them belonged to groups D and B1. MLST analysis of *K. pneumoniae* isolates identified four different STs with two new sequences: ST1617 and ST1618. Plasmid replicon typing indicates that *bla*<sub>CMY-4</sub> gene was located on broad host range A/C plasmid, while LVPK replicon was associated with *bla*<sub>DHA-1</sub>. All isolates carrying *bla*<sub>CMY-4</sub>, displayed the transposon-like structures ISEcp1/ΔISEcp1-blaCMY-blc-sugE.
Background
Vancomycin-resistant enterococci (VRE) are among the most common antibiotic-resistant nosocomial pathogens in health care in general and in intensive care units (ICUs), in particular.

Objectives
The main purpose of this study is to distinguish between lower respiratory tract (LRT) colonization and infection due to enterococci using semi-quantitative cultures.

Methods
Sixty-eight lower respiratory tract (LRT) specimens were collected from consecutive adult patients who underwent Lung transplantation and were developed ventilator associated pneumonia (VAP) from three teaching hospitals in Tehran, Iran, during December 2009 – September 2012. A semi-quantitative threshold of \( >10^4 \) cfu/ml was considered to be diagnostic for Enterococcal-VAP. Antibacterial resistance patterns of isolates to 13 antimicrobial agents were analyzed according to EUCAST. Pulsed Field Gel Electrophoresis (PFGE) was performed as described previously (Feizabadi, M.M. et al. J Microbiol Methods 2011; 84:144 –146).

Conclusions
Twenty-three and four isolates of \( E. \) faecalis and \( E. \) faecium showed \( >10^4 \) cfu/ml which corresponds to infection. Linezolid, teicoplanin and vancomycin were found to be the most effective drugs.
The results of MIC for vancomycin and teicoplanin are shown in the table. Seven and two different DNA banding patterns were identified among \( E. \) faecalis and \( E. \) faecium isolates, respectively.
Infection of LRT with enterococci seems to become serious and needs to be reconsidered as a real problem rather than colonization. In such cases clones of VRE is associated with high mortality.

<table>
<thead>
<tr>
<th>Number of isolates</th>
<th>MIC of vancomycin 64≥mg/mL</th>
<th>MIC of vancomycin 256≥mg/mL</th>
<th>MIC of teicoplanin</th>
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<tr>
<td></td>
<td>256 ≥ mg/mL</td>
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<tr>
<td><strong>E. faecalis</strong></td>
<td>26 7 5 4</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>E. faecium</strong></td>
<td>4 4 4 4</td>
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</table>
EVALUATION OF MULTIPLEX TANDEM REAL TIME PCR KIT FOR RAPID IDENTIFICATION OF GRAM-POSITIVE COCCI FROM BLOOD CULTURES AND DETECTION OF MECA AND VAN GENES

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Background

Currently available automated blood culture systems have significantly reduced bacteria detection time. However; so as to get optimum treatment, more rapid identification systems are required.

Objectives

Gram Positive 12 Kit (Aus Diagnostics, Australia) makes it possible to identify common Gram positive bacteria (Staphylococcus spp., Streptococcus spp., Enterococcus spp.) and to detect common antibiotic resistant genes (mecA, vanA, vanB). The aim of this study was to rapidly identify and to assess the performance of this test for sepsis agent Gram positive bacteria and resistance genes.

Methods

54 patients having positive blood cultures and Gram-positive cocci in Gram staining were included in the study. After the DNA isolation, identification and resistance genes were determined by using Easy Plex pipetting robot, and then Rotor GeneQ (Qiagen, Switzerland) real-time PCR device for pre-PCR and PCR set up phases with this multiplex tandem real time PCR kit. Results were compared to the results obtained from PHOENIX (BectonDickinson, USA) and Vitek 2 Compact (bioMérieux, France) automated systems.

Conclusions

42 results were completely as same as the results from PHOENIX and Vitek 2 Compact automated systems. Additionally; 12 polymicrobial infections which could not be detected by automated systems were successfully detected.

In conclusion; the kit which was evaluated proved to be both rapid (approximately 2,5 hours) and reliable in determining common Gram positive sepsis pathogens and resistance genes. According to our results, it could be concluded that the test could easily be used in patients who are suspected of sepsis in routine diagnosis.
RAPID IDENTIFICATION OF GRAM-NEGATIVE PATHOGENS FROM BLOOD CULTURES AND DETECTION OF RESISTANCE GENES

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Background

Currently available automated blood culture systems have significantly reduced bacteria detection time. However; so as to get optimum treatment, more rapid identification systems are required.

Objectives

Gram negative 12 Kit (Aus Diagnostics, Australia) makes it possible to identify common Gram negative bacteria (Enterobacteriaceae, Pseudomonas spp., Pseudomonas aeruginosa, Escherichia coli, Klebsiella pneumoniae) and to detect common antibiotic resistant genes. The aim of this study was to rapidly identify and to assess the performance of this test for sepsis agent Gram negative bacteria and resistance genes.

Methods

78 patients having positive blood cultures and Gram-negative bacilli in Gram staining were included in the study. After the DNA isolation, identification and resistance genes were determined by using Easy Plex pipetting robot, and then Rotor GeneQ (Qiagen, Switzerland) real-time PCR device for pre- PCR and PCR set up phases with this multiplex tandem real time PCR kit. Results were compared to the results obtained from PHOENIX (Becton Dickinson, USA) and Vitek 2 Compact (bioMérieux, France) automated systems.

Conclusions

All of the pathogenes were succesfully identified by using this test. Oxa-23 gene, aadA1 gene, pan-vim genes, aadB gene were found in 5, 15, 3, 2 samples; respectively.

In conclusion; the kit which was evaluated proved to be both rapid (approximately 2,5 hours) and reliable in determining common Gram negative sepsis pathogens and resistance genes. According to our results, it could be concluded that the test could easily be used in patients who are suspected of sepsis in routine diagnosis.
Background
Dermatophytosis is one of the most common fungal infections in humans. In vitro antifungal susceptibility testing is considered as a powerful method to seek the drug of choice among the available anti-dermatophyte agents as well as to optimize the therapy against dermatophytosis.

Objectives
we aimed the antifungal susceptibility of isolated dermatophytes from Iran against some common antifungal agents for the treatment of dermatophytosis.

Methods
The isolates were identified by traditional macroscopic and microscopic findings as well as PCR-RFLP. A total of 316 clinical isolates from patients with dermatophytosis including Trichophyton interdigitale (n = 156), T. rubrum (n = 60), Epidermophyton floccosum (n = 42), Microsporum canis (n = 29), Arthroderma benhamiae (n = 17) and T. tonsurans (n = 12) were tested for antifungal activities against terbinafine, itraconazole, griseofulvin and fluconazole by broth microdilution technique according to the recommendation of the Clinical and Laboratory Standards Institute (CLSI) M38-A2 document.

Conclusions
The MIC ranges were as follows: terbinafine, 0.002 to 0.25 μg/mL; itraconazole, 0.004 to 0.5 μg/mL; griseofulvin, 0.125 to 8 μg/mL and fluconazole 4 to 128 μg/mL. Terbinafine and fluconazole had the highest and lowest activity against dermatophytes isolates, respectively. Epidermophyton floccosum, Microsporum canis and Trichophyton tonsurans were the most susceptible species to terbinafine. Terbinafine had the lowest activity against Arthroderma benhamiae. E. floccosum showed the highest susceptibility to itraconazole. T. rubrum had the lowest
susceptibility to itraconazole. According to our data we profoundly propose to physicians to exclude fluconazole for treatment of dermatophytosis.
FEMS-2959
Antimicrobial resistance

SRL PATHOGENICITY ISLAND IS ASSOCIATED TO MULTIDRUG RESISTANCE IN SHIGELLA STRAINS ISOLATED FROM CHILE

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Background
Shigella spp., the etiologic agent of shigellosis, becomes a major problem of public health due to the increasing multidrug-resistance to antibiotic (MDR). In this context, the Shigella resistance locus (SRL) pathogenicity island (PAI) mediates resistance to streptomycin, ampicillin, chloramphenicol, and tetracycline, contributing to MDR phenotype.

Objectives
To characterize the presence and genetic organization of SRL PAI in Chilean Shigella spp. strains.

Methods
We selected 135 strains of S. sonnei and S. flexneri. We determined susceptibility to antibiotics using agar diffusion method, and detected seven specific SRL markers (orf9, orf58, oxa-1, tet, cat, and its sites of insertion in the chromosome) by PCR. Also, we assessed the presence of integrons. Then, we selected representative strains and analyzed them by Tilling-PCR for the complete island organization.

Conclusions
Our results indicate that most of these strains are MDR strains to ≥ 3 antibiotic class (> 78%), all of them had the SRL PAI. Besides, a high rate of the isolated was positive for class 1 integron (80%), which is concordant with the presence of the SRL PAI and negative for integron class 2.

Meanwhile, tilling-PCR for SRL PAI and sequencing indicated that in the 5’ end there are four variants with different insertions. Only one insertion was found in the 3’ end, compared to the reference strain, YSH6000. In conclusion, this work points out the diversity of variants of the SRL PAI in the Chilean Shigella spp strains, and its huge contribution to the MDR phenotype.

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MOLECULAR EPIDEMIOLOGY OF PLASMIDS MEDIATING OXA-48 CARBAPENEMASE DISSEMINATION

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Background

OXA-48 carbapenemase has become of major significance as a result of the progressive identification of nosocomial Klebsiella pneumoniae producing this enzyme. It has been reported mostly in Mediterranean countries, although it is increasingly detected in Northern Europe. The recent spread of a clonal OXA-48-producing K. pneumoniae as well as other Enterobacteriaceae in a Dutch hospital has resulted in one of the largest OXA-48 related outbreaks in Europe so far¹.

Objectives

The OXA-48 gene has been associated with 60- to 70-kb plasmids. Our aim was to perform a large-scale study using a next-generation sequencing approach to better understand the molecular epidemiology of plasmids carrying blaOXA-48.

Methods

A total of 89 OXA-48-producing bacteria isolated from the Dutch hospital outbreak¹ as well as from other locations in the Netherlands and endemic countries were selected. Plasmids were sequenced using the Illumina Miseq platform, and read sets were assembled de novo using Spades. Plasmid sequences were annotated with Prokka, and Samtools was used to identify SNPs and INDELs.

Conclusions

Sequencing data revealed 3 plasmid clusters of ca. 62kb, 30kb, and 13kb. All the plasmids shared an identical common region including the Tn1999 where blaOXA-48 is embedded, suggesting the presence of a highly modular plasmid ancestor. No link was found between plasmid clusters and origin of the strains. However, our results show several cases of plasmid transfer between bacterial species in the same patient over time, which represents a major driver for plasmid maintenance and spread despite control of the outbreak.
References

ANTIMICROBIAL ACTIVITY OF CHLORELLA VULGARIS EXTRACTS AGAINST FOODBORNE PATHOGENIC BACTERIA

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Background

Foodborne bacterial pathogens cause a major public health problem. In the last few decades, increased resistance of bacterial strains to antibiotics lead to search alternative effective compound against food borne pathogens.

Objectives

In this study was aimed at evaluating the antimicrobial activity of different extracts from Chlorella vulgaris on food borne pathogens. Systematic screening of them may result in the discovery of novel effective compounds.

Methods

Chlorella vulgaris (UTEX 2714) was grown in axenic cultures of 3N-Basal Bold Medium. Chloroform, methanol, ethanol, diethyl ether, acetone and ethyl acetate extracts were extracted directly from 5 mg dried C.vulgaris biomass. Staphylococcus aureus, Listeria grayii, L.monocytopogenes, L. innocua, Enterococcus fecalis and E. faecium strains was isolated from cheese, minced meat and poultry product and identified by molecular techniques. Disc diffusion methods were used to determine the antimicrobial activity of C. vulgaris extracts on the foodborne pathogens. Results compared with streptomycin and amikacin results which was used as positive control.

Conclusions

C.vulgaris extracts possessed potential antibacterial activities against foodborne gram positive bacterial strains with inhibition zones ranged for S.aureus, E. fecalis and E.faecium 7-14 mm, L.monocytopogenes 7-12 mm, L.grayi 7-14 mm, L.innocua 7-15 mm, Thus, it could be suggested that the C.vulgaris is useful biosystem for production of new antimicrobial compound that has an effect against gram positive bacteria.
Background
Carbapenemases are able to hydrolyze carbapenems and, in several instances, almost all other β-lactam antibiotics, making these enzymes a great threat to our ability to treat refractory bacterial infections.

Objectives
In this study, we aimed to determine the genetic context of carbapenemase genes and any additional acquired resistance genes in clinical isolates. We also wanted to study the potential of using whole genome sequencing (WGS) in predicting the antibiotic resistance profiles of multi-resistant isolates.

Methods
Six clinical carbapenemase-positive isolates of five different species (Acinetobacter baumannii, Escherichia coli, Klebsiella pneumoniae, Proteus mirabilis, Pseudomonas aeruginosa) were selected at the Sahlgrenska university hospital in Gothenburg, Sweden. WGS of selected isolates resulted in draft assemblies that were annotated for mobile antibiotic resistance genes and chromosomal mutations in antibiotic target genes.

Conclusions
We found that several different resistance genes and mutations had been acquired by the isolates. The detected carbapenemase genes included variants blaNDM, blaOXA-24, blaOXA-48, and blaOXA-51, and some isolates carried multiple variants. From the acquired resistance mechanisms and the intrinsic resistance profile of each species, we could predict a susceptibility profile. The phenotypic resistance profile was determined by measuring the MIC for 37 different antibiotics belonging to all major classes, using E-tests. The results showed good cohesion between the predicted phenotype and measured resistance profile. Our results suggest that WGS can be used to predict the resistance phenotype of carbapenem-resistant isolates. The analysis also underlines the importance of using a comprehensive and curated database of resistance factors for accurate identification the resistance genotype.
Background

Our work focused on ISCR, a family of insertions sequences encoding RCR transposases thought to catalyze the a rolling-circle (RC) mechanism transposition of their IS elements. The prevailing ISCR, termed ISCR1, are associated to variable combinations of antibiotics resistance genes. The recent discovery of free forms of ISCR1 elements suggests their contribution to the dissemination of these genes.

Objectives

Nothing is known about the expression of the RCR1-encoding gene rcr1. This gene displays a conserved s^70 promoter and a putative binding box (operator) for the LexA protein, the regulator of the SOS response. We therefore investigated the SOS induction of the expression of rcr1.

Methods

Translational fusions Prcr1-lacZ with a wild type or a mutated LexA box were constructed. The activity of the Prcr1 promoter was estimated using b-galactosidase assays performed in an E.coli ΔlacZ strain or its derivative ΔlexA, after treatment with an SOS-inducer or not.

Conclusions

An activity from Prcr1-lacZ could be detected either in the DlexA strain or upon exposure to mitomycin C unless in strains deficient for the SOS response (ΔrecA). RNA techniques allowed us to map the Prcr1 promoter that was also supported by mutation of the deduced -10 element.

Bioinformatics analyses suggested LexA could regulate more RCR transposases genes and we confirmed it with a recently identified transposase.
Our results highlight a SOS induction of \textit{rcr} genes, suggesting that the mobility of their cognate mobile elements could be triggered upon exposure to some antibiotics or single-strand generating horizontal gene transfer events.
EPIDEMIOLOGIC CHARACTERIZATION OF NDM-1-PRODUCING ESCHERICHIA COLI AND KLEBSIELLA PNEUMONIAE ISOLATES IN TURKISH HOSPITALS

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Background
Enterobacteriaceae isolates producing New Delhi metallo-beta-lactamase-1 (NDM-1) are considered as a major health threat. We investigated the epidemiology of NDM-1-producing isolates among carbapenem-resistant Escherichia coli and Klebsiella pneumoniae isolates submitted to a central laboratory from various Turkish hospitals.

Objectives
NDM-1 is a class B metallo-beta-lactamase conferring high-level resistance to beta-lactam antimicrobials including carbapenems and is transferable through plasmids. blaNDM-1 positive Enterobacteriaceae are being reported from Turkish hospitals but the epidemiological characterization of these isolates is lacking. This study investigates the prevalence and clonal relationship of blaNDM-1 positive isolates among carbapenem-resistant E. coli and K. pneumoniae collected from six major Turkish cities (Istanbul, Ankara, Bursa, Adana, Eskisehir, Kocaeli) between September 2011-March 2014.

Methods
We detected 143 carbapenem-resistant isolates (E. coli, n=28; K. pneumoniae, n=115) and a multiplex PCR assay targeting blaOXA-48, blaNDM-1, blaIMP, blaVIM and blaKPC genes was performed for these isolates. blaNDM-1 was identified in 24 (16.7%) isolates (E. coli, n=4; K. pneumoniae, n=20). Pulsed-field gel electrophoresis (PFGE) of XbaI-digested genomic DNA was used to assess the clonal relationship among blaNDM-1 positive isolates which revealed 3 and 8 distinct pulsotypes among E. coli and K. pneumoniae isolates, respectively (Figure 1).
Conclusions

bla_{NDM-1} was detected in 14.3% and 17.4% of carbapenem-resistant E. coli and K. pneumoniae isolates, respectively. The different patterns obtained by the PFGE showed that the infections with NDM-1-producing K. pneumoniae are due to intra-institutional spread rather than transfer between institutions.

Acknowledgement: This work was supported by TUBITAK for the project entitled ‘Observable Real-Time Electrophoresis” (Project No: 113M354).
Background

*Hyoscyamus* species contain important tropane alkaloids, such as Hyoscyamine, scopolamine and to lesser extend atropine which have antispasmodic, anticholinergic, analgesic and sedative effects. These compounds are produced mainly by *Hyoscyamus albus* L. This shrub is able to accumulated high amount of tropane alkaloids and these alkaloids indicated antimicrobial activity to microorganisms.

Objectives

In this study obtained extracts of some plant species which belongs to *Hyoscyamus* (*Hyoscyamus reticulatus* L., *Hyoscyamus leptocalyx* STAPF, *Hyoscyamus aureus* L., *Hyoscyamus albus* L.) (Solanaceae) family by various solvent systems were researched within the context of their antimicrobial activities.

Methods

Antimicrobial activities of hexane, ethylacetate and methanol extracts of *Hyoscyamus* species were tested by disc diffusion method. Gram positive and gram negative standard bacteria strains, clinical isolates and *Candida albicans* (ATCC-10231) yeasts over amoxicillin/clavulanic acid (2:1) ofloxacin, nystatin, imipenem and netilmicin standart antibiotics and 150 µg/6 mm paper disc plant extracts tested.

Conclusions

The antimicrobial activity of hexane, ethylacetate and methanol extracts of *Hyoscyamus* species showed different antibacterial activity against tested microorganisms. The diameters of growth inhibition zones of *Hyoscyamus* species ranged from 8 mm to 12 mm. Among all the *Hyoscyamus* extracts studied, methanol extract of *H. reticulatus* and *H. aureus* demonstrated the highest antimicrobial activity versus *S. aureus* (clinic-isolate). *S. aureus* (ATCC-25923), *S. aureus* (clinic-isolate),
and *S. pyogenes* (ATTC-19615) were sensitive to all *Hyoscyamus* extracts except *H. leptocalyx*. 
FEMS-2344
Antimicrobial resistance

ANTIMICROBIAL RESISTANCE- AND CONSUMPTION SURVEILLANCE – A 12-YEAR HOSPITAL EXPERIENCE
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Background
During the last years surveillance of Antibiotic resistance (AR) and Antibiotic consumption (AC) became a part of the efforts to contain AR.

Objectives
Our aim was to analyze the results of 2002 – 2014 AR and AC surveillance, conducted at a multiprofile tertiary care 300-bed hospital in Sofia.

Methods

Etiology of infections and AR were monitored as a part of the national surveillance program BulSTAR. AC was calculated by ABC calc (D. Monnet).

Results. A total of 44786 clinically significant bacteria have been isolated. The first 3 top isolates from urine were Escherichia coli (Eco), other Enterobacteriaceae (oE-ae), Enterococcus spp (Enc); from wounds - Staphylococcus aureus (Stau), Eco, Enc; from bacteremia - Eco and oE-ae, Stau and Enc; from the respiratory tract – Streptococcus pneumoniae (Spn) and Haemophilus influenzae (Hin). The rate of MRSA from wounds was 11 %, but from bacteremia 50 % to 78 %. The most problematic organisms were E-ae with ESBL: they increased from 3.8 % in 2002 through 12 % in 2007 till 26.05 % in 2014. Among Enc there were not VRE; among Spn – the rate of PNSSP was 16.4 %; 10 % of Hin were bla (+). The results from AC showed a fluctuation from 27 to 54 DDD/100 bed-day, with a significant higher AC in the surgical ICU (~ 250 DDD/100 bed-day).

Conclusions
Our results illustrated an increase of antimicrobial resistance (ESBL producing Enterobacteriaceae) despite the antibiotic policy attempts and suggest for more active actions both nationally and globally.
AMINO ACID SUBSTITUTIONS IN GYRA AFFECTS QUINOLONE SUSCEPTIBILITY IN SALMONELLA TYPHIMURIUM

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Background
Amino acid substitutions have generally been found within the quinolone resistance determining regions in subunit A of DNA gyrase (GyrA) of Salmonella Typhimurium, but direct evidence of the contribution of these substitutions to quinolone resistance remains to be shown.

Objectives
To clarify the significance of amino acid substitutions in S. Typhimurium GyrA to quinolone resistance, we expressed recombinant wild-type (WT) and S83F, D87N, D87G, D87Y and S83F-D87N mutant gyrases in Escherichia coli and characterised them in vitro.

Methods

WT and mutant gyrases were reconstituted in vitro by mixing recombinant subunits A and B. Correlation between the amino acid substitutions and resistance to quinolones ciprofloxacin, levofloxacin, nalidixic acid and sitafloxacin was assessed by the quinolone-inhibited supercoiling assay.

Conclusions
We concluded that amino acid substitutions at position 83 and 87 of GyrA, and quinolone structural variations at position 8 influenced the interaction between DNA gyrase and drugs. Furthermore, we suggest that sitafloxacin might be a choice for the treatment of salmonellosis caused by ciprofloxacin-resistant S. Typhimurium.
Background

*Klebsiella pneumoniae* (KP) is an opportunistic pathogen of *Enterobacteriaceae* family and is resistant to many antibiotics, especially the beta-lactams, due to extensive production of beta-lactamases. SHV-14 is one of the potent extended-spectrum serine beta-lactamases present chromosomally in *K. pneumoniae*.

Objectives

Physiological characterization of the membrane bound form of *bla*SHV-14 in *K. pneumoniae* and study its effect on antibiotic resistance and biofilm formation.

Methods

Membrane bound form of *bla*SHV14 gene was cloned in pBAD-18 Cam vector and transformed in *ampC* deleted CS109 *Escherichia coli* strain for *in vivo* studies. Minimum Inhibitory Concentration (MIC) was determined in accordance withClinical and Laboratory Standards Institute (CLSI) using the micro-broth dilution technique. Beta-lactamase activity of cloned *bla*SHV14 was checked by monitoring hydrolysis of a chromogenic substrate nitrocefin at particular time interval in *E. coli* CS109 cell lysate. Biofilm formation on polystyrene surface was also studied by staining the biofilms formed with 0.1% crystal violet stain. An elevated level of beta-lactamase hydrolytic activity was observed against penicillin and cephalosporin group of antibiotics in the presence of *bla*SHV14 in *ampC* deleted CS109. Bet-lactam resistance was increased by eight fold against cephalosporin and 14 fold against penicillin group of antibiotics as compared to the control lacking *bla*SHV14 gene. Biofilm formation index of *ampC* deleted CS109 harbouring *bla*SHV14 gene showed three fold increments as compared to control devoid of *bla*SHV14 beta-lactamase.

Conclusions

Therefore, the presence of *bla*SHV14 gene in *Klebsiella pneumoniae* makes the cell resistant to beta-lactam group of antibiotics and subsequently enhances biofilm formation.
ESCHERICHIA COLI FROM BLUE MUSSELS (MYTILUS EDULIS) AS INDICATORS FOR ANTIBIOTIC RESISTANCE

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Background

Antibiotic resistance in bacteria is an increasing challenge in all situations where such agents are applied. Blue mussels (Mytilus edulis) are suspension feeders and may retain particles with close to 100% efficiency. An adult blue mussel (60 mm in shell length) may clear between 12 and 240 litre of seawater per day for particles containing bacteria. According to the current EU regulations (EU Directive 854/2004), farm sites for cultivation of bivalves must be classified according to their suitability in terms of microbiological and chemical water quality. One of the parameters to be examined is the concentration of Escherichia coli in the soft parts and mantle water of the bivalves.

Objectives

E. coli derived from animals or man by sewage or land runoff, may subsequently be filtered out by Blue mussels and can be used as indicators for the antibiotic resistance status and detect spatial and temporal changes in resistance prevalence.

Methods

The Norwegian Food Safety Authority submit annually approximately 300 samples of Blue mussels from over 35 stations along the coast. The number of E. coli in these samples are determined by a multiple tube dilution method (Donovan’s method, ISO 16649-3). On average 60% of the examined samples harbour E. coli, and strains are currently being isolated, and will be characterised.

Conclusions

Isolation and characterisation of the antibiotic resistance profile of E. coli from Blue mussels may represent a potential indicator of the long term changes.
Background
Due to the rising incidence of carbapenem-resistant Enterobacteriaceae (CRE) in the UK and the increased mortality associated with such infections, it is imperative that adequate infection control measures are in place to limit their spread in hospitals. Despite targeted screening of patients at increased risk of carriage, the overall burden may be underestimated. Thus it is recommended that healthcare facilities with CRE conduct surveillance studies to establish the true burden of infection.

Objectives
Surveillance was performed using diarrhoeal stools from in-patients at a large London teaching hospital to assess whether current screening strategies adequately identify carriers, particularly those posing an increased risk of dissemination.

Methods
In addition to routine investigations, diarrhoea stool samples from in-patients were cultured onto Brilliance CRE medium. Blue or pink pigmented colonies that developed overnight were sub-cultured onto MacConkey agar with an ertapenem disc. Ertapenem-resistant colonies were identified by VITEK 2. Enterobacteriaceae with minimum inhibitory concentration values above the screening threshold for meropenem were referred to the national reference laboratory for further testing.

Conclusions
Over seven months, 1013 stool samples from 899 patients were screened. Thirteen colonised patients were identified; a carriage rate of 1.45%. Two carbapenemase producing Enterobacteriaceae (CPE) and 11 extended spectrum or AmpC β-lactamase producers with porin loss were isolated. Screening of diarrhoea stools detected a higher proportion of carbapenemase non-producing CRE than routine surveillance swabs (11/889 vs. 4/8663; P=0.001). CPEs remain rare in acute healthcare settings, even among patient populations with increased risk of transmission. Current guidance for CRE surveillance is considered adequate.
Background
Drug resistance mechanisms of bacteria are a menace to antibiotic therapy. Antibiotic inactivating enzymes and biofilm formation are prominent bacterial escape routes. The largest group of drug hydrolysing enzymes are beta-lactamases which may influence biofilm formation.

Objectives
Inducible beta-lactamase expression is regulated by the same genes that govern peptidoglycan recycling. We attempt to elucidate the role of these genes in biofilm formation by the beta-lactamase producing *E. coli*.

Methods

Single and double gene deletion mutants of *ampC, ampD, ampE* and *ampG* in *E. coli* 25113 were created by the one-step gene inactivation method. Serine beta-lactamases (TEM-1, OXA-2, CTX-M-15 and SHV-14) from the nosocomial enterobacterial isolates were cloned and expressed in the parent and mutant strains. The influence of deletions on beta-lactamase expression and subsequent effect on biofilm formation were studied by chromogenic cephalosporin hydrolysis and crystal violet staining, respectively. The drug resistance alteration was enumerated by standard microbroth dilution techniques. It was observed that serine beta-lactamases like TEM-1 and OXA-2 lowered the biofilm formation whereas ESBLs like CTX-M-15 and SHV-14 facilitated it. Deletion of *ampD* showed varied levels of beta-lactamase expression and similar effect was observed in biofilm formation. Nonetheless, overexpression of beta-lactamases in Δ*ampE* enhanced biofilm formation by 45% whereas a biofilm reduction of 29% was observed in Δ*ampG* mutant as compared to beta-lactamase deficient strain.

Conclusions
We infer that beta-lactamase expression not only enhances beta-lactam resistance but also influences biofilm formation; and the genes *ampD, ampE* and *ampG* affect the expression of Class A and D beta-lactamases in *E. coli*. 
First report of the presence of oxacillinases and metallo-β-lactamases in invasive isolates of carbapenem-resistant Acinetobacter baumannii in Vojvodina Province, Serbia

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Background
Acinetobacter spp. are the most common invasive isolates from blood in Vojvodina Province. The importance of these microorganisms is an exceptional ability of survival and spread in the hospital environment, as well as the extraordinary capacity to quickly acquire different mechanisms of resistance to a wide variety of antimicrobial agents, including carbapenems.

Objectives
To detect the presence of encoded genes for oxacillinases and metallo-β-lactamases in carbapenem resistant isolates of Acinetobacter baumannii in Vojvodina Province.

Methods
The identification to the species level of Acinetobacter baumannii was confirmed by PCR detection of blaOXA-51-like. In 50 carbapenem-resistant Acinetobacter baumannii isolates, detection of blaOXA and blaMBL gene was conducted by multiplex PCR, as well as the detection of insertion sequences ISAba1.

Conclusions
blaOXA-24-like was detected in 45 carbapenem-resistant Acinetobacter baumannii isolates. Twelve isolates were positive for blaOXA-58-like. The presence of blaOXA-23-like gene was not detected in any of the isolates. A simultaneous presence of two genes was detected in 43 isolates. Thirty eight isolates were positive for blaOXA-24-like and blaOXA-51-like genes and in 5 isolates blaOXA-58-like and blaOXA-51-like genes were detected. Seven isolates had three genes, blaOXA-24-like, blaOXA-51-like and blaOXA-58-like. All isolates were PCR positive for ISAba1. The presence of blaIMP and blaVIM was not detected in any of the isolates. The resistance to carbapenems in isolates of Acinetobacter baumannii in Vojvodina Province is caused by production of OXA-24 and OXA-58 oxacillinases, because metallo-β-lactamases were not detected.
COMPARATIVE EFFICACY OF THREE CHLORINE-BASED DISINFECTANTS TO CONTROL PLANKTONIC AND SESSILE ESCHERICHIA COLI

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Background

Sodium hypochlorite (SH) is the most common antimicrobial agent used in food industry even if it produces unhealthy and toxic by-products. New chlorine-based decontamination strategies are emerging for the decontamination of fresh products and disinfection of surfaces, such as neutral electrolyzed oxidizing water (NEOW) and chlorine dioxide (CD).

Objectives

The aim of the present work was to assess the effects of three chlorine-based disinfectants (SH, NEOW and CD) on the control of Escherichia coli in both planktonic and sessile states. The stability of the disinfectants was also assessed over time.

Methods

Four different concentrations (20, 50, 80 and 100 ppm) were tested on the control of planktonic and sessile E. coli cells. Planktonic tests were made by measuring growth inhibition over time. Biofilms were formed on stainless steel surfaces and the efficacy of chlorine-based products was assessed by colony forming units counting. The free chlorine content was also measured over time for each disinfectant tested.

Conclusions

NEOW was the most efficient disinfectant in the control of both planktonic cells and biofilms while SH was the least effective. In terms of free chlorine content in solution/disinfectant stability, SH was the most instable and NEOW was the most stable. In conclusion, NEOW is an effective disinfectant strategy for the control of both planktonic and sessile cells and disinfectant decay was negligible compared to the other disinfectants.
Background
Pseudomonas aeruginosa is a Gram-negative bacterium, commonly present in the environment, that causes serious health problems when it infects humans and is resistant to many antibiotics and treatments. Patient groups at risk for acquisition of P. aeruginosa infections include those with cystic fibrosis (which almost always leads to chronic airway infection), paraplegic, burn and immune systems deficiencies patients. Prevention of infection by vaccination is desirable until today no vaccine has yet obtained market authorization.

Objectives
This project is aimed at the development of the first peptide-based P. aeruginosa vaccine and to test planned techniques. Our goal is to produce high immunogenic response in human body.

Methods
We have selected short peptides from P. auruginosa surface localized proteins using bioinformatic tools. Now we are using genetic engineering and cloning to put those peptides into stable protein scaffolds. After stability tests we will try to induce response in animal model and obtain antibodies for further testing with serum samples from cystic fibrosis patients and Pseudomonas aeruginosa clinical isolates. To evaluate whether the specific antibodies are protective in vitro, we will use opsonophagocytosis assays.

Finally, we will perform in vivo infection experiments in animal models after immunization with targets to reveal whether the produces targets are protective against pathogen.

Conclusions
As this is rather new development method, we are facing problems and we are trying to establish protocols for our purpose, nevertheless project is going forward and we are expecting successful results soon.
ANTIMICROBIAL ACTIVITY OF NEWLY SYNTHESIZED Ni(II) COMPLEXES ON LABORATORY STRAINS OF MICROORGANISMS AND CLINICAL ISOLATES

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Background
The escalation of multidrug-resistant (MDR) bacteria is documented to be a serious problem that affects the choice of appropriate antibiotic therapy and increases the probability of unfavorable infection outcome. Resistance has emerged towards all classes of antibiotics leading to a continuous need for producing new drugs. One of the promising methods in coping with bacterial resistance is the use of alternative classes of antimicrobial agents.

Objectives
The aim of this study was to investigate the antimicrobial activity of seven newly synthesized Ni(II) complexes.

Methods
Minimum inhibitory concentrations (MICs) of complexes were determined by broth microdilution method according to Clinical and Laboratory Standards Institute guidelines (CLSI, 2007). The antimicrobial activity was investigated on eight laboratory control strains of microorganisms: Staphylococcus aureus ATCC 25923, Staphylococcus epidermidis ATCC 1228, Bacillus subtilis ATCC6633, Escherichia coli ATCC 25922, Klebsiella pneumoniae ATCC 13883, Pseudomonas aeruginosa ATCC 27853, Salmonella abony NCTC 6017, yeast Candida albicans ATCC 10231, and twenty-five clinical isolates of Escherichia coli and Pseudomonas aeruginosa. All tests were performed in Müller Hinton broth for the bacterial strains and in Sabouraud dextrose broth for the yeast. Complexes were dissolved in 1% dimethylsulfoxide (DMSO) and then diluted to the desired concentrations ranging from 31–1000 micrograms/ml with addition of 0.05% triphenyl tetrazolium chloride as a growth indicator.

Conclusions
Two of tested compounds exhibited strong activity against Gram-positive bacteria, moderate activity against standard strains of Gram-negative bacteria and weak activity against clinical isolates of Escherichia coli.
ELUCIDATING THE ROLE OF UPC2 AND NDT80 TRANSCRIPTION FACTORS IN CANDIDA PARAPSILOSIS AZOLE RESISTANCE

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Background

C. parapsilosis is responsible for hospital-acquired infections with remarkable risks especially to neonates and immunocompromised patients. In the past decades, this human pathogen has been ranked as second or third most isolated fungus, depending on the geographical region. Since its incidence is growing, the number of C. parapsilosis resistant to azoles has been followed the same trend. Despite C. parapsilosis high prevalence, molecular mechanisms associated to azole resistance remain poorly studied. In a pioneer study addressing azole resistance, an overexpression of ergosterol biosynthesis genes, namely ERG11, and their regulators, UPC2 and NDT80 genes, were identified as main players in posaconazole resistance.

Objectives

The aim of this work was to reveal the importance of UPC2 and NDT80 transcription factors in C. parapsilosis posaconazole resistance, through control of ergosterol biosynthesis.

Methods

For that, both genes were independently and simultaneously knockout in posaconazole resistant strain BC014R_psc, using SAT1-Flipper cassette. Thus, upc2Δ, ndt80Δ and upc2Δ/ndt80Δ mutants were generated. These strains were characterized regarding their azole susceptibility profile.

Conclusions

Deleted strains became susceptible to posaconazole, voriconazole and fluconazole. Minimal Inhibitory Concentrations for azoles in upc2Δ and upc2Δ/ndt80Δ mutants were similar and inferior to those obtained in ndt80Δ strain. We conclude that UPC2
and *NDT80* genes are both involved in *C. parapsilosis*azole resistance, however the first seems to play a major role.

This study was supported by FEDER, COMPETE and FCT (PTDC/DTP-EPI/1660/2012).
Background
Brucellosis is an important health problem in our country in aspect of long lasting treatment and tendency to relapse. Getting information about the pattern of brucella antibiotic susceptibility help us to treat patients more successfully.

Objectives
This study was done to assess antibiotic susceptibility pattern of brucella with E-test method in Kashan region.

Methods
This descriptive study was performed on 48 clinical samples were positive culture for brucella in Kashan shahid Beheshti hospital in 2013. Clinical samples got from patients suspicious to brucellosis and Minimum inhibitory concentrations was calculated for ciprofloxacin, doxycycline, rifamin, azithromycin, ceftriaxone, streptomycin and cotrimoxazol with E-test. For each sample a form contain of information about age, sex, clinical manifestation and E-test results was filled. All informations were entered in SPSS 16 software. Results were shown as descriptive statistics.

Conclusions
All brucella isolates were sensitive to antibiotics, without any report of resistance. Cotrimoxazol with MIC50:0.016, MIC90:0.064 was the most effective choice. Doxycycline the second one, streptomycin had the highest MIC. There was not any statistical relation between patient age, sex, type of reception and antibiotic susceptibility.

Cotrimoxazol and doxycycline as effective drugs on brucella are the best choice for treatment of brucellosis in our region. Streptomycin should be used cautiously because it had the highest MIC against brucella spp in our study.
Background

*Staphylococcus aureus* is an opportunistic pathogen that colonizes the anterior nares and skin, being one of the most common causes of skin, soft tissue and nosocomial infections.

Objectives

This study was undertaken to evaluate *S. aureus* carriage in public health students at the Faculty of Pharmacy (Granada, Spain), and phenotypical resistance of the strains to antibiotics.

Methods

This study was performed on 144 public health students. Sampling was done once between October 2013 and January 2014. Samples were collected from both nares inserting a sterile nasal swab into nostril. The swabs were cultured on Mannitol Salt Agar and incubated at 37ºC for 24h. Bacterial identification followed the standard phenotypic methods. Antibiotic susceptibility of isolates was tested by disc diffusion method (EUCAST, 2013). A wide array of antibiotics representing the main groups: penicillins, cephalosporins, aminoglycosides and others.

Conclusions

*S. aureus* was isolated from 20% (29/144) of the student population. Regarding the penicillin group, 3 of 29 strains did not show resistance to this group. 82.7% of *S. aureus* positive strains demonstrated multiresistances to penicillins and cephalosporins. Among the group of aminoglycosides, 93.1% of strains were multiresistant to them, and 2 strains were found to be sensitive for all antibiotics in this group. Furthermore, all strains of this study proved resistances to nalidixic acid. We may conclude that the strains studied exhibit a broad spectrum of multiresistances, which suggests that the use of some penicillins, cephalosporins and aminoglycosides antibiotics could be ineffective for the treatment of infections caused by *S. aureus*. 
ANTIMICROBIAL RESISTANCE AND MOLECULAR CHARACTERIZATION OF AVIAN PATHOGENIC ESCHERICHIA COLI IN BELGIUM.

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Background
Avian pathogenic Escherichia coli (APEC) causes colibacillosis and significant economic losses in the poultry industry.

Objectives
To isolate, perform antimicrobial susceptibility testing, serotyping, phylogrouping and molecular characterization of the APEC isolates from Belgium.

Methods
Antimicrobial susceptibility was tested by micro-broth dilution (TREK diagnostics sensititre plates) using the European Committee on Antimicrobial Susceptibility Testing (EUCAST) breakpoints. Multi drug resistance (MDR) was described as resistance to more than three classes of antimicrobials. O-antigen serotyping was done using 31 different antisera (LREC, Spain). Strains negative for all these serotypes were considered not typeable (NT). The isolates were grouped into A, B1, B2 and D phylo-groups. The microarray (CLONDIAG GmbH, Jena, Germany) system was used to test for virulence and antimicrobial resistance genes.

Conclusions
The 212 APEC isolates were resistant to ampicillin (61%), ciprofloxacin (58%) and tetracycline (49%). MDR was observed in 135 (64%) of the isolates. 13 (6%) isolates have been presumptively identified as ESBL producing. O-antigen grouping showed that 73 (34%) isolates collectively belonged to sero-group O78, O2, and O1; 65 (31%) to other sero-groups associated with colibacillosis in poultry, and 74 (35%) were NT. The most prevalent phylo-groups were D (42%) and A (40%); 15% were phylo-group B1, while 3% were B2. Genes coding for adhesion factors fimH (75%) and Iron acquisition systems iroC (57%), irp2 (52%), were mostly observed.
Antimicrobial resistance is high and a large percentage of the isolates are MDR. APEC strains showed high prevalence of virulence factors that might be associated with greater pathogenic potential.
FEMS-0992
Antimicrobial resistance

CLASS-I AND CLASS-II INTEGRONS IN MULTIDRUG RESISTANT ESBL AND CARBAPENAMASE PRODUCING ENTEROBACTERIACEAE FROM EGYPT
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Background

Escherichia coli and Klebsiella pneumoniae with Extended-Spectrum-β-Lactamase-(ESBL) and carbapenemase-mediated resistance are increasingly reported in hospital and community settings worldwide. ESBL CTX-M-15 and Class-D carbapenemase OXA-48 co-production, plus resistance to other antibiotics, limit treatment options, while integrons support resistance genes expression and movement.

Objectives

To characterize genetic determinants encoding β-lactam resistance in Enterobacteriaceae isolates collected from Egypt, and investigate their association with class-I and class-II integrons.

Methods

One hundred and sixty-nine ESBL-producing E. coli and K. pneumoniae from nosocomial and community-acquired infections were collected from 10 Egyptian university hospitals in two surveillance studies (Group-1: 65 K. pneumoniae 2000-2003, Group-2: 41 E. coli, 63 K. pneumoniae 2009-2011). Resistance genes and integron integrases were identified by PCR and DNA sequencing. Plasmid located blaCTX-M-15 was identified in 11% (n=7) of isolates from group-1 vs. 97% (n=101) from group-2. Non-metallo-carbapenamase blaOXA-48 was detected in 88% (n=57) of isolates from group-1 vs. 95% (n=99) from group-2. Plasmids displayed a range of different RFLP patterns. Integrase-1 was detected in 83% (n=54) of isolates from group-1 vs. 93% (n=97) from group-2, while Integrase-2 was identified in 97% (n=63) of isolates from group-1 isolates vs. 89% (n=93) from group-2.

Conclusions
We have shown that association between integrases and resistance genes carriage appear to increase over time. The linkage between integrases and blaCTX-M-15, and blaOXA-48 is currently being determined by investigating the first gene cassettes in integrons.
FIRST DESCRIPTION OF A FOSFOMYCIN RESISTANT CLINICAL ESCHERICHIA COLI ISOLATE HARBOURING PLASMID-BORNE FOSA3 AND BLACTX-M-15 IN EUROPE

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Background
The widespread of extended-spectrum beta-lactamase (ESBL)-producing Enterobacteriaceae has reigned the interest in old antibiotics, being fosfomycin one of the first therapeutic options for treatment of community acquired urinary tract infections. Resistance to fosfomycin is mostly linked to chromosomal mutations causing reduced fosfomycin uptake, though acquisition of plasmid-mediated fosfomycin resistance genes (fos), mainly fosA3 in isolates from Asia and non-clinical settings has been reported.

Objectives
To assess the frequency of plasmid-mediated fosfomycin resistance genes and their genetic background among fosfomycin resistant Enterobacteriaceae isolates from a community laboratory from Portugal.

Methods
Between August 2012-September 2013, 58 fosfomycin resistant isolates (29 K.pneumoniae, 26 E.coli, 2 Proteus mirabilis and 1 Morganella morgani) were identified among 461 ESBL-producing Enterobacteriaceae. Antibiotic susceptibility testing was performed by standard methods, including fosfomycin disks supplemented with glucose-6-phosphate. fos (fosA, fosA3, fosB, fosC2), blaESBL and rmtB genes were screened by PCR and sequencing. E. coli phylogroups and genetic context were investigated by PCR and PCR mapping and sequencing, respectively. Plasmid typing was assessed by PBRT, IncFII typing (FAB formula), I-Ceu-PFGE and hybridization.

Conclusions
Although with a low prevalence, we identified in the urine of a 61 year old patient, one E. coli (1.7%) isolate of phylogenetic group D1, from carrying the fosA3 gene together with blaCTX-M-15 in an F2:A6:B- IncFII plasmid, which corresponds to the first description of fosA3 in Europe. Although aac-lb-cr, rmtB and blaOXA-1 genes were not detected, the fosA3 gene was located in an IS26-composite transposon, similar to others previously reported in Asia suggesting a common platform.
Antimicrobial resistance

IN VITRO ACTIVITY OF AQUEOUS AND METHANOLIC EXTRACTS OF PSIDIUM GUAJAVA, NAUCLEA LATIFOLIA AND BROPHYLLUM PINNATUM AGAINST BACTERIA ISOLATED FROM CHICKEN MEAT

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Background
Medical knowledge derived from traditional societies has motivated searches for new bioactive molecules derived from plants that show potent activity against bacterial pathogens.

Objectives
The present study investigated the antibacterial activity of Psidium guajava, Nauclea latifolia and Brophyllum pinnatum on Staphylococcus aureus, Escherichia coli and Salmonella isolated from fresh and frozen chicken meat sold in Ibadan metropolis.

Methods
The samples were processed for isolation, identification and antibiotics susceptibility. The phytochemical components of aqueous and methanolic extracts of the three plants were obtained using column chromatography. Antimicrobial activity of the plants extracts was done using agar well diffusion method.

Conclusions
Most of the bacterial isolates were resistant to four or more commonly used antibiotics. All the plant extracts exhibit antibacterial activity; there was a significant difference in the antibacterial activity of the methanolic extract of the plants studied (P<0.05). Among the bacteria tested Escherichia coli was the most sensitive while Psidium guajava showed extensive inhibition zone. This study reveals that there is high prevalence of multi-drug resistant bacteria in the chicken meat tested and that Psidium guajava is potential source of antibiotics for drug development against the bacteria studied.
Background

The emergence of carbapenem resistance in Enterobacteriaceae is an important threat to global health. The reason of prevalence of bacterial resistance to antibiotics continues to increase, infections caused by carbapenem-resistant Enterobacteriaceae (CRE) are responsible for high rates of mortality and represent a major therapeutic challenge.

Objectives

The aim of this study was to investigate effective antibiotic combinations against carbapenem-resistant Enterobacteriaceae (CRE) strains.

Methods

The minimum inhibitory concentrations (MIC) of doripenem, colistin, tigecycline and tobramycin were determined by microbroth dilution method against 39 clinical strains isolated from patients in the intensive care unit (ICU). Carbapenem resistance mechanisms in these isolates were investigated by PCR technique. The in vitro synergistic activities of tigecycline, colistin, tobramycin and doripenem in double combination were determined by microbroth chequerboard technique and results were interpreted using the fractional inhibitory concentration (FIC) index.

Conclusions

Based on MIC90s, colistin (1 mg/l) was the most potent agent, followed by tigecycline (2 mg/l), tobramycin (32 mg/l), and meropenem (32 mg/l). According to PCR studies, carbapenem resistance in tested Enterobacteriaceae isolates is most often mediated by OXA-48 type carbapenemases. Additionally six isolates were positive for the
genes encoding VIM and one isolates was positive for the genes encoding KPC. Given that a FIC index of ≤ 0.5 is borderline, synergistic interactions were most frequent with tigecycline-tobramycin (31 %). No antagonism was observed. The findings of this study may play a useful role in selecting the appropriate combinations when a single agent is inadequate.
Background
Difficulties involved in treating drug-resistant pathogens have created a need for new therapies.

Objectives
In this study, we investigated the possibility of natural compounds as an adjuvant for antibiotics against Acinetobacter baumannii.

Methods
Measurement of the fractional inhibitory concentration (FIC) for OA demonstrated that it only synergizes with aminoglycoside antibiotics. Other classes of antibiotics (e.g., ampicillin, rifampicin, norfloxacin, chloramphenicol, and tetracycline) have additive effects with OA. Unlike OA, HBA and PP synergize only with chloramphenicol antibiotics. Microarray and quantitative reverse transcription-PCR analysis under OA indicated that genes involved in ATP synthesis and cell membrane permeability, the gene encoding glycosyltransferase, peptidoglycan-related genes, phage-related genes, and DNA repair genes were upregulated. OA highly induces the expression of adk, which encodes an adenylate kinase, and des6, which encodes a linoleoyl-CoA desaturase, and deletion of these genes increased FICs; these observations indicate that adk and des6 are involved in the synergism of OA with aminoglycosides. Data obtained using 8-anilino-1-naphthalenesulfonic acid, fluorescence-conjugated gentamicin, and membrane fatty acid analysis indicates that adk and des6 are involved in changes in membrane hydrophobicity and permeability. Proton-motive force and ATP synthesis tests show that those genes are also involved in energy metabolism.

Conclusions
We screened 28 natural compounds from plants and confirmed that a pentacyclic triterpenoid, oleanolic acid (OA) could be used only with aminoglycoside antibiotics and other natural compounds (4-hydroxybenzaldehyde and propyl 4-hydroxybenzoate) could be used only with chloramphenicol antibiotics. Taken together, our data show that OA boosts aminoglycoside uptake by changing membrane permeability and energy metabolism in A. baumannii.
ANTIBIOTIC SUSCEPTIBILITY TESTING OF SALMONELLA ENTERITIDIS ISOLATES FROM RAŠKA REGION OF SERBIA

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Background

Salmonella spp. are zoonotic agents widely spread amongst poultry, cattle and swine which are marked as one of the leading causes of gastroenteritis in humans in the modern world. Today’s open market and trade between countries are main reasons of propagation of Salmonella serotypes worldwide.

Heavy use of antibiotics in raising animals leads to selective pressure to Salmonella serotypes, allowing propagation of resistant serotypes and development or acquisition of genes, gene complexes or mutations which are holders of resistance. Multiresistant Salmonella serotypes can transfer resistance genes to other bacterial species.

Objectives

The aim of this investigation was to determine dominant Salmonella serotypes in poultry production in the region of Raška in Serbia and check their resistance to the commonly used antibiotics.

Methods

Salmonella Enteritidis was determined as one of dominant serotypes in the investigated period (2010/2011). Resistance testing was conducted on 41 Salmonella Enteritidis isolates. Test method was NLCCS Disk Diffusion Method. Antimicrobial panel comprised of 16 antibiotics.

Conclusions

Most of the samples were sensitive to all 16 antibiotics. Five isolates were resistant to flumequine and nalidixic acid. Six isolates showed intermediate sensitivity to neomycin and two were intermediate sensitive to tetracycline. This result is not surprising considering fact that these antibiotics are commonly used in poultry production in Serbia. These results suggest that it would be necessary to reassess use of flumequine and limit its use only to cases in which the causative agent had
exhibited resistance to other antibiotics and the bacteriological confirmation of diagnosis and susceptibility testing of pathogens justifies its application.
A NEW PROMOTER FOR EXPRESSION OF THE AAC(6')-IB-CR GENE IN CLASS 1 INTEGRONS.

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Background
Class 1 integrons are genetic elements able to acquire, exchange and express gene cassettes mainly encoding antibiotic resistance. GCs are usually promoterless and are expressed from a common promoter Pc. Thirteen Pc variants of different strenghts have been described, including a weak (PcW) variant. The aac(6')-ib-cr gene, conferring resistance to aminoglycosides, norfloxacin and ciprofloxacin, is mostly found at first position in class 1 integrons. A 101-bp sequence upstream of aac(6')-ib-cr, contains a putative promoter (named PA in this study) suggesting a an expression control by both Pc and PA.

Objectives
The role of respectively Pc and PA in expression of aac(6')-ib-cr will be evaluate.

Methods
A 484-bp fragment, from a clinical class 1 integron (Klebsiella pneumoniae C1911) and containing the PcW variant, the PA promoter and the first 194-bp of the aac(6')-ib-cr gene cassette, was cloned upstream the lacZ reporter gene. Wild-type and mutated promoters were used. Beta-galactosidase assays and quantification of lacZ transcripts were used to estimate the level of aac(6')-ib-cr expression. Mapping of the transcription start site by 5′ RACE and characterization the N-terminal amino acids of the protein are in progress.

Conclusions
Both promoters are involved in the transcription of aac(6')-ib-cr. However, the level of transcription is slightly increased (12% with begal assays) when PcW is mutated whereas it is highly decreased (84%) when PA is mutated.
IN-VITRO EVALUATION OF NISIN ALONE AND IN COMBINATION WITH POLYCATIONIC POLYMERS AGAINST HOSPITAL-ACQUIRED METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS (HA-MRSA) STRAINS

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Background

Nasal colonization by Methicillin-resistant Staphylococcus aureus (MRSA) is a problem in the hospital setting due to the required hygienic measures as well as the possibility of developing an infection. Polycationics such as Nisin or Chitosan are known to possess antimicrobial activity against a whole set of different bacteria including Staphylococcus aureus. The use of combinations may be one way to increase antimicrobial activity and to reduce side effects at the same time.

Objectives

The aim of this study was to investigate whether the use of the antimicrobial lantibiotic Nisin in combination with different types of polyethylenimine and chitosan show an increased antimicrobial effect towards MRSA.

Methods

The tested MRSA strains were provided by the Institute of Medical Microbiology and Hygiene Regensburg, Germany. Minimum inhibitory concentration (MIC) values were determined according to the CLSI microdilution broth method and at pH 5.5 to assess the efficacy of the antimicrobials. Likewise checkerboard assays were performed at pH 5.5.

Conclusions

Combining the antimicrobial peptide Nisin with the polycationic polymers shows promising activity against MRSA strains. Although no synergism could be detected, additive effects were clearly visible. This may be of advantage for the formulation of drug delivery systems for the treatment of nasal MRSA infections.
Antimicrobial resistance

MOLECULAR DETECTION AND CHARACTERISATION OF METALLO-ß-LACTAMASE GENES AND INTEGRONS OF IMIPENEM–RESISTANT PSEUDOMONAS AERUGINOSA

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Background

The increase in the prevalence of carbapenem-resistant Pseudomonas aeruginosa isolates, particularly caused by the metallo-ß-lactamases (MBL) is of great concern in the clinical settings worldwide.

Objectives

Here, we report the detection and identification of MBL producing P. aeruginosa in our hospital (Istanbul University Medical Center, Turkey), and the characterization of MBL cassette-containing integrons in 90 imipenem-resistant P. aeruginosa isolates. Of the isolates investigated in this study, 32 (94%) MBL-positive strains were identified, harboring either blaIMP or blaVIM genes. The genes detected from MBL-positive isolates were blaIMP-7, aacC1; blaVIM-11; aacA7, blaVIM-2, attI, and aacA7.

Methods

In this study, three phenotypic methods, i.e., combined IPM-EDTA disk test (CDT), double-disk synergy test (DDST), and IP/IPI E-test) were evaluated for MBL detection in imipenem-resistant P. aeruginosa (IRPA) clinical isolates, in comparison to PCR detection of MBL genes as the gold standard. Integrons and their associated gene cassettes were characterized and the genetic relatedness of the isolates was investigated using random amplification of polymorphic DNA (RAPD) analysis.

Conclusions

This study highlights the resistance to imipenem due to IMP- and VIM-producing P. aeruginosa and their associated class 1 integrons. Horizontal dissemination of the class 1 integron-associated MBL genes may contribute to the further emergence of carbapenem resistance in other Gram-negative bacteria. Therefore, appropriate surveillance and control measures are essential to prevent the further spread of MBL-producing organisms in hospitals. Further studies should be carried out to give a
better understanding of the impact of integrons on the dissemination of antimicrobial resistance in the clinical settings.
DETECTION OF AGR SYSTEM AND ANTIMICROBIAL RESISTANCE IN S. EPIDERMIDIS PRODUCER BIOFILM ISOLATED OF AN UNIVERSITARY HOSPITAL OVER 20 YEARS.

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Background

S. epidermidis isolated from blood culture is highly important in a hospital.

Objectives

To establish the \textit{agr} locus, the susceptibility of biofilm–producing \textit{S. epidermidis} specimens to antimicrobial agents, and to analyze the clonal profile of \textit{S. epidermidis} isolated from blood culture from an Universitary Hospital, Brazil, along a 20–year period.

Methods

Detection of biofilm–related genes (\textit{icaA}, \textit{icaB}, \textit{icaC} and \textit{icaD}, \textit{bhp} and \textit{aap}) by polymerase chain reaction (PCR) and expression by reverse transcriptase polymerase chain reaction (RT-PCR) showed 83.6\% of \textit{ica} operon, 11.5\% of \textit{bhp} gene, and 32.8\% of \textit{aap} gene for 61 \textit{S. epidermidis} specimens. The \textit{agrI} and \textit{agrII} loci were detected in 77\% and 19.7\%, respectively. Oxacillin–resistance was verified by detection of \textit{mecA} gene in 90.1\%, and the minimal inhibitory concentration (MIC) values assessed by E-test have shown 83.6\% resistance to oxacillin. Clonal profile determination showed 11 clusters, and the ST2 type was determined to be the major cluster.

Conclusions

The ability of \textit{S. epidermidis} to produce biofilm has been shown, with predominant \textit{agrI} locus, oxacillin–resistance, and highest prevalence of SCC\textit{mec} type III. Some isolates showed resistance to newer antimicrobial agents, and molecular type determination revealed clones persisting for up to 17 years. The major one was the \textit{S. epidermidis} ST2 invasive clone, pointing to the relevance of considering such
organisms as important pathogens with potential ability to cause serious infection in inpatients, as well as taking into account the potential dissemination of such resistant pathogenic clones in hospital settings.
RESPONSE TO ENVELOPE STRESS BY THE ZRAP/SR SYSTEM IN ESCHERICHIA COLI

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Background
To detect and respond to cell envelope damages, Gram-negative bacteria possess multiple envelope stress pathways. The \textit{E. coli}, ZraPSR system is composed of a two component system ZraSR that comprises a membrane associated sensor kinase ZraS and a cytoplasmic response regulator ZraR. This system has the particularity to encompass a third component, the periplasmic protein ZraP. The \textit{zraP} gene is up-regulated by ZraR in the presence of zinc. Moreover, the deletion of \textit{zraP} in \textit{Salmonella typhimurium} dramatically affected its resistance to polymyxin B, a cationic antimicrobial peptide (Appia-Ayme \textit{et al}, 2012).

Objectives
A connexion might exist between Zn(II) metabolism and antibiotics sensitivity, mediated by the ZraPSR system. In this study, we investigated this connexion and the ZraPSR signaling pathway in \textit{E. coli}.

Methods
WT and \(\Delta zraP\) strains were phenotypically tested for their drug susceptibility profiles.\(PzraP::lux\) and \(PzraSR::lux\) transcriptional fusions were monitored in the presence of metals and drugs. Minimal inhibitory concentrations of zinc, copper, lead and antibiotics were determined for \(\Delta zraP\), \(\Delta zraS\) and \(\Delta zraR\) \textit{E. coli} strains.

Conclusions
At high zinc concentrations, the expression levels of the two divergent operons \textit{zraP} and \textit{zraSR} increased by 100- and 40-fold, respectively. This suggested that the ZraPSR system is involved in zinc homeostasis. However we showed that high metal concentrations do not affect the survival rate of any \textit{zraP},\textit{zraS} or \textit{zraR} mutants. In contrast, these \textit{zraPSR} mutants were more sensitive to some envelope targeting antibiotics. This indicates that ZraPSR contributes to the envelope stress response of \textit{E. coli}. 

IDENTIFICATION OF DIVERSE CAPSULAR TYPES IN MULTIDRUG-RESISTANT (MDR) KLEBSIELLA PNEUMONIAE CLONES USING WZI SEQUENCING AND FTIR

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Background
Klebsiella pneumoniae polysaccharide capsule is a well-known determinant of virulence and pathogenicity, but their identification by conventional methods is laborious and time-consuming. A genotypic-based method has been proposed for capsular typing based on wzi sequencing, one of the genes from the cps cluster. Identification of capsular types in MDR K. pneumoniae high-risk clonal groups suggested intraclonal diversity but it has mainly been explored for carbapenemase-producing CG258 isolates.

Objectives
To assess inter and intraclonal diversity of capsular types in a representative collection of MDR K. pneumoniae isolates by genotypic, phylogenetic and spectroscopic approaches.

Methods

One hundred and four previously characterized MDR K. pneumoniae isolates from different clones or CG (ST14, ST15, ST39, ST54, ST101, ST147, ST336, CG258) producing diverse ESBLs and/or carbapenemases were studied. They represent a snapshot of MDR clones responsible for outbreaks or endemic situations in different countries (Brasil, Greece, Poland, Portugal, Romania, Spain). Characterization of capsular types included sequencing and phylogenetic analysis of wzi genes (including wzi sequences from genomes deposited in NCBI), and Fourier Transform Infrared Spectroscopy (FTIR) coupled with multivariate data analysis (PCA/PLSDA).

Conclusions

The diversity of capsular types identified among CG15, CG258 or ST11 isolates highlights an intraclonal diversity that might reflect a suboptimal resolution of MLST-based typing. The clustering of the isolates based on multivariate data analysis of
FTIR spectra according to sequence types and capsular types, which was corroborated by the topology obtained from the phylogenetic tree of wzi sequences, suggests the potential application of FTIR for discrimination of K. pneumoniae isolates.
EFFICACY OF BIOACTIVE COMPOUNDS FROM ACACIA ARABICA AGAINST MULTI-DRUG RESISTANT CLINICAL BACTERIAL PATHOGENS

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Background

The emergence of infectious diseases and development of drug resistance in pathogenic bacteria at an alarming rate is a matter of serious concern. Therefore, there is an urgent need to discover novel strategies and identify new antimicrobials from natural substances for development of next generation drugs. Hence screening of novel antibacterial agents was carried out from the bark of Acacia arabica with traditional value in ayurvedic medicine.

Objectives

The study aimed to identify bioactive compounds with antibacterial activity isolated from bark extracts of Acacia arabica. The inhibitory and cidal concentration of the compounds were evaluated. Further, synergistic effect of potent bioactive compounds with standard antibiotic was determined.

Methods

Bark extracts of Acacia arabica were prepared by cold percolation method using polar to non-polar solvents in succession. Antimicrobial activity of the extracts was evaluated using Agar well diffusion method against multi-drug resistant bacteria. Using chromatographic techniques, the bioactive compounds from the extracts were isolated and purified. The purity of the compounds was assessed using high performance liquid chromatography and the compounds were identified using GCMS and NMR spectroscopic techniques. The inhibitory and cidal concentration of the compounds were determined using Resazurin micro-titre assay plate method. The compound exhibiting maximum antibacterial activity with a minimum MIC value was evaluated for synergism with antibiotic.

Conclusions

The results showed that acetone extract was most effective in inhibiting the multi-drug resistant bacterial pathogens. The compounds identified from acetone extract showed positive synergism with antibiotic tested suggesting the possibility of new drug development.
Antimicrobial resistance

NANO-ENCAPSULATED COLISTIN SULPHATE: ANTIPSEUDOMONAL ACTIVITY

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Background

Pseudomonas aeruginosa infects the respiratory tract of cystic fibrosis (CF) patients. Multidrug resistant phenotypes and a high capacity to form stable biofilms are frequent. Despite colistin-resistance is infrequent, often emerges in patients treated with inhaled formulations for long periods. The use of nanoparticles can contribute to overcome drug resistance mechanisms. Some authors have reported that lipid nanoparticles loaded with aminoglycosides are more effective than free formulations against clinical isolates.

Objectives

The aim was to explore antimicrobial activity of nanoencapsulated colistin (SLN-NLC) versus free drug against P. aeruginosa clinical isolates from CF patients and to investigate their efficacy in biofilm eradication.

Methods

10 mg of colistin was mixed with a 5% (w/v) Precirol® ATO 5 dichloromethane solution. Then, the organic phase and Poloxamer 188 at 1% w/v and Polysorbate 80 at 1% w/v were mixed and emulsified by sonication, solvent evaporated and SLNs obtained filtered. NLC were made with Precirol® ATO 5 and Miglyol® 812 as lipid core. Susceptibility of planktonic bacteria to antimicrobials was examined by using the broth microdilution method and time-kill kinetic curves. Minimal Biofilm Eradication Concentration (MBEC) was determined to assess antimicrobial susceptibility of sessile bacteria. Treated and untreated biofilms were visualized by Atomic Force Microscopy (AFM).

Conclusions
Colistin nanoparticles presented slightly higher activity than free drug against planktonic bacteria. On the contrary, nanoencapsulated colistin was much more efficient in the eradication of biofilms than free colistin. Thus, these formulations have to be considered as a good alternative therapeutic option to treat *P. aeruginosa* infections.
CANDIDA GLABRATA PHENOTYPIC SWITCHING AND ITS ASSOCIATION WITH ECHINOCANDIN ANTIFUNGAL RESISTANCE.

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Background

*Candida glabrata* stands up as the second most prevalent pathogen of mucosal and systemic fungal infection, being the non-*albicans* species with the highest mortality rate associated. Since it possesses both innate and acquired resistance against azole antifungal drugs, echinocandins became the first-line therapy in many hospitals. Recently, *C. glabrata* pathogenicity was associated with phenotypic switching, an important virulence attribute in *C. albicans*, but yet poorly characterized in *C. glabrata*.

Objectives

We intend to reveal phenotypic switching as a predictor to uncover the emergence of echinocandins antifungal resistance in *C. glabrata*.

Methods

Equinocandin resistance induction assays were performed with CBS 138 strain and two clinical isolates. *C. glabrata* strains were daily exposed to the following echinocandins: caspofungin (CSF), micafungin (MCF) and anidulafungin (ANF). Every 5 days of incubation, minimal inhibitory concentrations (MIC) were determined, according to the CLSI protocol M27-A3 S4.

Phenotypic switching assessment was carried out during induction of antifungal resistance, by plating each strain in medium containing phloxine B.

Conclusions

At the beginning of induction assays susceptible colonies were Light Pink (LP) or Medium Pink (MP). In all strains, resistance to echinocandins was detected at day 5 and colonies were MP and Dark Pink (DP). The MICs of the different phenotypes were determined and differences in the susceptibility to echinocandins were found: LP and MP colonies remain susceptible and the DP were resistant.
The continuous exposure to serum levels of echinocandins promotes the acquisition of antifungal resistance in *C. glabrata* together with the emerging of phenotypic switching.
Antimicrobial resistance

HIGH VIRULENT CMY-2 AND CTX-M-2-TYPE-PRODUCING E. COLI CAUSING COLIBACILLOSIS IN COMMERCIAL TURKEYS FROM BRAZIL

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Background

A high diversity of AmpC and extended-spectrum (ESBL) beta-lactamases has been described in Brazil among isolates from nosocomial and community settings, but this is the first report of these enzymes in turkey Brazilian farms.

Objectives

The aim of this study was to characterize ESBL and plasmid-mediated AmpC-producing E. coli isolated from commercial turkey in Brazil.

Methods

In 2009, 227 E. coli strains were isolated from air sac of turkeys condemned by airsacculitis and screened for ESBL and pAmpC production. MICs were determined by microdilution, and beta-lactamases-encoding genes and virulence factors were investigated by PCR. Plasmids were extracted and transformed into Top10 electrocompetent E. coli strain (Invitrogen™) followed by direct sequencing. Genetic relatedness of beta-lactamase-positive strains was determined by ERIC-PCR fingerprinting.

Conclusions

The blaCTX-M-2 and blaCMY-2 genes were confirmed in four and seven APEC strains, respectively; which were clonally unrelated, showing high MIC to cefoxitin (MIC≥64 mg/L). The blaCMY-2 genotype was associated with the presence of an IncI1 plasmid of approximately 94kb, which showed 98% of similarity with pSTM709 plasmid (Genbank: HG428759.1), identified in Salmonella Typhimurium in Uruguay, confirming its spread in Latin America. Much more concerning is the high virulent profile exhibited by these beta-lactamase-producing APEC, which shared the same virulence factors of human ExPEC. In this regard, the dissemination of cephalosporin-resistant bacteria carrying blaESBL and/or blaCMY-2 genotypes has the potential to impact both veterinary and human therapeutic treatment options, which requires more surveillance of antimicrobial resistance in bacteria from food-producing animals.
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TESTING THE MUTANT SELECTION WINDOW (MSW) HYPOTHESIS WITH GRAM-NEGATIVE BACTERIA EXPOSED TO THE FLUOROQUINOLONES IN IN VITRO DYNAMIC MODELS

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Background

According to the MSW hypothesis, resistant mutants are selected at antibiotic concentrations between the MIC and the mutant prevention concentration (MPC). This hypothesis has been tested mainly with fluoroquinolone-exposed Gram-positive pathogens but not Gram-negatives.

Objectives

To test the MSW hypothesis with Escherichia coli, Klebsiella pneumoniae and Pseudomonas aeruginosa, ciprofloxacin pharmacokinetics were simulated in an in vitro dynamic model.

Methods

Four strains each of E. coli and Pseudomonas aeruginosa and three strains of K. pneumoniae with MPC/MIC ratio from 4 to 64 were exposed to twice-daily ciprofloxacin for three days over 50-380-fold ranges of the ratio of the area under the concentration-time curve (AUC) to the MIC. Peak antibiotic concentrations were simulated to be close to the MIC, between the MIC and the MPC, and above the MPC.

Conclusions

With all three bacterial species, resistant mutants were enriched at ciprofloxacin concentrations that fell into the MSW for most of the dosing interval but not at concentrations out of the MSW. Both AUC/MIC- and AUC/MPC-resistance relationships were bell-shaped, showing applicability of the MSW hypothesis. These relationships predict highly variable “anti-mutant” AUC/MIC ratios - 525-1070 h (E. coli), 560-1800 h (K. pneumoniae) and 225-1100 h (P. aeruginosa) – and AUC/MPC ratios – 25-100, 28-140 and 7-204 h, respectively.
BACTERIAL RESISTANCE STUDIES USING IN VITRO DYNAMIC MODELS: NEW INSIGHT IN THE DESIGN OF “ANTI-MUTANT” ANTIBIOTIC DOSING

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Background

In vitro models that simulate human pharmacokinetics of antibiotics allow pharmacodynamic comparisons. This approach can be used to examine the abilities of antimicrobials to suppress not only antibiotic-susceptible pathogens but also their resistant mutants.

Objectives

To assess the ability of fluoroquinolones, lipo- and glycopeptides to prevent the selection and/or to suppress resistant mutants of \textit{Staphylococcus aureus}, multiple dosing regimens of the antibiotics were simulated in an \textit{in vitro} dynamic model.

Methods

A series of monoexponential pharmacokinetic profiles that mimic once-daily administration of moxifloxacin, gatifloxacin, levofloxacin and daptomycin and twice-daily dosing of ciprofloxacin and vancomycin, were simulated with half-lives reported in humans. The simulated ratios of the area under the concentration–time curve (AUC) to the MIC varied in a wide range that includes clinically relevant AUC/MICs.

Conclusions

With each agent resistant \textit{S. aureus} mutants were enriched in a concentration-dependent manner. This enrichment was accompanied by concomitant loss in the susceptibility of antibiotic-exposed bacteria. Both in terms of the population data analysis and susceptibility testing, AUC/MIC-resistance relationships were bell-shaped. Based on these relationships, the threshold “anti-mutant” AUC/MIC ratios were predicted. With moxifloxacin and gatifloxacin but not levofloxacin and ciprofloxacin, and with daptomycin and vancomycin, the predicted thresholds were within the clinically attainable AUC/MIC ratios.
CHARACTERIZATION OF MUTATION IN STREPTOMYCIN-RESISTANT MYCOBACTERIUM TUBERCULOSIS AND THE ASSOCIATION BETWEEN BEIJING-LINEAGE AND DUAL-MUTATION IN GIDB GENE

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Background
In developing countries, resistance of Mycobacterium tuberculosis to streptomycin (STR) is a severe threat to tuberculosis treatment. Mutations in rpsl, rrs, and gidB genes are reported to be related to STR resistance, some of which are suggested to be associated with M. tuberculosis lineage, while this association is geographically varied.

Objectives
In this study, we aimed to investigate the mutation characteristics of STR resistance and the association between the polymorphism of drug-resistant genes and M. tuberculosis lineage in Sichuan, China.

Methods
A total of 180 STR-resistant and 47 susceptible M. tuberculosis clinical isolates were genotyped by spoligotyping followed by mutation sequencing of rpsl, rrs and gidB genes.

Conclusions
No mutation was released in rpsl and rrs genes among sensitive isolates, while in resistant ones, 150 harbored mutations in rpsl or rrs, suggesting a correlation between STR-resistance and mutations of these two genes. Mutation K43R in rpsl was found to strongly associate with high-level STR resistance (P < 0.01), while those in rrs and gidB potentially contributed to low-level resistance (P < 0.05). No significant geneal association was presented between STR-resistance and Beijing genotype, however, Beijing genotype was significantly correlated with high-level STR-resistance, as well as with K43R (P < 0.01), indicating that Beijing genotype has an evolutionary advantage under streptomycin pressure. Interestingly, a dual mutation of E92D and A205A in the gidB gene was detected in all isolates of Beijing genotype, suggesting this dual mutation have the potential to be used as an effective marker for identifying Beijing genotype.
Background
Yeast infections, caused by Candida albicans are health- and life-threatening, especially to patients with chronic diseases or immunodeficiency. Photodynamic inactivation (PDI) can be successfully used to overcome the problem of resistance of microorganisms to classical antimicrobial chemotherapies.

Objectives
The aim of this study was to compare the effect of nine imidazoacridinone (IA) derivatives used as photosensitizing agents in PDI against C. albicans ATCC and clinical strains.

Methods
The Quality Control and clinical strains of Candida albicans suspensions were treated with IA derivatives (50 µM concentration) and irradiated with 50 and 100 J/cm². The light source used was custom made LED lamp (Secure Media Poland), of maximum output power of 630 mW/cm², which emits blue light (405 nm). Cell suspensions kept in the dark were used as controls. After irradiation serial dilutions from 10⁻¹ to 10⁻⁴ were obtained from every sample and colony forming units/ml (CFU) were determined. Moreover, the accumulation of IAs in 3 ATCC strains was measured using spectrophotometric methods.

Conclusions
The use of IAs in PDI was able to promote a significant antimicrobial effect against the planktonic forms of C. albicans (>5 log₁₀ units reduction in survival). The PDI effect did not exclusively depend on the chemical structure of the photosensitizing compounds or their accumulation in C. albicans cells. We could observe very strong strain-dependent response to PDI treatment. Our results showed that PDI outcome did not strictly depend on physico-chemical processes, but biological background of particular strain played also an important role in this reaction.
MOLECULAR CHARACTERIZATION AND ANTIBIOTIC SUSCEPTIBILITY OF METICILLIN RESISTANT STAPHYLOCOCCUS AUREUS (MRSA) IN NORTH INDIA, WITH SPECIAL REFERENCE TO COMMUNITY ACQUIRED MRSA

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Background
Infiltration of community acquired meticillin resistant Staphylococcus aureus (CA-MRSA) strains into hospitals and hospital acquired (HA) MRSA into community has become a major hurdle leading to poor clinical outcomes. Role of Panton-Valentine leukocidin (PVL) and arginine catabolic mobile element (ACME), previously associated with CA-MRSA, still remains controversial.

Objectives
To study relative frequency of CA and HA- MRSA in outpatients, inpatients and to find and correlate presence of PVL and ACME.

Methods
Total 147 MRSA isolates were collected from clinical bacteriology lab of a tertiary care hospital. Isolates were characterized by PCR amplification of mecA, PVL, ACME genes and staphylococcal cassette chromosome mec (SCCmec) typing. Patient risk factors, antimicrobial susceptibility profiles and pcr results were compared. A total of 129 MRSA strains were mecA positive, and 34 (27.9%) were PVL gene positive. SCCmec III were 31 (24%), 41 were SCCmec IV (31.78%), and 32 were SCCmec V (24.8%). Of all SCCmecIII strains, multidrug resistant(MDR) - 70%. In SCCmec IV, 51.2% were MDR and in mecV strains - 31.25% MDR, further supported by presence of risk factors. PVL positivity in case of mecIII,IV,V was 2.94%,52.9%,41% respectively. ACME arc gene was found in 5(3.87%) cases, 2 of 5 were CA-MRSA non MDR isolates without risk factors. ACME OPP3gene was absent.

Conclusions
This study demonstrated the presence of CA-MRSA in hospital settings and vice versa. mecIII MDR strains are no longer restricted to hospital settings and mecIV, V are frequently found in hospital settings with MDR characteristics. But PVL is still present mostly in mecIV and V.
Antimicrobial resistance

CARBAPENEMASE-PRODUCING KLEBSIELLA PNEUMONIAE AND ESCHERICHIA COLI IN SERBIA

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Background

Emergence and dissemination of carbapenemase-producing Enterobacteriaceae is rising worldwide, representing a public health threat. Although most European countries have already established their national surveillance programs considering carbapenemase-producing enterobacteria, so far there are no reports from Serbia.

Objectives

The aim of the study was to determine the occurrence and types of carbapenemases in carbapenemase-producing Klebsiella pneumoniae and Escherichia coli in Serbia.

Methods

Strains of Klebsiella pneumoniae and Escherichia coli from 14 different hospitals resistant to at least one carbapenem (imipenem, meropenem, ertapenem) were collected from November 2013 to May 2014. In National Reference Laboratory for Susceptibility Testing carbapenem resistance was confirmed. PCR assays were performed for the detection of genes encoding carbapenemases (bla\textsubscript{OXA-48}, bla\textsubscript{VIM}, bla\textsubscript{NDM} and bla\textsubscript{KPC}).

Conclusions

Of 125 collected strains 117 (93.6%) were Klebsiella pneumoniae and 8 (6.4%) were Escherichia coli. Sixty seven (53.6%) strains were obtained from urine, 25 (20%) from blood, 19 (15.2%) from wound secretions and 14 (11.2%) from lower respiratory tract secretions. Seven (87.5%) Escherichia coli harbored bla\textsubscript{NDM} gene and in one isolate no carbapenemase encoding genes were detected. Gene bla\textsubscript{NDM} was found in 34 (29.1%) Klebsiella pneumoniae, bla\textsubscript{OXA-48} in 10 (8.5%), bla\textsubscript{KPC} in 1 (0.9%), and 7 (6%) strains harbored both bla\textsubscript{NDM} and bla\textsubscript{OXA-48}, while in 65 (55.6%) Klebsiella pneumoniae no carbapenemase encoding genes were detected.

In Serbia the most common type of carbapenemase in both carbapenemase-producing Klebsiella pneumoniae and Escherichia coli is NDM. Carbapenemase-
producing *Klebsiella pneumoniae* and *Escherichia coli* were most frequently isolated from urine.
MONITORING OF THE ENTEROCOCCI RESISTANCE MECHANISMS TO AMINOGLYCOSIDES

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Background

Enterococcus faecalis and Enterococcus faecium are opportunistic bacteria responsible for frequent nosocomial infections. Presently, resistance of enterococci to antibiotics, especially aminoglycosides, glycopeptides and oxazolidinones is becoming a severe clinical problem.

Objectives

The aims of the study were as follows: determination of phenotypic patterns of enterococci resistant to aminoglycosides, detection of genes responsible for above resistance, investigation of relationships between analyzed isolates.

Methods

125 E. faecalis and 49 E. feacium isolates resistant to aminoglycosides, obtained from clinical materials in one city hospital during period 2009-2012, were investigated. Besides estimation of antibiotic susceptibility patterns, the search for HLR (gentamicin MIC > 128mg/L and streptomycin MIC > 1024mg/L), HLGR (gentamicin MIC > 128mg/L) and HLKR (kanamycin and amikacin MICs > 512mg/L) phenotypes was undertaken. The presence of the following genes: aac(6')-le-aph(2')-la, aph(2')-lb, aph(2')-lc, aph(2')-ld, aph(3')-llla, ant(4')-la, ant(3')-la, ant(3')-la, responsible for resistance to aminoglycosides was detected by PCR technique. Clonal distribution and relationship between isolates was investigated by PFGE after bacterial DNA digestion with SmaI restructase.

Conclusions

80% of the analyzed isolates belonged to the HLAR group, remaining isolates possessed phenotype HLGR. HLSR phenotype was not detected. Genetic analysis showed that aac(6')-le-aph(2')-la gene responsible for the high resistance to
gentamicin and for cross-resistance to other aminoglycosides (except streptomycin) is the most frequently presented gene among the isolates of tested enterococci. The next common gene was \( \text{aph}(3')-\text{illa} \). Restriction genomic DNA fragments analysis showed large genetic diversity of tested isolates.
ANTIMICROBIAL RESISTANCE OF CLINICAL SALMONELLA STRAINS IN EDIRNE, TURKEY: FOUR YEAR TREND

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Background
Salmonella is one of the most common pathogens of the gastrointestinal tract for both humans and animals leads to food-borne outbreaks and various infections. Accurate diagnosis and treatment of this important pathogen is only achieved by knowing its serogroup and antibiotic susceptibility.

Objectives
In our study, we tried to reveal the distribution and antibiotic resistance of Salmonella strains in the community, also examined the effectiveness of multiplex polymerase chain reaction in the identification of Salmonella.

Methods
Bacterial identification and antibiotic susceptibility tests of 105 Salmonella strains, isolated from the clinical samples (between 2009-2013), were conducted with VITEK2 (Biomerieux, France) automatized system. Strains were grouped with both slide agglutination by using Salmonella polyvalent and group specific antisera (Plasmatec, UK) and multiplex PCR. Multiplex PCR was performed by using six sets of primers targeting O-antigen synthesising gene regions in A, B, C1, D and E serogroups commonly found in clinical isolates.

Conclusions
O-grouping results revealed that serogroup D (68%) and C1 (23%) are the most common causes of Salmonella oriented diarrhea in Edirne, respectively. Multiplex PCR results showed 100% compatibility with serologic diagnosis. The highest level of resistance found against to ampicillin (16%) among all antibiotics. Among four years there is an increasing resistance to cephalosporins, trimetoprim–sulfametoxazol and fluoroquinolones. Salmonella serogroup D is the most frequent serogroup isolated in Edirne and emerging resistance to several antibiotics might be a serious health problem in the future. According to our study, also, multiplex PCR is a reliable and reproducible method in O-grouping of Salmonella.
Background

It was recently proposed that bactericidal antibiotics kill bacteria through a common mechanism involving the production of reactive oxygen species (ROS). However, this mechanisms involving the production of hydroxyl radicals has become the subject of debate. Since the contribution of ROS to antibiotic mediated killing most likely depends on the conditions, differences in experimental procedures are expected to be at the basis of the conflicting results reported in the literature.

Objectives

The goal of this study was to compare methods used to measure the production of ROS and to study ROS production in *Burkholderia cepacia* complex (*Bcc*) biofilms and planktonic cultures.

Methods

Both direct (ROS-specific stainings) and indirect methods (gene-expression analyses, genetic and phenotypic experiments, detection of protein carbonylation and DNA oxidation) to measure ROS were tested and antibiotics belonging to different classes were included.

Conclusions

While some methods were not sensitive enough, others were found to be useful only in some conditions. Since fluorescein based stainings were shown to be pH sensitive and most *Bcc* species, due to resistance need to be treated with antibiotics in high concentrations (which has a marked influence on pH), these dyes cannot be added simultaneously with the antibiotics and control solutions should have the same pH as the antibiotic solutions tested.

Overall our results highlight some methodological key issues to be considered when evaluating the contribution of ROS in antibiotic mediated killing and although not one ideal method could be identified the different methods suggest that ROS is involved in antibiotic mediated killing in *Bcc* species.
Antimicrobial resistance

COLISTIN-RESISTANT MECHANISMS OF KLEBSIELLA PNEUMONIAE IN TAIWAN

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Background
Colistin is an important antibiotic for the treatment of carbapenem-resistant Klebsiella pneumoniae infection.

Objectives
The colistin resistance mechanism(s) of 26 colistin-resistant K. pneumoniae strains in Taiwan were studied.

Methods
We detected pmrH mRNA expression by RT-qPCR and analyzed sequences of pmrHFIJKLM regulators (MgrB, PhoPQ, PmrAB and PmrD).

Conclusions
Determination of the capsular types and multilocus sequence types (MLSTs) of colistin-resistant strains showed that K64 and ST11 were the predominant capsular type (50%) and sequence type (53.9%), respectively. The expression of pmrHFIJKLM operons in 25/26 (96.2%) colistin-resistant strains was significantly higher than in colistin-sensitive strains. The promoter and coding regions of mgrB were interrupted by a transposon in 5 and 3 strains, respectively, and mgrB was absent in 2 strains. Moreover, MgrB Cys28Tyr and Stop48Tyr mutations were observed in 1 and 3 strains, respectively. Site-directed mutagenesis revealed that MgrB Stop48Tyr and PhoQ Leu26Pro mutations in the colistin-sensitive strain NTUH-K2044 increased the mRNA expression of pmrH by 5.3-fold and 39.6-fold, respectively, and both mutations increased the NTUH-K2044 minimal inhibitory concentration for colistin by 32-fold. These results suggest that capsule type K64 is the prevalent type in colistin-resistant strains, and interruptions in mgrB which increased expression of the pmrHFIJKLM operon are major mechanisms contributing to colistin resistance in Taiwan. In addition, novel single amino acid changes in MgrB (Stop48Tyr) and PhoQ (Leu26Pro) were observed to contribute to colistin resistance.
Background

*Staphylococcus fleurettii* is a commensal of animals which can occasionally cause bovine mastitis. *S. fleurettii* strain JW205 isolated from bovine milk in Switzerland exhibited inducible resistance to macrolide, lincosamide and streptogramin B (MLS$_B$) antibiotics suggesting the acquisition of an erythromycin resistance methylase gene (*erm*), but no known Erm determinant was detected.

Objectives

To identify the MLS$_B$ resistance mechanism of *S. fleurettii* JW205 and determine if it is located on a mobile genetic element.

Methods

Whole genome sequencing of strain JW205 and MLS$_B$-susceptible strain M404 (Ion Torrent, Illumina); Comparative analysis of both genomes (progressiveMauve); Cloning and expression of a putative MLS$_B$ resistance gene in *Staphylococcus aureus* RN4220 to demonstrate functionality and inducible expression; Transformation assays by conjugation and electroporation.

Conclusions

The new MLS$_B$ resistance gene *erm*(45) showed 64% aa identity with the nearest related known methylase Erm(B). It was identified on an 11,513-bp novel genomic island in *S. fleurettii* JW205 (ENA acc. no. LN680996). When cloned into *S. aureus*, *erm*(45) confers inducible resistance to macrolide, lincosamide and streptogramin B. Although a circular form of the island was observed, the element could not be transferred into *S. aureus*. Detection of this new methylase emphasized the role of *S. fleurettii* as reservoir of antibiotic resistance genes and demonstrated once again the importance of animal bacteria as pool for novel genetic elements.
Antimicrobial resistance

RESEARCHANT HUB: A COMPREHENSIVE LITERATURE-BASED ANTIMICROBIAL SUSCEPTIBILITY DATABASE

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Background
Antimicrobial resistance is an ever-increasing global public health treat. This phenomenon has stimulated a vigorous research on the field of the antimicrobial susceptibilities and new antimicrobials for the treatment of the so-called superbugs. The decline in the efficacy of the traditionally used antimicrobials is accompanied by the increasing focus on less conventional antimicrobials agents such as plant extracts and experimental compounds. Scientific literature is a rich source containing thousands of either traditionally overlooked or recently discovered antimicrobial compounds. Unfortunately, the potential of these compounds is often not fully exploited. One reason is that literature information is scattered over various sources making the global assessments of the spectra or modes of the action of the compounds difficult.

Objectives
To alleviate the problem of the lack of a comprehensive and centralized antimicrobial susceptibility database, we developed ResearchAnt Hub. RAH is a manually curated database that contains a collection of literature reported antimicrobial susceptibility data. Our compound library contains thousands of experimental compounds with reported antimicrobial activity that hold promise to become the next-generation antimicrobials.

Methods
The antimicrobial susceptibility data was extracted from relevant peer-reviewed scientific journals by manual search and automatic mapping. The taxonomy of the organisms and physical properties of the known compounds were mapped against NCBI and PubChem, respectively.

Conclusions
RAH is the largest antimicrobial susceptibility database to date. It is a tool that can be used for sharing information, search, prediction and discovery of the next-generation antimicrobial agents.
THE INCIDENCE OF TOXIGENIC CLOSTRIDIUM DIFFICILE ASSOCIATED DIARRHEA IN MEXICAN CHILDREN AND ADULTS HOSPITALIZED.

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Background. Toxigenic Clostridium difficile is the most common cause of nosocomial diarrhea, and it is usually a consequence of antibiotic treatment. The incidence and mortality rate of C. difficile infection have increased remarkably worldwide during the last two decades. In Mexico, C. difficile was not considered an important pathogen in children or adults with nosocomial diarrhea. At this time we ignore the incidence and characteristics of the infecting strains.

Objective. To study the epidemiology and virulence of C. difficile from Mexican children and adults with nosocomial diarrhea.

Material and methods. Stools samples were collected from adults and children with nosocomial diarrhea in two tertiary level hospitals in Mexico City. Stools were cultured in cycloserine cefoxitin fructose agar (CCFA), the C. difficile toxin B was confirmed by neutralization with antitoxin, and PCR was performed to identify tcdA, tcdB, cdtA, and cdtB.

Results. We identified C. difficile and its cytotoxic activity in 21 (84%) feces from children and 85 (68.5%) from adults. The majority of the fecal supernatants were positive for toxins of C. difficile, and strains were positive for tcdA, tcdB, cdtA and B. At present, toxigenic C. difficile is a common hospitals-acquired infection in Mexico City.
ONE DECADE INVESTIGATION OF RIFT VALLEY FEVER (RVF) IN IRAN

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Background

Rift Valley Fever Virus (RVFV) is a member of Phlebovirus genus, Bunyaviridae family that cause acute zoonotic viral disease which mostly affect animals (livestock and specially sheep) but can also be a human threat.

Objectives

This disease becomes a concern in Iran after outbreaks in Saudi Arabia and Yemen in 2000s.

Methods

In the last decade human and livestocks sera have been screened; 1206 ovine, 405 caprine, 325 bovine and 28 camel samples and 37 suspected human sera were tested for RVFV in nine provinces in Iran (Sistan va Balooechestan, Khorasan, Tehran, Isfahan, Fars, Kerman, Kurdistan, Hormozgan and Booshehr).

Conclusions

None of these samples were IgG positive. Moreover, among 37 clinically suspected human cases with RVF symptoms, none of these samples were positive for RVFV. Despite no positive cases have been discovered until now, but as Iran is in a high risk region for RVF, the Laboratory of Arboviruses and Viral Hemorrhagic Fevers (National Ref Lab), Pasteur Institute of Iran, continues the surveillance of this acute hemorrhagic fever disease in Iran (in livestock and suspected human).
Background

Several Clostridium spp. strains cause serious, occasionally lethal illness in humans and animals and can be introduced into biogas plants by contaminated feedstock.

Objectives

In order to enlarge the knowledge on the existence and behavior of these pathogens in biogas plants, this study focused (I) on the incidence of Botulinum neurotoxin (BoNT-)producing Clostridium spp., C. difficile, C. novyi, C. haemolyticum, C. septicum and C. chauvoei in process chains of full-scale biogas plants (screening) and (II) on the quantitative development of C. botulinum in lab-scale biogas digesters (sentinel chamber experiments).

Methods

The pathogens were detected by methods combining cultural enrichment with biomolecular analysis using quantitative Real-time PCR. For the screening, 154 samples were analyzed containing 29% plant substrates (e.g. silages), 11% animal substrates (e.g. manure), 27% main digester slurries and 33% digestion products (second stage digesters, storage tanks).

Conclusions

Pathogenic Clostridium spp. were not detected in any of the samples, except for C. novyi (4% of the samples contaminated) and C. difficile, which was found in approx. 50% of the samples, predominantly in animal substrates, digester slurries and digestion products. Sentinel chamber experiments resulted in reduction of C. botulinum by 1.3 to 2.8 log_{10} units after 63 d (95.4% to 99.8% inactivation, D-value: 34.6 ± 11.2 d) at 38°C and by at least 99.7% already after 3 d (D-value: 1.0 ± 0.2 d) at 55°C, proving that the sanitary quality of the digestate compared to untreated manure concerning C. botulinum is improved by mesophilic and thermophilic biogas processes.
CASE REPORT: FIRST REPORT OF PULMONARY INFECTION CAUSED BY HORMOGRAPHIELLA ASPERGILLATA IN ITALY.

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Background
Hormographiella aspergillata is a basidiomycete and belongs to the Psathyrellaceae family. It is an anamorph of Coprinopsis cinerea, which is normally found in compost and sewage. It has been rarely found in human infections, namely a few cases of pulmonary infections and isolated cases of endophthalmitis, endocarditis and sinusitis have been described, mainly in patients with haematological malignancies.

Objectives
We report on a case of pulmonary infection caused by H. aspergillata in a 26 years old male patient affected by “mixed phenotype acute leukemia” admitted to the University Hospital G.B.Rossi in Verona, Italy. This is the first case of infection caused by this pathogen reported in Italy.

Methods
Despite antifungal prophylaxis during hospitalization, Beta-D-Glucan assay resulted in increasingly positive values, followed by a positive result with the Galactomannan assay. Since the patient suddenly complained a severe pain in scapular region, thoracic RX and CT were performed. The CT revealed the presence of a dense mass compatible with a presumptive Aspergillosis in the right lung. After thoracotomy, a proper diagnosis of H. aspergillata was reached by morphological analysis of the growth on Sabouraux agar from biopsy specimen and identification of fungal pathogen was confirmed by molecular tools: sequencing and alignment were conducted on PCR products performed using primers for ITS regions and 26S rRNA gene beyond the D1-D2 hypervariable region.

Conclusions
In diagnosis of rare etiologic agents in immunocompromised hosts, mostly when no experienced medical microbiologist is promptly available, molecular tools are becoming essential for accurate species identification and proper antifungal therapy.
Background
Intestinal parasites are two major groups including protozoa and helminths that inhabit the gastrointestinal tract in humans. They are one of the major health problems of especially poor and under-developed countries.

Objectives
This study was undertaken to determine the prevalence of intestinal parasitic infection in patients with diarrhea and non-diarrhea at the Gulhane Military Medical Academy, Military Hospital in Turkey.

Methods
This retrospective study reviewed the hospital records of 45,638 patients admitted over a ten-year period from 2001 to 2014. During the study period, 2.33% (1859/79,537) of stool samples were tested positive for sixteen species of intestinal parasites by using standard parasitological techniques. *Giardia intestinalis* (1.47%) was the most common parasitic cause of diarrhea among the patients. Its prevalence appears to be decreasing in recent years. *Blastocystis hominis* (0.98%), *Entamoeba coli* (0.74%) ranked second and third in prevalence, respectively. *Taenia* spp (0.20%) was more common in non-diarrheal samples. The stool *E. histolytica* antigen ELISA method had 20% sensitivity and microscopy had 11.80% sensitivity compared to the results of a real-time PCR method. Majority of patients were aged between 14-50 years. Prevalence of intestinal parasitic infection was lowest in winter (11.20%), gradually increased during the spring, reached peaks of 57.34% between June and October, and gradually decreased to 2.28% in December.

Conclusions
These data will help provide accurate estimates of prevalence of intestinal parasites, which are crucial for the development of policies and strategies to enhance their effective control in Turkey.
EMERGENCE OF CLOSTRIDIUM DIFFICILE PCR RIBOTYPE 027 IN HOSPITALS OF SILESIA, POLAND

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Background

According to 2014 ECDC surveillance an reemergence of complicated Clostridium difficile infection (CDI) caused by PCR 027 ribotype strains was found.

Objectives

The aim of this study was to determine the incidence of CDI in 20 Silesian hospitals and to characterize C. difficile isolates from clinical cases.

Methods

In March 2013 to May 2014, 166 stools from patients with clinically suspected CDI were tested by ESCMID algorithm. GDH was tested by CDIFF QUICK CHEK COMPLETE. GDH-positives were examined for toxins by EIA (toxin A/B) and molecular method (IllumiGene) and cultured anaerobically. Antibiotic susceptibility (MIC for 11 antibiotics) was tested with E-tests, (EUCAST interpretation).

Conclusions

Of 160 GDH-positive samples, 140 C. difficile strains were isolated, 110 were PCR ribotyped: 91 (82.7%) belonged to PCR 027 ribotype, three (2.7%) of 176 and 014, two (1.8%) of 010 and 1 strain (0.9%) each of 001, 018, 020 and 046. Seven (6.4%) isolates were non-typeable. All 91 PCR 027 strains were susceptible to vancomycin, amoxicillin/clavulanate and piperacillin/tazobactam and resistant to moxifloxacin, ciprofloxacin, imipenem and erythromycin; 42,9% (39/91) showed resistance to clindamycin. Of 91 isolates tested, 31.9% had MIC> 2 mg/l to metronidazole (ECOFF MIC=2 mg/l) and were resistant to rifampicin.

We confirmed the importance of an appropriate algorithm for rapid CDI diagnostics. The high incidence of C. difficile 027 ribotype is in agreement with recent reports from other countries, though elevated MIC values to metronidazole warrants further exploration.
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ASSOCIATION BETWEEN PCR DETECTION RATES AND SEROLOGICAL RESULTS IN PATIENTS WITH SYPHILIS

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Background

Syphilis is a sexually transmitted disease caused by Treponema pallidum subsp. pallidum. Serological methods are considered to be the standard in syphilis diagnostics. PCR based tests for detection of treponemal DNA are being used more frequently although not routinely.

Objectives

To determine whether the results of PCR detection of treponemal DNA correlate with the results of syphilis serology.

Methods

Swab (n = 179) and whole blood (n = 260) samples were collected from syphilis-seropositive patients during 2004 to 2013 in the Czech Republic and underwent PCR for detection of treponemal DNA (polA, tmpC, TP0136, TP0548 and 23S rRNA genes). Several methods (RPR, VDRL, TPHA, TPPA, ELISA IgM and IgG tests and Western blot IgM and IgG tests) were used to test patients serology. Statistical analysis was conducted for all serological tests and results of PCR for both swabs and whole blood samples.

Conclusions
PCR detection of treponemal DNA was more frequent in swabs (78%) compared to whole blood samples (41%). Western blot analysis revealed that patients with PCR–positive swabs (n = 118) were more frequently IgM positive (p = 0.0002). Analysis of 95 patients where both whole blood and swab samples were available showed that most patients with PCR–positive whole blood had also PCR–positive swabs (89%), however, only 56% of patients with PCR-positive swabs had also PCR-positive whole blood (p = 0.0074). Success rate of detection of all 5 tested loci correlated in swabs with lower RPR titer values. In contrary, whole blood showed the opposite.
REAL-TIME PCR BASED DETECTION OF TROPHERYMA WHIPPLEI CARRIERSHIP IN THE NETHERLANDS

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Background
Whipple’s disease is a rare, potentially fatal multisystemic disorder resulting from infection with the rod-shaped, gram-positive bacterium Tropheryma whipplei. Asymptomatic carriersonship is a precondition of this disease that is thought to be relatively common, but reliable data on the prevalence of carriersonship are sparse.

Objectives
To investigate the prevalence of asymptomatic T. whipplei carriersonship in the Dutch population.

Methods
364 fecal samples were tested for presence of T. whipplei DNA using three different real-time PCR’s: PCR 1, a PCR targeting a repetitive chromosomal locus, (Fenollar e.a. 2004. JCM 42:401-403), PCR 2, targeting the rpoB-gene (Moter e.a. 2013. JCM 51:3858-3861), and PCR 3, a newly developed PCR targeting a unique VNTR region in the genome of T. whipplei (this study). To solve discrepancies between PCR reactions the identity of the amplicons were determined by sequence analysis.

Conclusions
The number of positive samples varied significantly between the three PCR reactions with PCR 1 finding 46/364 (12.6%), PCR 2 23/364 (6.3%), and PCR 3 21/364 (5.8%) positives. Only 19/364 (5.2%) samples were found positive in all three PCR tests. Sequencing data of the samples suggests that none of the 27 discrepant samples of PCR 1 contained sequences that were of T. whipplei origin. Also 3/4 discrepant samples for PCR 2 did not contain T. whipplei derived PCR products, while the remaining discrepant sample did. For PCR 3 sequencing of the two discrepant revealed one true positive (shared with PCR2) and one false positive. This suggests a true prevalence of 20/364 (5.5%).
Background

Outer membrane vesicles (OMVs) are released spontaneously during growth by many Gram-negative bacteria. OMVs have pathogen-like properties inducing a broad protective immune response in hosts.

Classical vaccines were generally ill-defined products but recent recombinant subunit vaccines are developing towards well-characterized pathogen-like structures. Progress in immunology and microbiology can facilitate selection of components that should be or not be present in a platform OMV vaccine.

Objectives

Evaluation of OMVs from Neisseria meningitides as platform vaccine product.

Characterization of the vaccine OMV product with identified structure-function relationship.

Methods

A number of different assays, such as ELISA, DLS, SDS-PAGE and Biacore aid to predict the OMV quality in the accepted potency assay, the Serum Bactericidal Assay (SBA).

For most cases, genetic engineering of the OMV-producing bacteria can be used to improve their application as vaccines. Used modifications include (I) removal of capsular structures and anchor proteins, (II) overexpression of specific antigens, (III) lipopolysaccharide (LPS) modification, (IV) retention of antigens in the membrane that can be secreted, (V) removal of unwanted immune-modulating components.

Examples of these approaches will be given.

Conclusions
The increased knowledge in microbiology and immunology of OMV from bacterial pathogens will enable the development of well-characterized OMV platform vaccine products, as illustrated for Neisseria meningitis OMVs.
THE INCIDENCE OF INFECTION BY BORRELIA BACTERIA AFTER A TICK BITE IN AREAS OF SWEDEN AND FINLAND


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Background
Lyme borreliosis (LB) is a common and increasing tick-borne disease in Europe caused by the Borrelia bacteria. The risk of acquiring a Borrelia infection after a tick bite is not fully known. The transmission of Borrelia bacteria from ticks to humans may depend on many factors such as tick-species, life-stage, duration of tick-feeding, Borrelia species, number of Borrelia bacteria, and site for infestation.

Objectives
We investigated the incidence of Borrelia infection after a bite by a Borrelia-positive tick and if the number of Borrelia bacteria and/or the duration of tick-feeding influenced the risk of contracting a Borrelia infection.

Methods
During 2008-2009, ticks and blood samples were collected from 1546 tick-bitten persons from Sweden and the Åland Islands, Finland. Follow-up blood samples were taken three months after the tick bite. The duration of tick feeding was microscopically
estimated and *Borrelia* bacteria was detected and quantified in ticks by real-time PCR. Anti-*Borrelia* antibodies were detected in sera using ELISA tests and immunoblot.

**Conclusions**
One out of four of the participants were bitten by a *Borrelia*-positive tick but only 10% of these developed signs of a *Borrelia* infection (seroconversion and/or LB diagnosis). Those who seroconverted removed “their” ticks significantly later than those who did not. The number of *Borrelia* bacteria in the ticks did not explain the risk of seroconversion. The risk of developing a *Borrelia* infection after a bite by a *Borrelia*-positive tick is small but increases with the duration of tick feeding.
Background
Cyanobacteria produce various harmful secondary metabolites, which pose a serious global threat to aquatic ecosystems and human health. Biodegradation is a hot topic of water purification research and offers especially an environmentally friendly remediation strategy.

Objectives
The water fungus *Mucor hiemalis* EH5 (*M. hiemalis* EH5) represents a promising biodegrading candidate, as previous studies have shown its ability to breakdown the herbicide isoproturon based on natural metabolic processes and further has been characterized to be resistant to environmental perturbations, such as low temperatures and H2S/sulfide pollution, which would facilitate its application even under extreme environmental conditions.

Methods
Herein we elucidate the effect of three different cyanotoxins, the hepatotoxin microcystin-LR (MC-LR), the neurotoxin β-N-methylamino-l-alanine (BMAA) and the cytotoxin cylindrospermopsin (CYN), on the susceptibility of *M. hiemalis* EH5 using an adaptation of the Kirby-Bauer disk diffusion assay, and the influence on fungal growth and biomass production via radial extension and dry weight measurements. Additionally, we established an optimized strategy for the individual cyanobacterial toxin extraction from the vegetative part of *M. hiemalis* EH5.

Conclusions
The fungal microorganism showed a fast adaptation behavior and strong resistance towards the toxins. No significant differences in terms of growth were perceived compared to an untreated control. This indicates that the cyanobacterial toxins have no toxic effect on the fungus and that the organism can grow and develop undisturbed in their presence. Our results suggest that *M. hiemalis* EH5 is an ideal organism to be tested as a biodegrading system for the remediation of cyanobacterial toxins from contaminated waters.
Background

Forest ecosystems play an essential ecological function by storing up to two thirds of the organic carbon. Before being stored in below-ground a large part of this carbon is contained in litter. Therefore, degradation of plant biomass is an essential process for the proper functioning of forest soils and terrestrial carbon cycling. Soil microorganisms and especially saprotrophic fungi are the principal actors of this process as they produce a wide range of extracellular enzymes involved in the decomposition of the main plant organic polymers. Nature of litter is different depending on tree species and affects diversity and composition of fungal community. We therefore hypothesized that fungal communities selected by the different tree species express specific lignocellulolytic enzymes.

Objectives

We simultaneously analyzed the effect of tree species (beech vs spruce) on the taxonomic composition of the active fungal community in soils using the Elongation factor I (EF1alpha) and on the diversity of expressed fungal genes encoding three main families of Carbohydrate-Active Enzymes (CAZymes) involved in the degradation of the lignocellulose (GH7, GH11 and AA2).

Methods

Environmental cDNA, generated from soil extracted RNA, were used as template to amplify the targeted genes using family-specific primers. All amplicons were sequenced by high-throughput sequencing using Illumina Miseq technology.

Conclusions

Results obtained revealed that only a few numbers (less than 20 percent) of taxonomic and functional clusters were detected in both forest soils. Statistical analysis (AMOVA) showed a significant tree species effect on the diversity of the transcript analyzed.
DIURNAL VARIATION OF BACTERIAL COMMUNITIES IN DRINKING WATER SYSTEMS OVER SMALL SCALES.

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Background

The application of next-generation sequencing techniques to the study of microbial communities that inhabit Drinking Water Systems (DWSs) has revealed the presence of a diverse and abundant microbiome. To our knowledge, the seasonal, weekly and monthly variations of the DW microbiome have been explored, while information on diurnal variation is still lacking. Similarly, sampling efforts mostly focus on covering large spatial scales in the DW, while information on DW microbiome over small spatial scales is still limited.

Objectives

To investigate the variabilities in the drinking water microbiome originating from: (a) diurnal water use patterns, and (b) small-scale spatial effects.

Methods

Triplicate samples of disinfected bulk water (15 to 18L) were taken from 5 sampling locations in Glasgow over 24 hours, in six time periods to capture diurnal effects. The samples were filtered through sterile 0.22um pore size filters and subject to DNA extraction, triplicate 16S rRNA targetted PCR amplification followed by Illumina MiSeq sequencing. Sequence processing was conducted in Mothur using a previously reported protocol. Statistical analyses were conducted using Mothur and R.

Conclusions

Significant differences in richness over a 24-hour period were observed in 3 of the 5 sampling locations. Across sampling locations, a significant difference for the time periods corresponding to higher demand and flows in the system was observed, while differences were not significant for time periods corresponding to lower demand and flows in the system. For all the sampling locations, a change in the community structure and membership throughout the day was observed.
Background

There is a growing need to characterize the effects of environmental stressors at the molecular level on model organisms with the ever increasing number and variety of anthropogenic chemical pollutants. The herbicide 2,4-dichlorophenoxyacetic acid (2,4-D), one of the most widely applied pesticides in the world, is known to have undesirable effects on nontarget species, but the specific molecular details of these effects at sublethal levels are unknown.

Objectives

The model bacterium *Escherichia coli* and the nitrogen fixing, beneficial soil organism *Rhizobium leguminosarum* viciae 3841 (*Rlv*) were used to characterize the specific stress response mechanisms during sublethal 2,4-D exposure.

Methods

We designed a unique platform combining advanced microscopy with metabolomics to determine the sublethal effects of 2,4-D on cell morphology, surface molecular organization and physical properties, and specific target pathways in the metabolic network.

Conclusions

*E. coli* BL21 and a selection of genotypically diverse *E. coli* strains isolated from the environment exhibited filamentous phenotype and envelope remodelling during low level 2,4-D exposure. Sublethal 2,4-D altered vital pathways of cellular metabolism most of which could be attributed to oxidative stress, consistent with increased reactive oxygen species (ROS). *Rlv* exposed to 2,4-D also showed accumulation of ROS and carbonylated proteins coinciding with adaptive changes to integral pathways of cellular metabolism, envelope remodelling and the potential to assimilate 2,4-D. This study identifies biomarkers of 2,4-D exposure in *E. coli* and *Rlv* and offers valuable insights into the mode-of-action of 2,4-D at sublethal levels in soil bacteria.
EFFECT OF OCEAN ACIDIFICATION ON RHODOLITH HOLOBIOT: FREE-LIVING CORALLINE ALGAE AND THEIR ASSOCIATED MICROBIOTA

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Background
Rhodoliths are free-living coralline algae (Rhodophyta, Corallinales) that are ecologically important for the functioning of marine environments. They form extensive beds distributed worldwide, providing a habitat and nursery for benthic organisms and space for fisheries, and are an important source of calcium carbonate. Rhodolith beds are large carbon sinks, but the growth of the Rhodolith holobiont may be affected by lower pH, which is predicted to occur in the near future. The term holobiont refers to any organism and all of its associated symbiotic microbes (parasites, mutualists, synergists and amensals), including endobionts and epibionts that perform diverse ecological roles. A holobiont occupies and adapts to an ecological niche, and is able to employ strategies unavailable in any one species alone when challenged by environmental perturbations.

Objectives
The main goal of this research was to perform an experiment to test the effects of higher pCO₂ (lower pH) on Rhodoliths holobionts in order to analyze whether the microbial community associated with Rhodoliths is perturbed with a change in pH.

Methods
The effect of climate changes, indeed increasing acidification of oceans, on Rhodolith holobiont growth may be due to dissolution of their calcium carbonated skeleton or a change in the microbial community and this was investigate through physiological assays (photosynthesis and calcification) and metagenomics (taxonomical and functional diversity of the holobiont).

Conclusions
The outcomes of the research are an increased understanding of microbes associated with Rhodoliths and new hints on how the holobiont might respond to global climate changes.
Background

Replacement of crude oil with its synthetic and biological equivalents is important for environmental and economical reasons. It is essential to assess synthetically produced alternative fuels in terms of bio/degradability and storage conditions in comparison to crude oils.

Objectives

This study investigates the growth of a model marine microorganism in seawater in the presence of different naval fuels and its impact on corrosion of 1018 carbon steel storage tanks.

Methods

Cultures of Marinobacter isolate J5B1 were incubated in sterilized Key West seawater augmented with different naval fuels, i.e., F76, JP5, FT-F76 and JP5-Camelina. All cultures contained 1018 carbon steel coupons. Bulk phases of each incubation were sampled periodically for cell count and metabolomes over a period of 80 days. Scanning electron microscopy was employed to characterize surfaces of exposed coupons. Polar organic compounds in neat fuels and their water extracts were analyzed using HPLC-Mass Spectrometry.

Conclusions

Marinobacter isolate J5B1 grew in the presence of all tested fuels. HPLC-MS analysis revealed that all neat fuels shared 84 compounds while 21 compounds were common in water extracts of fuels. Fuel F76 was the most and JP5-Camelina was the least diverse in terms of the abundance of polar compounds. Seawater alone could maintain the growth of Marinobacter and water soluble compounds of JP5-Camelina were readily utilized by this isolate. Quaternary ammonium compounds detected in F76 fuel were most likely the reason for stunted bacterial growth. While general corrosion of steel coupons was noted in all incubations the highest level of localized attack was observed in cultures augmented with F76.
Background

Pectobacterium spp. and Dickeya spp. are necrotrophic bacterial pathogens of many important crops worldwide. The effective strategies to control pectinolytic bacteria have not yet been developed. Consequently, the management of Pectobacterium spp. and Dickeya spp. is based mainly on the exclusion of infected plant material and the use of hygienic practices during potato cultivation and storage.

Objectives

This study reports on the isolation and characterization of broad host lytic bacteriophages able to infect the dominant Pectobacterium spp. and Dickeya spp. affecting potato in Europe viz. Pectobacterium carotovorum subsp. carotovorum (Pcc), P. wasabiae (Pwa) and Dickeya solani (Dso) with the objective to assess their potential as biological disease control agents.

Methods

The lytic bacteriophages were isolated from potato samples collected in different potato fields in Poland. Transmission electron microscopy was used to study bacteriophage morphology. The phages were characterized for optimal multiplicity of infection, the rate of adsorption to the bacterial cells, the latent period and the burst size. They were also genotypically characterized with RAPD-PCR and RFLP techniques. The structural proteomes of both phages were obtained by fractionation of phage proteins by SDS-PAGE. Phage protein identification was performed by liquid chromatography-mass spectrometry (LC-MS) analysis. Pulsed-field gel electrophoresis (PFGE), genome sequencing and comparative genome analysis were used to gain knowledge of the length, organization and function of the (phi)PD10.3 and (phi)PD23.1 genomes.

Conclusions

This is the first study dealing with isolation and characterization of broad host lytic bacteriophages able to infect dominant Pectobacterium and Dickeya spp. in potato in Europe.
Background

Deinococcus bacteria tolerate very high doses of gamma and UV radiation and other massive DNA damage- and oxidative stress-generating conditions.

Objectives

To better understand the mechanisms underlying extreme radioresistance, we characterized the transcriptome of Deinococcus deserti. Furthermore, we investigated how IrrE, a protein essential for radioresistance, is involved in upregulation of various genes following irradiation.

Methods

Differential RNA sequencing was performed, allowing genome-wide identification of transcription start sites (TSSs). The function of IrrE, which contains a conserved domain of unknown function (COG2856/DUF955), was analysed using several genetic and biochemical experiments.

Conclusions

Strikingly, 60% of the identified mRNAs in D. deserti is leaderless, with the TSS exactly at or very close to the translation initiation codon (1). Such proportion of leaderless mRNAs is unprecedented for a bacterial species. Numerous additional leaderless mRNAs predicted to encode small peptides were identified, providing a new explanation for the accumulation in the cell of small peptides important for protection of proteins against oxidation and thus for radioresistance. Several novel highly radiation-induced genes were also found. The mechanism by which IrrE is involved in upregulation of DNA repair and other genes has remained unknown for many years. We have now demonstrated that IrrE is a metalloprotease that cleaves and inactivates repressor protein DdrO (2). In D. deserti, this cleavage is stimulated by irradiation.

NEXT GENERATION SEQUENCING FOR THE IDENTIFICATION OF FILAMENTOUS BACTERIA IN INDUSTRIAL ACTIVATED SLUDGE PLANTS

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Background
Filamentous bacteria are mostly identified in activated sludge. Under specific conditions these organisms can outcompete other bacteria inducing process disturbances underlining the importance of a better comprehension and understanding on the growth of these bacteria to develop effective control measures.

Objectives
Identification of filamentous bacteria present in industrial WWTPs and evaluation of influencing factors on the growth of these bacteria.

Methods

Activated sludge samples from two industrial WWTPs in Germany were collected.

DNA-sequencing based PCR-product from all bacteria primer (104F, 515R) was conducted via Illumina platform.

Sequencing data was generated using a developed bioinformatics pipeline and it was further analyzed by BLAST.

Biodiversity analyses using MEGA 5.1 and R studio 3.1.0 were performed.

Conclusions
In total 432 different gene sequences were identified showing 17 sequences related to bulking and foaming bacteria as described in Guo and Zhang (2012).

The abundance of filamentous bacteria was calculated as the percentage of total reads. Chryseobacterium sp. Iso-52 (24.82 %), Candidatus Microthrix parvicella (M. parvicella) clone OTU-5-40m.ABB (21.77 %), Chloroflexi bacterium ET1 (15.61 %) and Gordonia sp. YIM 100324 (11.67 %) were the most dominant filamentous organisms found.

Phylogenetic analysis revealed a close relationship of M. parvicella and Gordonia.

Canonical correspondence analysis (CCA) indicated positive correlations to sludge age and wastewater influent for M. parvicella and Gordonia, while dissolved oxygen (DO), effluent phosphorous and ammonia concentration, nitrate sludge loading showed a negative impact. It was also observed that the occurrence of M. parvicella and Gordonia is connected to high sludge volume index (SVI) and floating sludge
fraction (AVA) values.
Background

Pig manure is usually stored in pits before spreading on agricultural land. It may also be biologically treated, leading to a liquid fraction stored in a lagoon (lagoon effluent). The impact of the storage on the persistence of pathogens depends on environmental factors and on intrinsic factors of the pathogens including their ability to enter into a viable but non cultivable (VBNC) state.

Objectives

This study aims to compare the behaviour of *Listeria monocytogenes* which can enter VBNC state, in pig manure and in lagoon microcosms.

Methods

Two strains of *L. monocytogenes* belonging to different serogroups were isolated from two farms. They were inoculated in flasks containing two manures and two lagoon effluents stored at 8 and 20°C during 63 days. The bacteria were quantified by cultural method, qPCR and qPCR associated with Propidium MonoAzide which quantifies both cultural and VBNC cells.

Conclusions

Regardless their origin and their serogroup, both strains showed similar kinetics of decrease. The lower temperature increased the persistence of the strains. An effect of the matrix was observed not only between manures and lagoon effluents but also within a same type of matrix with similar chemical composition, suggesting that biotic factors may play a role in the persistence of *L. monocytogenes*. The VBNC appeared within the first hours of contact regardless the matrix. Nevertheless, their proportion increased over time in the effluent lagoon.

This study highlights that lagoon effluents used for watering plants, creates more favorable conditions than manure for VBNC cells of *L. monocytogenes*. 
DENITRIFICATION AND NITROUS OXIDE PRODUCTION IN SOILS IN THE HAMPSHIRE RIVER AVON CATCHMENT IN THE UNITED KINGDOM

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Background

Microbial denitrification can lead to a loss of N from soils to atmosphere as N\textsubscript{2} or N\textsubscript{2}O. However, the magnitude of the soil N sink and environmental factors regulating denitrification are essentially unknown in UK catchments.

Objectives

This study aims to examine seasonal and spatial variation of denitrification and N\textsubscript{2}O production and environmental factors regulating the processes and constrained the magnitude of N loss by denitrification.

Methods

Rates of denitrification and N\textsubscript{2}O production were measured in the laboratory by \textsuperscript{15}N isotope enrichment technique in grassland soils in three sub-catchments with contrasting geologies (clay, greensand and chalk) in the River Avon catchment at four seasons from August 2013 to May 2014.

Conclusions

Denitrification and N\textsubscript{2}O production varied seasonally and spatially with annual mean rates of denitrification of 0.025, 0.265, and 0.210 \mu g N.g\textsuperscript{-1} dry soil.day\textsuperscript{-1} respectively, and of N\textsubscript{2}O production of 0.870, 0.547 and 0.620 ng N.g\textsuperscript{-1} dry soil.day\textsuperscript{-1} respectively at clay, Greensand and chalk sites. Denitrification rates were higher in August and November and higher rates of N\textsubscript{2}O production were observed in August and May.

Rates of denitrification and N\textsubscript{2}O production, measured at four temperatures (4, 10, 20 and 30°C) and three levels of soil water concentration (57.5%, 77.5% and water saturation), increased with temperature and soil water content.

The dominating environmental factors for denitrification and N\textsubscript{2}O production were soil pH, soil water content and temperature. It is estimated that annual losses of N by
denitrification were 11.8, 125.8 and 99.7 kg N per hectare at clay, greensand and chalk sites respectively.
EFFECT OF ENERGY CROP CULTIVATION AND FERTILIZATION ON SOIL ARCHAEAAL COMMUNITY AND ITS POTENTIAL OF METHANOGENESIS IN ABANDONED PEAT EXTRACTION AREA

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Background

Peat is an important resource and is being extracted widely from regions rich in peatlands, which inevitably leads to the problem of abandoned peatlands. These areas emit greenhouse gases (CH₄, N₂O, CO₂) for decades if restoration action is not implemented. The potential of methane production of abandoned peatlands can be evaluated by the abundance of mcrA genes which encodes the alpha-subunit of the enzyme that catalyses the last step in CH₄ synthesis in all methanogens.

Objectives

The purpose of this study is to assess the effect of energy crop (Phalaris arundinacea L.) cultivation and fertilisation on the abundance of soil bacterial and archaeal 16S rRNA genes, and to evaluate the methanogenic potential of the microbial communities in the soils of the experimental plots on the abandoned peat extraction area.

Methods

The experiment was carried out in 2012-2014 and samples were collected from 12 experimental plots at the vegetation period. Besides the widely used statistical methods, linear mixed effects modelling was used to test relationships between gene parameter values and chemical variables and in cases of different grouping factors.

Conclusions

Various physicochemical parameters changed the abundance of peat methanogenic community in vertical profiles. Methane emission rates correlated with the abundance of peat deeper layers methanogenic community in uncultivated plots while without Phalaris cultivation there are less constraining factors. These findings are important in the assessment of global warming potential while using abandoned peat extraction areas to produce biomass for energy.
Background

Release of effluents from textile dyeing and printing industries prior to proper treatment is a vital source polluting water and soil. A number of structurally dissimilar industrial azo dyes are present in textile effluents.

Objectives

To isolate, screen and characterize a bacterial isolate having potential to decolorize and degrade industrial textile dyes from textile effluent. To investigate the effects of various environmental and nutritional parameters on decolorization activity of that isolate on a specific dye under lab condition. To study phytotoxic effects caused by decolorized and degraded dye products.

Methods

Decolorization and degradation study was of selected dye was performed using UV Visible spectroscopy and HPLC analysis. Germination assay of seeds of some selected crops was performed to investigate the toxic effects of decolorized and degraded dye products.

Conclusions

We were able to screen a bacterial isolate with decolorization and degradation potential from the textile effluents. The complete decolorization of dye (150 mgL-1) was achieved within 24 hrs of incubation under static culture condition (at pH7, 37°C). The maximum decolorization was found in presence of Yeast extract as a source of carbon (at pH7±0.2, 37°C) Static culture condition. Seeds irrigated with decolorized dye sample (under static culture condition) show inhibition in germination. Inhibition of seed germination was not observed in case of seeds irrigated using dye treated under sequentially static-shaking culture conditions did not show.
Background

Microbial communities show the greatest organisms diversity on earth. Culture-independent molecular approaches targeting ssu ribosomal RNA genes have revealed this extraordinary diversity. However, amplicons cannot establish the link between microbial communities structure and realized metabolic functions, limiting the comprehension of micro-organisms roles. Despite the advent of the current ultra-high throughput sequencing, efficient assembly of sequencing data from metagenomic samples remains difficult and complete genome reconstruction has been principally realized for dominant micro-organisms. Single cell sequencing strategies have also been developed to overcome these limitations and to access less abundant micro-organisms but are not always easily practicable.

Objectives

Based on a solution hybrid selection method combined with next generation sequencing, we developed an innovative gene capture approach to enable the reconstruction of large genomic regions or even complete microbial genomes from complex environments.

Methods

A first gene capture using highly specific short probes targeting 16S rRNA gene was applied on metagenomic DNA to obtain large specific probes (several kbp) spanning the biomarker and the unknown flanking regions of the microbial species of interest, *Roseobacter denitrificans*. Generated long probes were then used to capture very large DNA fragments of *R. denitrificans* within a mix of bacterial gDNA. Real-time PCR performed on captured DNA revealed a significant enrichment in targeted 16S rRNA gene, surrounding genes and even several Mbp away genes, suggesting the capture of *R. denitrificans* complete chromosome.

Conclusions

Sequencing of captured DNA allows identification of unknown sequences and reveals new genetic associations. This promising approach will facilitate linking identity and associated functions.
THE IMPACT OF SEPTIC SYSTEMS AND WATERSHED LEVEL CHARACTERISTICS ON STREAM FAECAL POLLUTION IN SUBURBAN WATERSHEDS IN GEORGIA, USA

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Background

A quarter of housing units in the USA use septic systems for wastewater treatment and disposal. As widely as septic systems are used, their watershed level impact on water quality is poorly understood.

Objectives

The objective of this study was to determine the impact of septic systems on water quality and in so doing identify watershed level characteristics that influence their impact.

Methods

Water samples, collected from 24 well characterized watersheds with varying septic density, were analyzed for fecal indicator bacteria to quantify pollution levels and identify seasonal trends. The watersheds represent a gradient of land-use conditions from low to high density of septic systems, as well as developed to undeveloped uses. Pollutant sources were then tracked using Bacteroides sp. genetic markers.

Conclusions

Our findings indicated statistically significant interaction between septic density and season for enterococci count (P = 0.005) and stream yield (P = 0.04). Seasonal variations in bacterial count and stream yield were also observed, with significant differences between spring–winter and summer–winter. Multiple linear regression models suggested that 4 watershed characteristics (septic density, median distance of septic systems to stream, % developed area and forest cover) and water temperature could explain about half of the variability in bacterial count and yield in spring and summer. There was a significantly positive correlation (R = 0.67) between septic density above 77 units/km² and human-specific marker yield (indicator of septic contribution). Overall study conclusion was that septic density strongly influences fecal pollution but its impact is dependent on season and watershed characteristics.
ANAEROBIC DEHALOGENATING BACTERIA RELATED TO DESULFOLUNA SPONGIIPHILA FORM A COSMOPOLITAN GROUP WIDELY DISTRIBUTED IN ORGANOHALIDE-CONTAINING SPONGES

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Background
The marine environment is a rich source of biogenic organohalides produced by a diversity of marine organisms, such as sponges that produce a vast array of halogenated bioactive compounds as secondary metabolites. These organohalogen compounds in turn appear to select for bacteria that can utilize them as a source of energy.

Objectives
Our objectives are to investigate the dehalogenating bacterial populations within the sponge animal with the overall goal of understanding the roles and metabolic activities of this endomesohyl microbiota.

Methods
We enriched for bromophenol dehalogenating bacterial symbionts from a range of sponge species collected at different sites in the Mediterranean, Atlantic and Pacific, and applied an integrated cultivation and molecular analysis-based approach to analyze the bacterial communities.

Conclusions
We demonstrated the widespread distribution of anaerobic reductive debrominating bacteria in different marine sponges. We isolated a novel bacterial species, \textit{Desulfoluna spongiiphila}, from Mediterranean sponges that grows by respiratory reductive dehalogenation. Using a cultivation and molecular analysis-based approach we demonstrated that \textit{D. spongiiphila} and its close relatives form a cosmopolitan group widely distributed in organohalide-containing sponges across different geographic locations. The sponge-associated dehalogenating bacteria can operate \textit{in vivo} and impact the fate of brominated organics. This new bacterial species group is an excellent model system to study a "chemically-driven" endosymbiotic relationship and one possible origin of organohalogen respiration. Organobromine-rich sponges
appear to provide a specialized, possibly ancient, habitat for organohalide-respiring microbes, which mediate a cycling of organohalide compounds within the sponge animal.
Background

Quantitatively methanesulfonate (MSA) is very relevant in the global biogeochemical sulfur cycle. MSA utilization by bacteria as a source of carbon and energy has been described and a specific enzyme, methanesulfonate monooxygenase (MSAMO), has been found to perform the first catabolic step of its oxidation. Other proteins, seemingly involved in the import of MSA into bacterial cells, have been reported.

Objectives

In this work we looked for the genes encoding the large subunit of the MSAMO (msmA) and the periplasmic component of the import system (msmE) in the genomes of MSA-enriched bacteria and also in metagenomic DNA from seawater, in order to extend our knowledge on these genes as ecofunctional genetic markers of MSA degradation.

Methods

PCR approaches were implemented and the resulting products were cloned and sequenced. Two novel marine MSA-isolates were also submitted to whole genome sequencing.

Conclusions

In conclusion, novel msmA and msmE sequences were obtained, and two full msm operons were found in the genomes of the novel marine strains. Clearly, the msmA genes (and derived proteins) show less sequence variability than msmE. This, added to the apparent conservation of a peculiarly long Rieske-associated motif, reinforces the value of msmA as ecofunctional indicator for methanesulfonate cycling by bacterial natural communities. In the line of previous observations, our results also show that high-GC strains are somehow favoured during enrichment and isolation of MSA-utilizing bacteria, illustrating the misrepresentation of natural populations that
culturing may entail.
Background
Heavy metal pollution is of great concern in countries like Pakistan where a huge proportion of human population is exposed to it. Bioremediation is one of the most viable and efficient solution for the problem.

Objectives
Purpose of the current study was to isolate endophytic fungi from corn plants and screen them for phytostimulation under heavy metal stress.

Methods
Endophytic strain (MHR-23) was identified as *Paecilomyces formosus* by 28 S rDNA and internal transcribed spacer (ITS) sequence homology. The strain effectively tolerated up to 2000 µg mL\(^{-1}\) of these heavy metals. Adverse effects of metals were obvious in the form of gradual decrease of fungal growth by increasing the concentration of heavy metals. The strain was able to remove 60-87 % of metals (300 µg mL\(^{-1}\)) from PDA broth. The strain removed metals by biotransformation and/or accumulation of heavy metal in its hyphae. Growth of corn stressed with heavy metals was restored by the application of MHR-23. Accumulation of heavy metals by corn stem and leaves was significantly reduced (p < 0.05) by the application of MHR-23.

Conclusions
It may be concluded that MHR-23 is an excellent candidate to be used as biofertilizer in fields affected with heavy metals.
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GENOME DATA MINING IN DETERMINATION OF GENES INVOLVED IN THE ANTIMICROBIAL ACTIVITY OF PSEUDOMONAS SP. P482 AGAINST PLANT PATHOGENIC BACTERIA AND FUNGI
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Background
To survive in a competitive environment, the plant-associated Pseudomonads produce a set of biologically active compounds, many of them known for their antimicrobial properties. Pseudomonas sp. P482, a tomato rhizosphere isolate, shows antagonistic activity towards a spectrum of plant pathogenic bacteria and fungi.

Objectives
In this study, we aimed to apply a genome data mining approach to determine the genetic background of the antibacterial activity of the P482 strain.

Methods
The genome sequencing of P482 was performed. Bioinformatics tools such as the antiSMASH and BAGEL3 were employed to analyze the resulting data and to determine the ‘candidate’ genes potentially involved in the antimicrobial activity of the P482. The ‘candidate’ genes were subjected to site directed mutagenesis with the use of a suicide vector. The obtained mutants were tested for the lack of antimicrobial activity towards selected strains of plant pathogenic bacteria and fungi.

Conclusions
A draft genome sequence of P482 was obtained and automatically annotated. The BAGEL3 analysis yielded no putative bacteriocins. The antiSMASH analysis revealed 5 biosynthetic clusters encoding biosynthetic genes potentially involved in the synthesis of antimicrobial factors. Knock out of the 5 major biosynthetic genes, one per each predicted cluster, has shown that these genes are not essential for the antimicrobial activity of P482 against the soft rot bacteria. Thus, genome mining is a powerful approach yet when novel mechanism of antagonism are at stake, as in seems to be in case of the P482, it should be used in combination with alternative methods.
ELUCIDATING THE GENETIC BASIS FOR SULFONAMIDE DEGRADATION IN ISOLATES WITH DIVERSE PHYLOGENETIC BACKGROUND

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Background

Sulfonamides antibiotics are among the most frequently prescribed antibiotics in human and veterinary medicine, and, at the same time quite recalcitrant to degradation. Therefore, sulfonamides can be commonly detected in wastewater worldwide in trace amounts. Up to now, few sulfonamide degrading bacteria stemming from different phyla have been isolated. While the degradation mechanisms in most strains have not been conclusively elucidated yet, most cases were shown to lead to analogous metabolites

Objectives

The intention of this study is to understand how sulfonamide degradation genes are propagated in the environment and whether the isolated sulfonamide strains have in common a similar apparatus of genes encoding degradative enzymes.

Methods

A set of draft genomes of sulfonamide degrading bacteria was obtained by next generation sequencing of genomic DNA of these strains and from public available data sets. The nucleotide sequence of a gene cluster related to sulfonamide degradation in Microbacterium sp. BR1 was compared to similar regions found in other strains.

Conclusions

It was found that some sulfonamide degraders contained in their genome DNA regions highly similar to that found to be responsible for degradation in Microbacterium sp. strain BR1, while for others, similarities to this region were too low to safely attribute the same function to the encoded enzymes. It remains to be solved
whether these distantly related genes confer degrading activity in these strains or whether completely different enzymes are involved.
ROLE OF VIBRIO CHOLERAE ADHESINS IN SURVIVAL OF PREDATION BY HETEROTROPHIC PROTISTS

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Background
Vibrio cholerae is an opportunistic pathogen and a natural inhabitant of aquatic ecosystems. Predation by heterotrophic protozoa has a major impact on the environmental survival of V. cholerae. Adhesins are important virulence factors required for V. cholerae pathogenesis but may have evolved during long-term interactions with protozoa.

Objectives
Here, we investigated the role of V. cholerae adhesions (TcpA, MshA, GbpA, LapA and VC1804) in resistance to predation by the surface-feeding Acanthamoeba castellanii.

Methods
V. cholerae wild type stains and adhesin mutants were exposed to predation by A. castellanii. The grazing resistance was determined by V. cholerae selective grazing assays, and quantification of V. cholerae cells attached to the surface and/or internalized by A. castellanii as assessed by CFUs and microscopy.

Conclusions
Data show that only VC1804 made a significant contribution to grazing resistance as the mutant was grazed more than the parent wild type. Compared to wild type strains, the MshA and VC1804 mutants exhibited significantly lower colonization efficiency for the A. castellanii surface, which also resulted in less cells surviving intracellularly in the amoebae. These findings further elucidate the role of V. cholerae adhesins in persistence in the environment and support the hypothesis that some virulence factors provide for increased fitness in the environment.
THE VALUE OF P. AERUGINOSA IN HEALTH AND MICROBIOLOGICAL QUALITY CONTROL OF RECREATIONAL WATERS USING IN PUBLIC SWIMMING POOLS
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Background
Pseudomonas aeruginosa is the most significant bacteria capable of multiplying in water especially in recreational waters. Nowadays most public swimming pools use recreational water.

Objectives
To assess if these swimming pools are health risk to users, eleven of public swimming pools in East and North-East of Tehran were monitored. Useful information such as flow rate of filtration of water in the pools, clarity, temperature and pH of the water were recorded at the same time of sampling too.

Methods
According to the Standard Method 21th edition, Bacteriological tests such as isolation, identification and colony counts of Escherichia coli, coliforms, and Pseudomonas aeruginosa were performed on the samples. P. aeruginosa was isolated from 9(81.8%) of the pools. Also P. aeruginosa was the only predominant organism isolated from 7(63.6%) of the pools, while in the remained 2(18.2%) other, in addition to P. aeruginosa, high rates of total bacterial count, total coliforms and faecal coliform counts were also found.

Conclusions
Results of the contaminated swimming pools revealed that chlorination was often inadequate to disinfect mentioned bacteria especially P. aeruginosa when high population of people had to over use of the pools. Thus results of this study, showed the importance and the value of assaying presence of indicator of P. aeruginosa routinely in quality control of recreational waters using in public swimming pools.
THE EFFECT OF NUTRIENTS ON CARBON AND NITROGEN FIXATION BY THE UCYN-A-HAPTOPHYTE SYMBIOSIS
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Background
Symbioses between phytoplankton and N₂-fixing microorganisms play a pivotal role in the ocean's carbon and nitrogen cycle. The abundant and widespread unicellular cyanobacteria group A (UCYN-A) was recently found to live symbiotically with a haptophyte.

Objectives
Here, we investigated the effect of nitrogen (N), phosphorus (P), iron (Fe) and Saharan dust additions on nitrogen (N₂) fixation and primary production by the UCYN-A-haptophyte association in the subtropical eastern North Atlantic Ocean.

Methods
In order to determine single-cell rates of N₂ fixation and primary productivity, we used stable isotope incubations combined with nanoSIMS measurements. nifH gene expression analysis was carried out to determine the molecular regulation of N₂ fixation under the different nutrient conditions.

Conclusions
N₂ fixation by UCYN-A was stimulated by the addition of Fe and Saharan dust whereas CO₂ fixation by the haptophyte was stimulated by the addition of ammonium nitrate as well as Fe and Saharan dust. However, there were no detectable differences in nifH gene expression in any of the incubations. Independent of the nutrient treatment, we observed a mutual exchange of carbon and nitrogen between the UCYN-A and the haptophyte suggesting a tight coupling. However, it appears that the transfer of carbon from the haptophyte to UCYN-A requires a transfer of nitrogen from UCYN-A indicating an obligate symbiosis of this globally important diazotrophic association.
RESPONSE OF GEOLOGICAL RESIDENT MICROBIAL COMMUNITIES TO ADDITIONS OF NITRATES AND/OR ACETATE, MIMICKING LEACHATES OF BITUMINIZED INTERMEDIATE-LEVEL RADIOACTIVE WASTE

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Background
Clay deposits such as the Opalinus Clay formation are studied as host rocks for geological disposal of high- and intermediate-level long-lived radioactive waste in several European countries. Bituminized intermediate-level long-lived radioactive waste contains, besides bitumen and radionuclides, also large amounts of organics, nitrates and sulphates. Over time, these salts will dissolve, leach and diffuse into the surrounding clay host rock, together with water soluble organic substances. To mimic the potential effect of such inorganic-and organic leachates, pulses of nitrate and/or acetate have been injected in intervals of an in situ experiment, called Bitumen-Nitrate-Clay interaction (BN) experiment, running at the Mont Terri geological laboratory (St. Ursanne, Switzerland).

Objectives
To elucidate whether the microbial communities, present in the BN experiment, are affected or involved in the observed biogeochemical changes.

Methods
Beside classical microbial analyses, at pivotal moments (i.e. before, during and after these pulse injection tests), priority was given to DNA-based molecular biology analysis methods, as these methods provide very accurate information on the composition, the metabolic capacity and possible evolution of bacterial communities in response to the nitrate and/or acetate injections.

Conclusions
The changes observed in the bacterial populations appeared to correlate well with the imposed physico-chemical changes. As soon as nitrate was added an overwhelming community shift appeared to nitrate reducing bacteria. If in parallel acetate was offered, the community composition did not alter that much, but the speed of nitrate reduction was increased twentyfold. This high nitrate removal speed was maintained as long as the easily consumable carbon source was abundant.
FUNCTIONAL CHARACTERIZATION OF METAGENOMES FROM MICROBIAL COMMUNITIES IN PRISTINE AND CHRONICALLY-POLLUTED HARBOR WATERS IN RELATION TO HYDROCARBON POLLUTION

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Background
The phylogenetic composition of bacterial communities in chronically-polluted harbor waters in Mallorca Island (Spain) has been shown to differ from those in nearby clean waters, with a characteristic predominance of bacteria from the Roseobacter clade (Nogales et al., 2007). We have also observed that bacteria of this group proliferate after artificial pollution with diesel oil (Lanfranconi et al. 2010).

Objectives
Given these facts, and the evidences relating roseobacters with aromatic hydrocarbon degradation, our aim was to explore the functional diversity of bacteria in polluted harbor waters and waters polluted artificially, with special attention to genes for the catabolism of hydrocarbons, and relate it with the presence of members of the Roseobacter clade.

Methods
Eight metagenomes have been generated (20.6 Gb in total) using Illumina technology. Sequences filtered using quality criteria are analyzed using MG-Rast (Meyer et al., 2008), and in house assembly and annotation.

Conclusions
Phylogenetic affiliation of sequence reads confirmed the abundance of roseobacters in the metagenomes from polluted samples. Genes coding for peripheral and central pathways for aromatic hydrocarbon degradation have been observed in all metagenomes analyzed. However, at a broad level of resolution we cannot evidence significant differences in number of reads between metagenomes from polluted and control samples. Recruitment of sequence reads against protein databases from Roseobacter genomes and for catabolism of hydrocarbons would help us to define the fine catabolic differences between polluted and unpolluted samples.

BIOFLOCCULANT PRODUCTION BY ACTINOBACTERIA CONSORTIA AND MEDIA OPTIMIZATION THROUGH RESPONSE SURFACE MODEL

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Background
Flocculants aggregates suspended particles in solutions thus, reducing turbidity. The industrial application of this process includes water/wastewater treatment. Conventionally used flocculants have been implicated in deleterious health problems hence, the need for a safe alternative. Bioflocculants are valuable alternative as they are non-toxic and biodegradable.

Objectives
The consortia of \textit{Brachybacterium}, \textit{Cellulomonas} and \textit{Streptomyces} were evaluated for biofloculant production and their nutritional requirements optimized via response surface model.

Methods
\textit{Cellulomonas}, \textit{Streptomyces} and \textit{Brachybacterium} species were evaluated in pair wise consortium for enhanced production of bioflocculant. Bioflocculant production, flocculation activity testing and media optimization through response surface model were in accordance with standard methods.

Conclusions
Nutritional preferences for bioflocculant production were: glucose (56%; 2.78±0.15 g/l), (NH\textsubscript{4})\textsubscript{2}NO\textsubscript{3} (53%; 2.81±0.37 g/l) and CaSO\textsubscript{4}.H\textsubscript{2}O (47%; 2.19±0.13 g/l) against \textit{Brachybacterium} and \textit{Streptomyces} species, glucose (92%; 4.06±0.62 g/l), peptone (89%; 3.88 ± 0.432 g/l) and MgCl\textsubscript{2} (83%; 3.99±0.79 g/l) against \textit{Cellulomonas} and \textit{Brachybacterium} species. Lastly, sucrose (91%; 4.01±0.22 g/L), peptone (82%; 3.21±0.13 g/L) and magnesium chloride (78%; 3.29±0.42 g/L) were preferred by the consortium of \textit{Streptomyces} and \textit{Cellulomonas} species. Plackett-Burman design revealed that the carbon, nitrogen and cation sources were the critical media components essential for the production of bioflocculant. High regression coefficients respectively obtained for the consortia was indicative of the model adequacy (P ≤ 0.05). The bioflocculant compositional analysis and functional group determination (FTIR spectrometry) indicates a heteropolysaccharides bioflocculant while SEM imaging revealed an amorphous loosely arranged fluffy structure with interstice less than 1µm. The high flocculation activity shown suggests suitability for industrial processes.
THE INFLUENCE OF AI-2 MOLECULES ON RHAMNOLIPIDS SECRETION EXHIBITED BY PSEUDOMONAS AERUGINOSA WITHIN MICROBIAL COMMUNITY

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Background

Rhamnolipids are a group of biosurfactants produced by Pseudomonas species. It was proven that rhamnolipids participate in biofilm formation and play an important role in microbial motility. Because of their biodegradability and biocompatibility, they are also considered as promising alternatives to widely used synthetic agents in surfactant-mediated bioaugmentation. The cooperation and competition between indigenous and exogenous microorganisms plays an essential role in the final bioaugmentation effect and is based on quorum sensing (QS) mechanism, which enables cell to cell communication. The autoinducers 2 (AI-2) are QS signal molecules observed in both Gram-positive and Gram-negative bacteria. We hypothesize that AI-2 molecules play important role in secretion of rhamnolipids by Pseudomonas aeruginosa within microbial community.

Objectives

1. To isolate biosurfactant-produced bacteria and investigate the structures of secreted compounds.

2. To investigate the relationship between AI-2 activity and secretion of rhamnolipids.

Methods

Isolation: samples originating from cow, chicken and pig feces; screening of bacterial communities by: (i) oil spreading, (ii) drop collapse, (iii) hemolytic activity, (iv) emulsification activity; bioreactor cultivation: Sartorius Stedim, Germany; surface tension: Krüss, Germany; bioluminescence assay: Vibrio harveyi BB170 and Vibrio harveyi BB152; rhamnolipids identification: HPLC-MS.
Conclusions

Only monorhamnolipids were detected in the consortium and Pseudomonas aeruginosa supernatants. The maximal rhamnolipids production by Pseudomonas aeruginosa and Al-2 activity were observed at the same time. The results indicate that QS molecules secreted by associated microorganisms have significant influence on Pseudomonas aeruginosa gene expression responsible for rhamnolipids production.

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THE INFLUENCE OF ZEROVALENT IRON ON THE DEGRADATION OF HEXABROMOCYCLODODECANE BY MICROCOSM FROM RIVER SEDIMENT

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Background
Brominated flame retardants (BFRs) have been widely used in many industries over decades. Hexabromocyclododecane (HBCD) is one of the most commonly used BFRs, but it has just listed on the table of persistent organic pollutants (POPs).

Objectives
In order to develop cost-effective remediation methods for this emerging pollutant, we evaluated the HBCD degradation ability of one anaerobic microcosm from river sediment, as well as the influence of zerovalent iron (MZVI) on the biodegradation.

Methods
An anaerobic sediment sample was acclimated by HBCD. The supernatant solution which contained microcosm was used to conduct further experiments. MZVI (2.5 g L⁻¹) was used to degrade HBCD alone or together with the microcosm.

Conclusions
The microcosm started to degrade HBCD at day 7, and 30% of HBCD was eliminated in the following 4 days. Sixty percent of HBCD was degraded by MZVI or the co-incubated system after 11 days of incubation. The HBCD biodegradation ability was inhibited by MZVI. MZVI degraded HBCD through reductive debromination, however, this reaction did not occur in the biodegradation setting, suggesting that the microcosm eliminated HBCD through other mechanism. The fluctuation of bacterial community during the degradation processes was monitored. Cluster analysis shown that the microbial composition was less close to that of the MZVI added ones. The inhibition of biodegradation from MZVI may due to the alteration of medium characteristics, which in term changed the microbial composition and function. Our studies provided an insight into the influence of MZVI on the biodegradation of HBCD, which would facilitate the bioremediation of BFRs.
BACTERIA ISOLATED FROM DIATOM BLOOMS SEA WATER ADAPT AND GROW IN THE PRESENCE OF POLYUNSATURATED ALDEHYDES

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Background

Diatoms commonly grow in association with heterotrophic bacteria (Amin et al., 2012) and some species produce polyunsaturated aldehydes (PUAs), with antimicrobial activity (Ianora et al., 2004). Therefore, diatom-associated bacteria may have evolved resistance to PUAs (Ribalet et al., 2008; Balestra et al., 2011), possibly involving adaptive mechanisms of membrane fatty acids (Heipieper and de Bont, 1994; Heipieper et al., 2007).

Objectives

The objectives of this study were the isolation of PUAs-resistant bacterial strains from sea water during diatom blooms and the investigation of bacterial adaptive mechanisms to PUAs.

Methods

Enrichment cultures in mineral medium with PUAs \(\text{\(2E,4E/Z\)-heptadienal, \(2E,4E/Z\)-octadienal, \(2E,4E/Z\)-decadienal}\) were inoculated with sea water and incubated at 22°C. PUA-resistant bacteria were isolated, identified by 16S rDNA, grown in the presence of the PUAs and analyzed at membrane fatty acids level.

Conclusions

Six PUAs-resistant isolates, i.e. 1A, 5a, 5b, 7a, 11 and 12b, were assigned to the genera \textit{Pseudomonas}, \textit{Sufflavibacter}, \textit{Halomonas}, \textit{Vibrio}, \textit{Idiomarina} and \textit{Labrenzia}, respectively. Strains 1A, 5b and 7a showed higher growth in the presence of PUAs with respect to the others. Preliminary data on adaptive responses to the PUAs on membrane fatty acid level revealed an increased saturation.

References

INVESTIGATION OF BIOTIC AND ABIOTIC FACTORS THAT AFFECT SURVIVAL OF THE PATHOGENIC BACTERIUM LISTERIA MONOCYTOGENES IN SOIL

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Background

Agroecosystems are dedicated to the provision of high quality, safe food and sustainability is a major issue. Biological contamination of the food supply raises food safety issues. Prevention of contamination of the food supply requires a thorough understanding of the ecology of pathogenic microorganisms. Listeria monocytogenes is a food pathogen found in many habitats including soil, plants, water systems and food processing factories.

Objectives

We designed microcosm experiments to decipher extrinsic factors that drive the fate of this model of Human pathogen in soil.

Methods

We analysed population dynamics over 80 days after inoculation of a set of 100 soil microcosms. Population dynamics in unsterilised and sterilised soil microcosms were compared. Detailed information such as soil characteristics and land use were integrated into an analysis of variance.

Conclusions

The population of L. monocytogenes decreased over time but survival depended on the soil under scrutiny. Chemical properties and soil texture affected survival of L. monocytogenes. The endogenous soil microbiota had a major impact on the decrease of the population of L. monocytogenes. This prompted us to investigate the relationship between the biodiversity of the soil microbiota and the inhibition of the population of L. monocytogenes. We investigated the consequences of soil diversity erosion on the fate of a human bacterial pathogen in the telluric environment. Diversity analysis showed that both soil diversity and phylogenetic composition affected survival of L. monocytogenes. Overall, our results demonstrate that soil biodiversity is a major driver that limit invasion of soil by pathogens.
Background
The treatment of wastewater, before their dumping to the aquatic ecosystems, constitutes a topic of current importance.

Objectives
The objective of the research was to evaluate the physiologic and metabolic diversity of the rhizobacteria from hydrophyte plants of natural wetlands.

Methods
Individual bacterial strains were tested in front of different pollutants. Microbial consortia were faced to water simulating a domestic effluent and synthetic water similar to an industrial effluent. This allowed the selection of the autochthonous strains Bacillus sp. (T-119, T-117 and T-229), Acinetobacter sp. (T-118), Pseudomonas sp. (T-1111) and Exiguobacterium sp. (T-316) for their capacities in the elimination of organic matter, ammonium and phosphate. Also, the strains Bacillus sp. (T-119, T-117, T-1115 and T-1113) were selected for the lead, chromium and mercury biosorption from synthetic residual waters. Two consortia designed with these strains achieved 75%, 85% and 100% removal of organic matter, ammonium and phosphate and 70%, 9% and 17% of lead, chromium and mercury, respectively. The effect of factors associated to the process of pollutants elimination was determined, which it contributed to the design and bioaugmentation with the consortia of artificial wetlands to laboratory scale. With these systems were reached 100%, over 70% and 55% removal of organic matter, ammonium and phosphate and elimination levels over 95%, 80% and 50% of lead, mercury and chromium, respectively.

Conclusions
It was settled down an integral strategy for obtaining effective bacterial consortia in the bioaugmentation of constructed wetlands, applied in the removal of inorganic and organic pollutants.
ON THE EVOLUTION OF WRINKLY SPREADERS, WRINKLEALITY AND FITNESS
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Background
Bacterial adaptive radiation has been extensively studied using Pseudomonas fluorescens SBW25 which gives rise to the Wrinkly Spreader (WS), a class of adaptive mutants producing a wrinkled colony morphology and air-liquid interface biofilms in static microcosms. The WS phenotype is the result of mutations targeting different diguanylate cyclases that up-regulate intracellular levels of c-di-GMP and induce biofilm-formation. Although variations in colony morphology have been noted previously, variations in the WS phenotype (wrinkleality) have not been quantified, nor related to the fitness advantage WS isolates have over other competitors in static microcosms.

Objectives
In this work, we investigated the relationship between wrinkleality and fitness using WS isolates evolved in microcosms containing different growth media.

Methods
Wrinkleality was quantified using a combined biofilm assay (measuring growth, biofilm strength and attachment levels), competitive fitness determined using the non-biofilm–forming reference strain SM-13, and data analyzed by ANOVA and fitness modeled using a GLM approach with environment (media) as the main factor and wrinklealities as covariates.

Conclusions
Significant variation in wrinkleality and fitness was observed between and within WS isolates evolved in different environments. Environment, growth, strength and attachment were all found to have significant effects on fitness. This analyses has shown that growth media selected for subtly different classes of WS, and the underlying differences in phenotype could also help to explain the variation seen in fitnesses. These findings also provide evidence to suggest that WS mutations might have pleiotropic effects and explain why some Wrinkly Spreaders are more successful than others.
ASSESSMENT OF MICROBIAL CONTAMINATION LEVEL OF SOLID WASTES FROM HOSPITALS IN UNIVERSITY OF PORT HARCOURT, NIGERIA.
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Background
Hospital solid waste disposal has become a major problem in that people indiscriminately dump this waste which pose health danger in the city and therefore there is need to assess these wastes in order to fully understand the micro organisms of health significance which could be transmitted through the improper disposal of these waste materials.

Objectives
This study was to evaluate the microbial load of solid health care waste from healthcare institutions in University of Port Harcourt. Solid waste samples were collected randomly from selected units from University of Port Harcourt teaching hospital and LULU Briggs health center

Methods
The method adopted was that of Faith Efosa Oviasogie et al.(2010), modified.

Conclusions
Bacterial species isolated from the health care solid waste include; Klebsiella, Pseudomonas, Staphylococcus, Proteus, Streptococcus While Klebsiella, Salmonella and Pseudomonas were isolated from the control samples. The fungal species isolated from the health institutions of University of Port Harcourt include Aspergillus and candida, where-as Candida was isolated from the control sample. The total heterotrophic bacterial count from all samples collected ranged from $1.8 \times 10^5$ to $6.8 \times 10^8$ Cfu/100g and total fungal count ranged from $2 \times 10^2$ to $7x10^4$ Cfu/100g. The predominant organisms were Staphylococcus aureus, Klebsiella pneumonia and Pseudomonas aeruginosa. Some of the bacterial isolates have been incriminated in nosocomial infections.
Environmental microbiology

FRESHWATER ANAEROBIC OXIDATION OF METHANE ASSOCIATED WITH SULFATE REDUCTION IN A NATURAL GAS SOURCE

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Background

Anaerobic oxidation of methane (AOM) coupled to sulfate reduction (SR) occurs in marine sediments and the responsible microorganisms are communities of anaerobic methanotrophic archaea (ANME) and sulfate reducing bacteria (SRB) of the SEEP-SRB1 clade. AOM coupled to SR is rarely documented in terrestrial environments where sulfate concentrations are generally low.

Objectives

The possible occurrence of AOM coupled to SR was investigated in a natural freshwater gas source. This was done to know where and if AOM coupled to SR occurs in terrestrial environments and which organisms execute this process.

Methods

¹³CO₂ formation and methanogenesis was monitored at long term (>168 days) in slurries incubated with ¹³CH₄ and different electron acceptors (sulfate, nitrate, ferrihydrite, humic acids). ¹³CH₄, ¹²CH₄ (produced during methanogenesis) and ¹³CO₂ were quantified using gas chromatography coupled to mass spectrometry. Electron acceptors were analyzed by ion Chromatography and iron and sulfide by colorimetric methods. Microbial community analysis was done using 16S rRNA pyrosequencing and quantification of specific microbial groups was done using quantitative PCR.

Conclusions

Slurry incubations gave evidence that AOM occurred only with sulfate as electron acceptor. Sulfide production occurred simultaneously with ¹³CO₂ production and no methanogenesis occurred. Pyrosequencing analysis showed presence of ANME-2a/b and ANME-2d 16S rRNA genes. Higher abundance of ANME2a/b in incubations with methane and sulfate was confirmed with qPCR analysis. Bacterial pyrosequencing showed presence of SRB sequences belonging to SEEP-SRB1. This is the first report...
that unequivocally shows that AOM is associated with SR in a freshwater environment.
Background
Mitigation of arsenic (As) pollution is a topical environmental issue of high R&D priority. *Pteris vittata* is a proven arsenic hyperaccumulator and has capability of remediating arsenic from water and soil through phytoextraction. There are a few reports on rhizospheric microflora of *Pteris vittata* which could probably mediate As uptake by the plants, but there are no reports of such study on this Indian ecotype of *Pteris vittata* identified by us.

Objectives
This investigation was carried out to isolate and study role of rhizospheric microbes of the Indian ecotype *Pteris vittata*.

Methods
A glasshouse experiment was conducted to study rhizosphere microbes of *P. vittata* grown in soil amended with 25 mg arsenic Kg$^{-1}$ soil in the form of sodium arsenate for sixty days. A total of 12 bacteria cultures were isolated and marked as As1-As12 and they were further studied for Minimum Inhibitory Concentration, Arsenic transformation test and Plant Growth Promoting Characteristics (PGPRs) i.e. Siderophore, IAA and ACC deaminase.

Conclusions
The isolated microbes showed arsenic tolerance up to 1000ppm and have dual capacity of AsV reduction and AsIII oxidation. Out of the twelve isolates, As 1, 2, 11, 12 exhibited siderophore positive. Except As 3, 4, 5, 9 & 12, all produced IAA while ACC deaminase was absent in all isolates. These finding show the interaction between plant and microbes for As detoxification and remediation by this ecotype of *P. vittata* endogenous to India. Our study opens up further scope of using the isolated arsenic tolerant microbes for enhancing the As remediation process through phytoremediation.
EFFECT OF ATTENDANCE ON THE SEDIMENT MICROBIAL COMMUNITIES OF 14 SANDSTONE CAVES IN THE PROTECTED LANDSCAPE AREA LABSKÉ PÍSKOVCE (CZECH REPUBLIC)

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Background

In the Protected Landscape Area Labské pískovce (Czech Republic) there is a series of sandstone caves accessible with climbing equipment prevailing; their access is subjected to permission and attendance recorded in the log-books.

Objectives

Investigation of the effect of attendance on the quantity, rough composition, stress indicators, and activities of microbial communities in cave-sediments in 14 caves using number of visitors and step-compression as the main tested factors.

Methods

Microbial communities were characterized by phospholipid fatty acid (PLFA) profiles and activities of extracellular oxidases, peroxidases, phosphatases, proteases, and glucosidases.

Conclusions

Sediment microbial communities were generally poor (low PLFA content, low activities). Step-compression increased significantly ratios of fungal/bacterial and G+/G- PLFA as well as PLFA of actinobacteria and activities of phosphatases, glucosidases, and peroxidases while decreased trans/cis PLFA indicator of microbial
stress. Number of visitors correlated positively with G+/G- ratio and negatively with soil moisture, trans/cis PLFA stress indicator, and activity of peroxidases. Overall the effect of attendance to sediment microbial communities was found significant and it is likely related to input of nutrients into caves. The results might serve as a supporting material for reconsidering of the cave attendance policy.
STUDY OF NEMATODE-BACTERIA INTERACTION IN PINE WILT DISEASE

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Background

Pine wilt disease (PWD) is one of the most threatening diseases for worldwide forestlands, and is caused by the plant parasitic nematode *Bursaphelenchus xylophilus*. Bacterial communities associated with the *B. xylophilus* are suggested to play a role in PWD development.

Objectives

Our aim is to understand the contribution of nematode-associated bacteria (in particular, *Serratia quinovorans* FX1 and *Serratia marcescens* PWN146) in PWD, as well as their behavior towards *B. xylophilus*.

Methods

Previously, we have characterized the phenotype of *S. quinovorans* FX1 and *S. marcescens* PWN146, and their performance under *in vitro* oxidative stress conditions, alone and in association with *B. xylophilus*. We are now analyzing both genomes and comparing both with endophytic and plant pathogenic bacteria.

Conclusions

Both *Serratias* present phenotypic plasticity in traits that favour their fitness inside of the host tree (Vicente et al. 2011; Vicente et al., 2012), and under certain circumstances can help the nematode to overcome the hostile tree environment (Vicente et al., 2013). We believe that the beneficial association nematode-bacteria may be facultative and co-dependent of other unknown features (Nascimento et al. 2014).

DIVERSITY AND BIOTECHNOLOGICAL POTENTIAL OF THRAUSTOCHYTRIDS FROM THE CHINA’S COASTAL WATERS
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Background

Thraustochytrids are unicellular and heterotrophic fungal-like marine protists with known low diversity and have long been known to produce diverse interesting bio-products. However, their diversity and biotechnological potential have been barely explored in the China’s coastal waters.

Objectives

Investigate the diversity of thraustochytrids using molecular and cultivation methods; explore the isolated strain in the production of biotechnological potential.

Methods

Flow Cytometer, pyrosequencing, plate cultivation, fermentation, and HPLC analysis

Conclusions

Thraustochytrids abundance displays significant spatial and temporal variations in the coastal waters and can surpass that of bacteroplankton. The molecular diversity of thraustochytrids is much higher than what we have known. Samples from contaminated areas tend to contain higher richness and diversity than those uncontaminated areas. The majority of recovered reads are members of the family Thraustochytriaceae. Thus, thraustochytrids were the dominant components of labyrinthulomycetes and presumably play significant role ocean carbon cycling the coastal waters. The culturable diversity of thraustochytrids is low in coastal waters. However, most of cultivated strains produces carotenoids, polysaccharides and proteases. Some of the stains produce significant amount polyunsaturated fatty acids (PUFA) (e.g. DHA) and saturated fatty acids. Thus, thraustochytrids can be used for the production nutraceuticals and biodiesel.
Background: Terra preta do indio is a typical hortic anthrosol in the amazonas region derived from long-term human landscape cultivation. This distinct type of soil is characterized by a high storage capacity of plant nutrients as well as a high content of carbon enabling agriculture even on soils with low humus and/or nutrient content. Due to certain similarities in consistence and composition, biochar is recently discussed as a value artificial soil amendment to obtain similar positive effects on soil fertility like terra preta. In addition to the structural and compositional analysis of biochars the interaction of the chars with living microorganisms is studied, to explain the effect of biochar enriched soils from a microbiological point of view. Due to seclusion of a biogas system and the consequent creation of a separate world the study of effects by biochar colonization with microorganisms by minimizing the confounding factors is more easier compared with soil tries.

Objectives: In this study the microbial interaction caused by colonization, biofilm formation and degradation with two types of char (hydrochar and pyrolized char), which differs in degradation stability, is examined using a mesophilic gas lift biogas fermentation system.

Methods: In samples taken from mesophilic to thermophilic biogas system at various time points the bacteria:archaea ratio was determined by DNA-based quantitative PCR according to Yu et al. [1995]. In addition, the treated chars were investigated with respect to their surface properties by gas adsorption method (BET).

Conclusions: Caused by feedstock and process condition during carbonization the pyro-char shows higher stability regarding degradability while fermentation. This leads due to the colonization of the char to a measurable reduction of pore surfaces in the pyro-chars, whereas the instability of the hydrochars lead to an increase in the specific surface area compared with the unfermented sample. Most interesting is the dynamic not stable population in the microbial community over time under anaerobic digestive conditions.
COMPARATIVE ANALYSIS OF THE MICROBIAL COMMUNITIES ON THE SURFACE OF MINERAL ENHANCED BIOCHARS

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Background
Biochar is a kind of porous material made by the pyrolysis of biomass under complete or partial exclusion of oxygen. It has applications in different areas, with the most important one being as an amendment to improve the quality of agricultural soils. Several studies reported significant changes for the microbial communities in bulk soil with biochar-treatment, whereas microbiota colonised on the surface of biochar is largely neglected.

Objectives
In this study, two types of mineral enhanced biochars, including kaolinite (Kao) and bentonite (Be), and non-enhanced biochar (Bam) were applied into soil with (F) or without (NF) traditional fertilisation (urea, superphosphate and potash) on a pot trial. We compared the differences of microbial communities on the surface of biochar and amended soil after 4-month incubation.

Methods
Total DNA was extracted from biochar particles and soil respectively and sequencing using the Illumina MiSeq platform.

Conclusions
Analysis of 2001281 quality-filtered sequences revealed high microbial diversity in these samples. Bray-Curtis distance-based non-metric multidimensional scaling (nMDS) analysis showed that the microbiota of biochar particle and soil were clearly separated into four clusters (soil vs. particle, F vs. NF). Samples in the NF cluster were observed to vary by biochar types. Acidobacteria, Actinobacteria, Bacteroidetes, Proteobacteria was detected as the dominant phyla regardless of particle and soil. Furthermore, unique OTUs in each samples were identified, three OTUs from the class Gammaproteobacteria, the family Oxalobacteraceae, the genus Thiobacillus were found significantly more abundant in biochar particles.
CHARACTERIZATION OF ENDOLYSINS FROM A VIBRIO PARAHAEMLYTICUS BACTERIOPHAGE VPP-KF1

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Background

Vibrio parahaemolyticus is a halophilic gram-negative bacterium in seawater and estuaries. It is one of the leading causes of seafood-related illness through the ingestion of raw or undercooked seafood and cross contamination from unsanitary cooking environments. Endolysin, a phage-encoded enzyme that breaks down bacterial peptidoglycan at the terminal stage of the phage reproduction cycle, is reported to be effective for the control of pathogenic bacteria as well as bacteriophage itself.

Objectives

The purpose of this study was to examine lytic properties of endolysins from bacteriophage Vpp-KF1 infecting Vibrio parahaemolyticus.

Methods

Lytic bacteriophage Vpp-KF1 infecting Vibrio parahaemolyticus was isolated from west coastal area of Korea, and its whole genome was sequenced and analyzed. Endolysin genes were identified in the bacteriophage genome, and then the endolysin genes were cloned and overexpressed in Escherichia coli. The endolysins were purified by Ni-NTA affinity chromatography. The effects of temperature and pH on the lytic activities, and the lytic spectrums of endolysins were determined by measuring changes in the optical density of cell suspension.

Conclusions

Endolysin genes were successfully expressed in E.coli expression system, and the enzymes were purified as soluble forms. The endolysins were relatively stable and active over broad temperature (from 24˚C to 55˚C) and pH (from 2 to 10) ranges. The lytic spectrums were relatively narrow because those endolysins were effective mainly in Vibrio species when treated without EDTA. These results suggested that the bacteriophage Vpp-KF1 endolysins have potential as an alternative to antibiotics or antibacterial agents in controlling bacterial contamination.
THE METABOLIC EFFECTS OF PHAGE INFECTION: PHAGE SPECIFIC OR NOT?

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Background
Bacterial viruses are fully dependent on their host cell metabolic resources to fuel their propagation. Through millions of years of co-evolution, many bacteriophages have acquired host genes in their genomes which can overcome biochemical bottlenecks in specific metabolic processes, and therefore were termed ‘auxiliary metabolic genes’ (Thompson et al, 2011).

Objectives
As such, the metabolic response to phage infection would vary depending on the number of AMGs in the genome of the phage. To gain additional insight in this topic, we studied the metabolic variations in an ubiquitous, gram-negative bacterium, Pseudomonas aeruginosa, under attack of six distinct lytic dsDNA phages.

Methods
We used state of the art metabolomics techniques (Fuhrer et al, 2011). A total of 6006 ions were measured, of which 918 were annotated to P. aeruginosa metabolites. After filtering 375 unique masses were retained, corresponding to 810 P. aeruginosa metabolites.

Conclusions
Using hierarchical clustering we saw that the metabolic response triggered by phage infection is not a general response but depends on the requirements of a specific phage. A pathway enrichment analysis showed that not all pathways were influenced equally. An example is the clear up regulation of the pyrimidine metabolism, which can be linked to a study from Enav et al in 2014 that showed that AMGs with a function in the pyrimidine and purine metabolism are enriched in the viral gene pool. These results indicate that unique AMGs might play a crucial role in the specific host metabolic response induced by different phages.
Background

*Lactococcus garvieae*, one of the most important pathogens in the aquaculture sector, has been recently recognized as an opportunistic human pathogen. However, little is known about the factors controlling its growth.

Objectives

The aim of this project is to isolate and characterize temperate and virulent phages of *L. garvieae*.

Methods

Forty-five *L. garvieae* strains isolated from various ecological niches were tested for the presence of inducible prophages. Virulent phages were also isolated from different environments. Morphological and genomic characteristics of the new phages were studied through transmission electronic microscope (TEM), DNA restriction profile, protein profile, host range, and in few cases, genomic sequences.

Conclusions

Twenty temperate phages belonging to the *Siphoviridae* family and five new virulent phages able to infect *L. garvieae* strains have been isolated and partially characterized. Temperate phages, integrated into the genome of the host cell, promoted genome plasticity and appeared to be involved in gene mobilization. Virulent and strain–specific phages may be useful as alternatives to antibiotics to treat *L. garvieae* infections.
ANALYSIS OF CDT-I PHAGE PRODUCTION IN ESCHERICHIA COLI

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Background
Cytolethal distending toxin (CDT)-producing Escherichia coli (CTEC) have been isolated from patients with diarrhea. Previously, we have reported that cdt-I genes in E. coli are located in genetically diverse lambdoid phages, and CDT-I phage (CDT-IΦ) could be transferred horizontally between E. coli in pathotype-specific manner.

Objectives
In this study, CDT-IΦ was comprehensively analyzed to understand underlying mechanisms of its production.

Methods
Culture supernatants of 46 CTEC-I strains were examined for CDT-IΦ production by plaque assay with E. coli C600 both in the presence and absence of mitomycinC (MMC). Plaque assay was also performed with DrecA-E. coli (strain TOP10F') and its recA complement as a recipient strain. Presence of other lambdoid phages was examined by using 23 known cI gene-probes.

Conclusions
Among analyzed strains belonging to diverse EPEC and non-EPEC, only EPEC O127 and O142 could produce CDT-IΦ plaques. All the CTEC-I O142 produced CDT-IΦ both in the presence and absence of MMC, whereas CTEC-I O127 produced only in the presence of MMC. However, under identical host background (C600), these phenomena disappeared. CDT-IΦ plaque was not produced against DrecA-E. coli, whereas produced when complemented with recA gene. CTEC-I O127 strains harbored 6 or 7 cI genes, whereas O142 had 2~4. CDT-IΦ production was differently regulated in CTEC-I O127 and O142, suggesting involvement of host factors, which might contribute to inhibit or stimulate spontaneous CDT-IΦ induction in these serogroups in a RecA-dependent pathway. Moreover, intra-prophage interactions with O127-specific prophages could be related to CDT-IΦ production.
Phage therapy has been accepted as an alternative to chemotherapy in aquaculture systems especially in shrimp larval production. Uniqueness of bacteriophages as therapeutics is their self-replicating character and specificity to the target organisms. However, concerns exist with their use in vivo as they may turn out to be the agents of generalized transduction in the environment mediating transfer of genes among bacteria. This is important because phages selected for therapeutic purpose must not carry virulence genes or properties conferring antimicrobial resistance and toxin genes to the host bacteria. Moreover, therapeutic phages should not become latent as well during their life cycle. Using an enrichment method, bacteriophages were isolated from sediment and water samples from Vembanad estuary, Cochin, India employing 12 isolates of Vibrio harveyi. By this method four phage lysates were obtained, purified by soft agar overlay method, titred and stocked at 4°C and stability checked over a year. Host range of the phage was assessed against 162 bacterial isolates spanning 14 species to which they showed broad spectrum lytic activity. One phage was lytic to both V. alginolyticus and V. parahaemolyticus potentially pathogenic to shrimp and other aquatic animals in culture systems. Lytic efficiency of all phages was checked on their respective host bacterial species. In all cases the phages were able to arrest growth of V. harveyi for about 5-6 hours after which phage resistant forms emerged and began to dominate. By electron microscopy, three of the phages were identified as Myovirus-like with icosahedral head, short collar and contractile tail. The fourth one was Siphovirus-like with icosahedral head and long, flexible, helical non-contractile tail. It was infective to 62% V. harveyi isolates tested and to its near taxonomical neighbours, V. alginolyticus and V. parahaemolyticus. Development of resistance or lysogenization is an impediment in the development of phage therapy in aquaculture necessitating deeper understanding on their genome and phage-host dynamics in the environment.
HIGH FREQUENCY OF MUTATOR STRAINS INDUCED BY PHAGE GHOST ATTACHMENT

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Background
Lytic phages induce bacterial mortality and contribute to the global carbon cycle. Because their main impact on the host is lysis of the cell, short-term contact between the host and phage is considered genetically less important unless genetic exchange occurs.

Objectives
In order to understand the genetic impact of a lytic phage on the host, we examined the fate of host cells after attack by a lytic phage and the mutant frequency of cells that did not undergo lysis.

Methods
After incubating Escherichia coli with phage T4 or the ghost for 10 min, the cells were placed on LB plates and allowed to grow. Colonies that escaped lysis were randomly selected. Mutant frequency to nalidixic acid resistance in the escaped colonies was examined. The insertion of phage DNA and mutations on the genomes of mutator strains which showed elevated mutant frequencies were examined by whole genome sequencing analysis. Role of DNA polymerase IV gene (dinB) was examined with RT-PCR and dinB-deficient mutant.

Conclusions
After the attachment of the phage T4 to E. coli cells, 1% E. coli cells showed an approximately 40-fold increase in mutant frequency. Phage ghosts increased the incidence of mutator strains. Phage DNA was absent in the mutator strains. The attachment of the phage ghosts to the host induced the mRNA expression of the dinB, resulting host genome mutagenesis. These results suggested that attachment of a lytic phage alters the genome of the host that has escaped lysis at high frequency, without relying on gene transfer.
Background

Viral hemorrhagic septicemia virus (VHSV) is known as the most common cause of diseases in fisheries. VHSV infects fishes of fresh water as well as ocean. Natural VHSV may cause of disease by introducing into fish farm. A huge amount of fry are stocked into the coast.

Objectives

Therefore development of VHSV detection method is required. For the detection of VHSV, enzyme immunoassay using polyclonal antibodies, western blotting, and electron microscopy are mainly used. However these methods have limitation to detect latent infection and an initial viral disease. Recently nucleotide amplification test using polymerase chain reaction (PCR) has been used widely for rapid and specific detection of viruses.

Methods

In this study, TaqMan-based real-time PCR assays to detect VHSV was developed and validated. Specific primers for amplification of VHSV were selected, and VHSV was quantified by use of TaqMan probe.

Conclusions

The sensitivity for VHSV real-time PCR was calculated to be $8.10 \times 10^{-1} \text{TCID}_{50}/\text{mL}$. The TaqMan probe real-time PCR assay was proven to be reproducible and very specific to VHSV. Therefore, it was concluded that this rapid, specific, and robust assay could replace the conventional assay for detection of VHSV.
COMPARATIVE ANALYSIS OF AVIAN PATHOGENIC ESCHERICHIA COLI PHAGES

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Background

Avian pathogenic Escherichia coli (APEC) causes colibacillosis and significant economic losses in the poultry industry.

Objectives

To isolate and characterize poultry fecal phages that are active against Escherichia coli serogroups O1, O2, and O78; the main causative agents of avian colibacillosis.

Methods

Lytic phages (71) were isolated from poultry fecal material and their lytic spectrum of activity was determined against 61 APEC (O1, O2, O78 serotypes) strains collected from Belgium. The phages with the broadest activity spectrum were characterized by, restriction-fragment length polymorphism (RFLP). Based on their obligate lytic characteristics and broad host spectrum of activity, some were selected for whole genome sequencing using the Nextera XT™ kit (Illumina, San Diego, USA). The morphology of the phages was determined by negative staining and Transmission Electron Microscopy (TEM).

Conclusions

Phages PhK7, PhK10S, PhK10L, PhK12B, PhK21, PhK22, PhK12A, PhK13, PhK26, PhK2, PhK3, PhK6, Ph431, PhC3 and PhK33C were isolated and further characterized. The most active phages were Ph431 active against 70% (43/61) of the APEC strains, PhC3 against 68% (42/61), PhK33C against 41% (25/61), PhK12A against 26% (16/61), Ph12A 19% against (12/61).
In all samples, particles were observed consisting of a non-enveloped head with icosahedral symmetry and a tail with helical symmetry; characteristics attributing the particles to the bacteriophage order of the Caudovirales and belonged to families Myoviridae and Siphoviridae.

Genome analysis of some of phages indicated there were close relatives of Siphoviridae present under the unclassified Tunalikevirus (PhK2, PhK3, PhK6) and unclassified Siphoviridae (PhK10S, PhK7, PhK10L, PhK12B, PhK21, PhK22, Ph12A, PhK26, PhK13).
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OBSERVATION OF PHAGE-ANTIBIOTIC SYNERGY IN VARIOUS COMBINATIONS OF BACTERIA, BACTERIOPHAGE, AND ANTIBIOTICS

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Background
When phages infect bacteria cultured in the presence of sub-lethal doses of antibiotics, sizes of phage plaques significantly increase. This phenomenon is known as phage-antibiotic synergy (PAS).

Objectives
We extended observation of PAS to a wide variety of bacteria-phage pairs with four different classes of antibiotics.

Methods
In the presence of sublethal dose of various antibiotics, phage growth and burst was measured.

Conclusions
PAS was observed in both Gram-positive and Gram-negative bacteria. Cell wall synthesis inhibitors and DNA metabolism inhibitors generally induced PAS. Some protein synthesis inhibitors induced PAS, while others did not. The use of sub-lethal dose of ampicillin, cefotaxime, ciprofloxacin, or mitomycin C allowed formation of highly visible plaques of increased sizes when various bacteriophages infected Gram-positive bacteria such as Staphylococcus aureus, Bacillus cereus, and Enterococcus feacalis. In Gram-negative bacteria, Escherichia coli and Pseudomonas aeruginosa phages showed increased plaques in the presence of sub-lethal dose of cefotaxime, ciprofloxacin, or mitomycin C. We also confirmed that cells stressed with β-lactam and quinolone antibiotics filamented or swelled extensively. Burst sizes of bacteriophages also increased in the presence of antibiotics. Increase of production of phages in the presence of antibiotics was also shown in vivo using Caenorhabditis elegans as a model animal. Induction of SOS response was observed in many cases of PAS, but not all. Enlarged production facility allowed increased DNA replication, transcription, and translation of phage leading to PAS.
IMMUNOGENICITY EVALUATION OF HEPATITIS B (HB) VACCINES UTILIZED IN THE EXPANDED PROGRAMME ON IMMUNIZATION (EPI) IN SOUTH-EASTERN NIGERIA

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Background
Vaccination effectively enables the control of many infectious diseases. However, we cannot always avoid the problem of adverse reactions accompanied by vaccination. The most common cause of vaccine-related side-effects is human errors linked to logistics problems.

Objectives
This study investigated
a. whether or not the Hepatitis B (HB) Vaccines in use in South-East Nigeria have the features and characteristics needed for them to satisfy stated or implied needs.
b. the efficiency of cold-chain facilities in South-east, Nigeria

Methods
The immunogenic quality of the vaccines was determined by quantitation of the antibody titer (using ELISA method) in vaccinated laboratory animals. The mean antibody titers for the control, Enugu/Ebonyi, Imo, Anambra and Abia States’ vaccines were 8.7, 398.2, 353.7, 396.3 and 369.6 mIU/mL respectively. The vaccine used as the control did not produce sufficient protective effect (antibody titer was less than 10 mIU/mL). One-way analysis of variance shows that the means of all the samples, including the control, are significantly different at P < 0.05. Also, Bartlett's test for equal variances showed that the variances are also significantly different. The vaccines do not all produce equal protection (P value was < 0.0001), their antibody titers being statistically different. The Dunnett's Multiple Comparison Test shows that the mean antibody titers produced by the vaccines from the States significantly differ from the value produced by the control vaccine at P < 0.05.

Conclusions
All the vaccines tested produced the required amount of protective antibodies. The cold-chain facilities, at the time of vaccines collection, are in good condition.
Background: Bacteriophage endolysins are phage-encoded peptidoglycan hydrolases by which bacteriophage are released from bacterial cells and which are effective in bacterial cell wall lysis, when applied exogenously. The gene product of orf50 from actinophage µ1/6 of Streptomyces aureofaciens, producing tetracycline, is endolysin Lyt µ1/6. It shows a two-domain modular structure, consisting of an N-terminal catalytic and a C-terminal binding domain (CBD). BLAST analysis of Lyt µ1/6 CBD disclosed the presence of a peptidoglycan binding domain type 1 (PG_binding_1) which was also confirmed by in vitro experiments.

Objectives: The main aim of this study was to determine the part of the Lyt µ1/6 CBD responsible for binding activity towards the Streptomyces cell wall, as well as to study concrete truncated forms of it in fusion with green fluorescent protein (GFP).

Methods: The sequence of the Lyt µ1/6 CBD and truncated forms of it were cloned and expressed in Escherichia coli. All recombinant proteins were isolated and partially purified with conventional chromatographic methods. The ability of the CBD truncated variants fused to GFP to bind to the substrate consisting of Streptomyces aureofaciens NMU cells was demonstrated by specific binding assays. Subsequently, the binding of the CBD fusion products was visualized by fluorescence microscopy.

Conclusions: It could be assumed that the binding activity of the CBD is located within the three helices of the PG_binding_1 domain. This fact is supported by a negative control which is a construct of the N-terminal part of the Lyt µ1/6 CBD, showing no binding activity.

Keywords: bacteriophage endolysin, Streptomyces aureofaciens, cell wall binding domain (CBD), green fluorescent protein (GFP), truncations, binding activity

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THE ROLE OF AVIN08930 IN BIOSYNTHESIS OF POLY-B-HYDROXYBUTIRATE AND ALKYLRESOLCINOLS IN AZOTOBACTER VINELANDII

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Background

\textit{Azotobacter vinelandii} is a Gram-negative bacterium able to synthesize polyhydroxybutyrate (PHB), a polyester to produce biodegradable plastic. \textit{A. vinelandii} can undergo a differentiation process to produce resistant-desiccation cysts. During the encystment, the bacterium synthesizes the phenolic lipids alkylresorcinols (ARs), which replace the membrane phospholipids. The operons \textit{phbBAC} and \textit{arsABCD} encode the enzymes necessary for PHB and ARs synthesis and their transcription are activated by PhbR and ArpR, respectively.

The two component regulatory system GacS/GacA is essential for biosynthesis of PHB and ARs, since a \textit{gacA} mutation abrogates their production. GacA activates the transcription of genes encoding sRNAs \textit{rsmZ1-7} and \textit{rsmY}, which bind to RsmA; a small protein that blocks the translation of \textit{phbR} and \textit{arpR} mRNAs. Because inactivation of \textit{rsmA} in a \textit{gacA} mutant did not restore the PHB and ARs synthesis, we hypothesize that there is another pathway by which GacS/GacA controls the biosynthesis of PHB and ARs in \textit{A. vinelandii}.

Objectives

To characterize the role of Avin08930 on PHB and ARs in \textit{A. vinelandii}.

Methods

A bioinformatic analysis of \textit{A. vinelandii} genome allowed us to identify a consensus binding site for GacA in the Avin08930 promoter. By using qPCR and transcriptional fusions, we determined if Avin08930 expression is dependent of GacA. An Avin08930 mutant was constructed and PHB and ARs synthesis as well as \textit{phbB} and \textit{arsA} expression were determined in this strain.

Conclusions

The transcription of Avin08930 is activated by GacA. In the Avin08930 mutant the \textit{phbB} and \textit{arsA} expression as well as the PHB and ARs levels were reduced.
Global regulatory networks

A POTENTIAL NOVEL GENE TRANSCRIPTION REGULATORY SYSTEM IN STAPHYLOCOCCUS GENUS
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Background
Gene transcription regulatory pathways are crucial for adaptation to constantly-changing environmental conditions. These include pathogen-host interactions. Up until now, few such systems have been characterised for Staphylococcus aureus and simultaneously demonstrated as relevant for virulence.

Objectives
This study presents a comprehensive analysis of a three-gene operon saoABC, which likely constitutes yet uncharacterised staphylococci-specific gene expression regulatory system. The research is focused on the operon structure and conservation among species of Staphylococcus genus, DNA-binding properties of SaoC protein, changes in saoABC operon genes expression triggered by stress stimuli, links between saoABC operon and known gene transcription regulatory systems as well as on differences between phenotypes of the wild type and the null mutants of saoA and saoC genes.

Methods
The applied methods include comparative genomic analysis, heterologous SaoC protein expression and its purification, Real-Time PCR gene expression quantification, allelic replacement, virulence analysis in chicken embryo model and intracellular survival within human gingival fibroblasts and keratinocytes.

Conclusions
The revealed features of saoABC operon convincingly suggest that it is a novel operon of a transcription regulation system, likely of a yet uncharacterised alternative sigma factor. The changes in the operon genes expression specifically induced by starvation and acidification point out the possible role of saoABC in intracellular survival of staphylococci in host cells.

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Background
In Caulobacter crescentus cell cycle is controlled at multiple levels by a plethora of regulators, such as GcrA, a putative transcriptional regulator with no functional characterization. Another regulator involved in controlling gene expression is CcrM, a GAnTC-specific N6-adenosine methylase active only at the end of S-phase; as a consequence, at every round of replication a fully methylated promoter of a gene gives two different hemi-methylated copies.

Objectives
Here we investigated in vivo and in vitro the role of GcrA during the S-phase on the transcription of important cell cycle genes that are controlled by CcrM.

Methods
We used a combination of ChIP (chromatin-immunoprecipitation), biochemical and biophysical experimentation and genetics to show that GcrA is a dimeric DNA-binding protein that preferentially targets promoters harbouring CcrM methylation sites in Alphaproteobacteria. After tracing CcrM-dependent promoter marks at a genome-wide scale, we showed that these marks recruit GcrA in vitro and in vivo. Moreover in the presence of a methylated target, GcrA recruits the RNA polymerase to promoters, consistent with its role in transcriptional activation. In vitro transcription assay using purified GcrA and RNAP on several GcrA-regulated genes along the chromosome showed that GcrA is a modulator of transcription that activates or represses depending on the methylation state of promoters. We confirmed this observation in vivo using RNA-FISH, revealing in addition a potential role of GcrA in spatial regulation of transcription.

Conclusions
In conclusion from our results we propose a novel model of methylation-dependent transcriptional regulation that modulates the expression of important genes during cell cycle.
Background

PAS-LuxR regulators are highly conserved proteins devoted to the control of antifungal production by binding to operators located in given promoters of polyene biosynthetic genes. These regulators have been found to be encoded in all known biosynthetic gene clusters of polyene polyketides, and hence considered pathway-specific modulators.

Objectives

The canonical operator of PimM, archetype of this class of regulators, has been used to search for putative targets of orthologous protein PteF in the genome of Streptomyces avermitilis.

Methods

We have used bioinformatic and electrophoretic mobility shift techniques to identify PimM binding sites along the genome, and mutants in pteF for the functional characterization of selected operators.

Conclusions

Contrary to the stablished opinion, PAS-LuxR regulators may control directly many different processes previously unforeseen, such as genetic information processing, DNA replication and repair, energy, carbohydrate, and lipid metabolism, morphological differentiation, transcriptional regulation, and other secondary metabolites biosynthesis. To prove that, we demonstrate that PteF, is able to cross-regulate the biosynthesis of the ATP-synthase inhibitor oligomycin in addition to that its canonical target filipin. Our results place these regulators as wide-domain modulators in regulation networks, and open new possibilities for the enhancement and awakening of metabolite production in Streptomyces.
Background

*S. clavuligerus* is a versatile and industrially important strain characterized by its ability to produce several economic relevant bioactive compounds such as cephamycin C and the β-lactamase inhibitor clavulanic acid. *S. clavuligerus* genome harbours 48 gene clusters encoding potential secondary metabolites. Under experimental conditions these clusters are not expressed (“silent clusters”), are expressed at very low levels producing undetectable amounts of the compound and in many cases the final product structure encoded by the cluster is unknown (“cryptic clusters”). A large part (21%) of the wild type genome is located in a 1.8 Mb megaplasmid that greatly influences secondary metabolites biosynthesis even if the secondary metabolites are chromosomally encoded.

Objectives

The objective is to analyze the differential expression between the wild strain and a pSCL4-deleted mutant and identify the activation of cryptic gene clusters in this mutant.

Methods

Deletion of pSCL4 was performed on the wild type *S. clavuligerus* ATCC 27064 using the Redirect Method. Microarray (*Agilent 8x15K*) data were normalized and analyzed with the *Bioconductor package limma*.

Conclusions

We demonstrated that the absence of linear megaplasmid pSCL4 induced holomycin overproduction, (*S. clavuligerus* is holomycin non-producer). The transcriptomic results have shown the expression of several cryptic gene clusters in *S. clavuligerus* pSCL4-. Therefore, the extrachromosomal-loss elements may be a strategy for the cryptic clusters activation in this microorganism.
CHARACTERIZATION OF SFBR, AN AUTOREGULATOR RECEPTOR FROM STREPTOMYCYES FILIPINENSIS

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Background

Gamma-butyrolactones are small signaling molecules that are mainly produced by Streptomyces species. They have been shown to affect secondary metabolism and/or morphological differentiation by binding to specific cytoplasmic receptors and therefore inhibiting their binding to target DNA sequences.

Objectives

Identification and characterization of a gamma-butyrolactone receptor (GBR) encoding gene from Streptomyces filipinensis, the industrial producer of the polyene antibiotic filipin.

Methods

A genomic library of S. filipinensis DSM 40112 was screened with a probe designed from a HTH DNA-binding motif conserved among autoregulator receptors. A putative receptor encoding gene (sfbR) was found and sequenced. Characterization of this gene was performed using gene replacement and complementation techniques.

Conclusions

The sfbR gene encodes a putative gamma-butyrolactone receptor protein which shows homology with other described GBRs such as AvaR from S. avermitilis or Brp from S. clavuligerus. Contrary to what occurs with many GBR-encoding genes, an adjacent gene encoding a gamma-butyrolactone biosynthetic enzyme was not found. Gene deletion resulted in a significant decrease in filipin production while complementation restored parental production, thus suggesting that SfbR positively controls the filipin biosynthetic genes. Moreover, sfbR deletion resulted in altered morphological differentiation when the mutant strain was grown on different solid media. Our results indicate that the SfbR protein regulates both sporulation and filipin production in S. filipinensis.
Background

*Pseudomonas fluorescens* spp. are present in all major natural environments, and able to quickly adapt to environmental modifications. Physiologically this adaptation depends primarily on membrane lipid composition and on its capacity to adjust membrane lipids in terms of environmental modifications. Previously we have already characterized the phospholipid composition of *P. fluorescens* [1].

Objectives

The aim of this work was to establish if the response to increasing temperature involves changes in the structure and the composition of *P. fluorescens* phospholipids at different phases of growth. The lipidomes of two *P. fluorescens* strains from different ecological niches: one clinical isolate MFN1032 and one airborne MFAF76a, were investigated at 28°C and 37°C.

Methods

To investigate *Pseudomonas* lipidome, MALDI-TOF MS Imaging was coupled to HPTLC. Additionally, NMR spectroscopy led to precise the phospholipid head group identification.

Conclusions

In stationary growth phase *P. fluorescens* produces a new phospholipid class: phosphoaminolipid (PAL).

Results provide evidence that next to the increasing of the temperature *P. fluorescens* is able to modify its phospholipid composition using two pathways. First the increasing of degree of unsaturation of fatty acid chains controls the membrane fluidity. Additionally the increase of the synthesis of anionic phospholipids (e.g. phosphatidylglycerol) results in the membrane charge changes. Physiologically, these two pathways allow *P. fluorescens* to adapt to stress conditions.
References

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Global regulatory networks

FINE TUNING OF METABOLISM: THE ROLE OF CRC/CRCZ-CRCY SYSTEM IN INTEGRATING GLUCONEOGENIC AND GLYCOLYTIC METABOLISM IN PSEUDOMONAS PUTIDA

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Background

In *P. putida*, the Crc protein and the small RNAs CrcZ and CrcY are key players in carbon catabolite repression (CCR), which facilitates the assimilation of the most efficient carbon sources, improving growth rate and fitness. Succinate and glucose are good carbon sources for *P. putida*, but elicit a weak CCR. Little information is available on the regulatory and metabolic interactions when both compounds are present. The choice between them is crucial on driving a metabolism focused on glucose consumption (glycolytic), or an inverse configuration directed to gluconeogenesis (succinate consumption and synthesis of sugars).

Objectives

The aim was to clarify whether there is a preference for succinate or glucose, and whether the Crc-dependent CCR has a role in coordinating metabolism when cells grow with a mixture of succinate and glucose at non-limiting concentrations.

Methods

We undertook an integrative approach combining metabolic, transcriptomics and constraints-based metabolic flux analyses.

Conclusions

Succinate was consumed faster than glucose and allowed a higher growth rate. However, both compounds were co-metabolised when provided simultaneously. CrcZ and CrcY levels were lower when both substrates were present than when only one of them was provided, suggesting a role for Crc in coordinating metabolism. Flux distribution analyses suggested that, when both substrates are present, Crc works to organize a metabolism in which carbon compounds flow in two opposite directions: from glucose to pyruvate, and from succinate to pyruvate. Therefore, Crc serves not only to favour the assimilation of preferred compounds, but also to balance the carbon fluxes, optimising metabolism and growth.
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Global regulatory networks

SEVERAL ALTERATIONS IN PHYSIOLOGY OF YERSINIA ENTEROCOLITICA O:3 HFQ MUTANT ARE MEDIATED BY OVEREXPRESSION OF LRHA.
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Background
In previous studies we observed that Hfq strongly downregulates the LrhA (LysR homologue A) expression in Yersinia enterocolitica. Hfq was first identified as a host bacterial factor required for the replication of bacteriophage Qβ RNA. Now it is known that Hfq plays a crucial role in global post-transcriptional regulation network as an RNA chaperone that binds and stabilizes small regulatory RNAs. On the other hand, LrhA was shown to regulate negatively motility, chemotaxis and flagella synthesis in Escherichia coli.

Objectives
In this work we wanted to elucidate the role that LrhA plays in the phenotype of the hfq mutant of Y. enterocolitica.

Methods
The hfq and lrhA genes were knocked out to create single and double mutants for subsequent analyses. The mutant strains were complemented in trans by functional hfq and lrhA genes. LrhA was overexpressed from an IPTG-inducible plasmid. The strains were analysed for motility, biofilm formation, growth rate and mannitol utilization affected in the hfq single mutant.

Conclusions
In the lrhA hfq double mutant we could observe partial or full restoration of such features as growth rate, motility, biofilm production and mannitol utilization. On the other hand, overexpression of LrhA in wild type bacteria caused hypermotility and different colony morphology on mannitol-containing Yersinia selective agar plates. Our results thus showed that several alterations in physiology present in the hfq mutant of Y. enterocolitica are mediated by increased expression of LrhA.
Background

The obligate anaerobe Azoarcus anaerobius degrades resorcinol by an oxidative reaction where the aromatic ring is hydroxylated at position 4 to render hydroxyhydroquinone (HHQ). The three enzymatic steps are transcribed in three different operons controlled by two bacterial enhancer-binding proteins (bEBP) that share 97% identity, RedR1 and RedR2.

Objectives

The work aims at understanding the coordinate expression of the pathway genes in response to the presence of the substrate by elucidating the specific activation mechanism of each regulator.

Methods

Genetic and biochemical approaches were used to define the activation mechanism of the two regulators.

Conclusions

Despite the high sequence identity between RedR1 and Red2, they are controlled by different activation mechanisms: in the absence of effector, RedR2 is kept inactive through binding to the membrane protein BtdS, the small subunit of HHQ dehydrogenase. The presence of the substrate favors electron flow through BtdS, allowing unbinding of RedR2 in an active conformation to induce low expression levels of the enzymes. RedR1 then becomes activated by the pathway intermediate hydroxybenzoquinone. Both regulators are required for maximum pathway expression, suggesting an amplification regulatory cascade is required for coordinate expression of the entire pathway. The structure of the pathway promoters and
regulator binding have been characterized.

References
REGULATION OF CSRA EXPRESSION BY TWO COMPONENT SYSTEMS IN E. COLI
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Background

Global regulators play major roles in response to a variety of intracellular and extracellular signals, allowing bacteria to adapt to environmental changes.

We are particularly interested in CsrA, a post-transcriptional global regulator acting by binding target mRNAs and thereby modulating translation either negatively or positively.

CsrA is important for cell growth and regulates central carbon metabolism and social behaviors.

Transcriptional regulation of csrA expression remains largely unknown although it was shown that Sigma70 and RpoS directly regulate csrA expression. At the post-translational level, activity of CsrA is regulated by 2 major sRNAs, csrB and csrC, which bind CsrA and sequester it away from its targets. Expression of csrB and csrC is regulated by the BarA/UvrY two-component system.

Objectives

Since CsrA controls pathways that are influenced by environmental conditions, it is likely that csrA expression is regulated by two-component systems.

Methods

A screen using lacZ transcriptional fusions with the csrA promoter was set up to test two-component system mutants. In addition, bioinformatics analysis was performed to determine response regulator binding sites in the csrA promoter.

Conclusions
Our results indicate that *csrA* expression is positively regulated by the Cpx two-component system. The Cpx system responds to a broad range of stimuli including pH, salt, lipids and misfolded proteins that cause perturbation in the envelope.

Bioinformatics analysis predicted potential CpxR binding sites in the *csrA* promoter region. EMSA and mutagenesis experiments will be performed to whether Cpx directly regulates *csrA* expression.
CHARACTERIZATION OF THE ESCHERICHIA COLI SIGMA-S CORE REGULON BY CHROMATIN IMMUNOPRECIPITATION-SEQUENCING (CHIP-SEQ) ANALYSIS.

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Background

In bacteria, selective promoter recognition by RNA polymerase is achieved by its association with σ factors, accessory subunits able to direct RNA polymerase “core enzyme” (E) to different promoter sequences.

Objectives

The aim of this work was to identify promoters bound by Eσ⁵, an RNA polymerase form mainly active during slow growth and in response to stress conditions, by using a Chromatin Immunoprecipitation-sequencing (ChIP-seq) approach.

Methods

ChIP-seq experiments were performed in an Escherichia coli strain carrying a chromosomal rpoSHis⁶ allele grown to stationary phase in rich medium.

Conclusions

We identified 78 Eσ⁵-binding sites, 63 of which (81%) were located in intergenic regions, mostly overlapping known or putative promoters. The majority of the Eσ⁵-bound intergenic regions was located upstream of genes (encoding either ORFs or non-coding RNAs) showing some degree of dependence on the σ⁵-encoding rpoS gene. The remaining Eσ⁵-bound intergenic regions overlapped promoters recognized by RNA polymerase associated with the housekeeping σ⁷⁰ factor, or even with the alternative σ⁶ and σ¹ factors. Transcription of this latter set of genes does not appear to be affected by rpoS inactivation. Thus, consistent with previous observations, our
results indicate significant overlap in promoter recognition between $\sigma^S$ and other σ factors.

In particular, our results suggest that, in conditions leading to $\sigma^S$ accumulation, E$\sigma^S$ significantly contributes to transcription of genes involved in LPS biosynthesis not belonging to the \textit{rpoS} regulon during transition from exponential to stationary phase.

Finally, our results highlight a direct role of E$\sigma^S$ in the regulation of non-coding RNAs
Background
Helicobacter pylori is a human pathogen surviving in the prohibitive gastric environment where it can cause cancer. The ability to counteract environmental stresses and host responses is a key feature for persistence and depends on the rapid and fine-tuned expression of the genetic repertoire orchestrated by a shallow Transcriptional Regulatory Network. Metal-dependent regulators NikR and Fur are key hubs in this network; they directly control metal ion homeostasis and indirectly impact on several processes involved in H. pylori virulence and survival.

Objectives
In this work we combined ChIP- and RNA-sequencing to gain a comprehensive view of the NikR regulome. Moreover we investigated NikR regulation in presence or absence of Ni$^{2+}$ in wild-type and in nikR or fur knock-out strains, aiming to elucidate the plasticity of the system and the level of cooperation and antagonism between these two transcription factors.

Methods
For each experimental condition we determined NikR binding regions and defined the link between each binding site and its putative gene/operon target. Then we performed a differential peak calling to classify basal, Ni$^{2+}$ driven and Fur coordinated bindings. In parallel differentially expressed genes, operons and ncRNAs were identified. Finally, we discriminated direct and indirect effects of NikR action by integrating ChIP-seq and RNA-seq results.

Conclusions
Our study confirms known targets identified through single-gene analysis and sheds light on the entire framework of dynamic interactions between NikR and its regulatory targets, unraveling new explanations for its pivotal function in the persistence and pathogenicity of the bacterium.
Global regulatory networks

COMPLEMENTATION AND PHENOTYPIC CHARACTERIZATION OF MUTANTS FOR NUCLEOID ASSOCIATE PROTEINS (NAPS) IN SALMONELLA ENTERICA SEROVAR ENTERITIDIS

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Background
Salmonella Enteritidis is the most prevalent serotype in foodborne outbreaks since 1990 associated to poultry meat, eggs and products. The phagotype 4 is among most commonly isolates associated with diarrhea and systemic infection. Due to importance of this pathogen in human and animal health, researchers are searching ways to control S. enterica infections using vaccines. Despite the efforts of many groups, a few live vaccine strains are released for use in the salmonellosis control and there are no human vaccines to serovar Enteritidis yet.

Objectives
Based on these data, and tacking account that motility is an important virulence trait to Salmonella enterica, the goal of this study was evaluate the motility phenotype of S. Enteritidis PT4 null mutants to genes coding to Naps and their complemented strains in comparison to wild-type strain (WT).

Methods
The single mutants to NAPs were constructed by λ Red methodology and the complementation was performed using the cloning vector pACYC184. The motility was determined by measurement of bacteria migration from the center to the periphery of Luria Bertani 0.35% agar plates.

Conclusions
The data were analyzed by test-T using the software Assistat version 7.7. The S. Enteritidis PT4 mutants showed a significant decrease in motility (20-55%) compared to wild-type strain (p<0.01). The complemented mutants reversed the phenotype, presenting motility similar to WT Thus, we concluded that deletion of these genes affects motility, suggesting that they act directly or indirectly in the regulation of motility-related genes.
Background
In the context of the fast development of aquaculture, infections by Flavobacterium psychrophilum are a major sanitary concern for the salmonid farming industry worldwide. Juvenile salmonids are particularly susceptible to this Gram-negative aerobic bacterium, which is able to survive outside the fish during long periods of starvation in freshwater environments.

Objectives
Transcriptional regulatory networks are crucial for bacteria to maintain their integrity and to adapt to changing growth conditions. Alternative sigma factors confer promoter selectivity by binding to the bacterial RNA polymerase and allow switching between various adaptative responses. How F. psychrophilum adapts to environmental conditions has been investigated at the molecular level by analysis of its primary transcriptome and in silico prediction of regulons controlled by various sigma factors.

Methods
The transcriptional landscape and the transcription start sites (TSS) of the virulent strain THCO2/90 isolated from a Coho salmon, have been established using RNAseq technology by analysing a pool of bacterial RNAs extracted from cells undergoing various growing conditions.

Conclusions
Expressed genes as well as putative expression signals and regulatory elements were listed (i.e. TSS, 5'-untranslated regions, non-coding and antisense RNAs). In silico searches for consensus promoter sequences upstream of identified TSSs revealed more than a thousand sigma-70 dependent promoters as well as three other promoters groups responding likely to alternative sigma factors. Building this detailed transcriptional map represents the first attempt to characterize regulatory networks of F. psychrophilum, an important pathogen of the Bacteroidetes phylum.
Background

*Burkholderia xenovorans* LB400 is a model aromatic compounds-degrading bacterium. Aerobic metabolism of aromatic compounds may produce oxidative stress in bacteria. However, the molecular mechanisms involved are not well known.

Objectives

The aims of this study were to characterize bacterial stress response in *B. xenovorans* LB400 during oxidative stress and aromatic metabolism.

Methods

Stress-related genes were predicted using bioinformatic tools and functional assays were performed by proteomics using high-resolution mass spectrometry and by transcriptomics using qRT-PCR. Oxidizing conditions included exposure of LB400 cells to paraquat and hydrogen peroxide. In addition, transcriptional analysis of stress-related genes from strain LB400 grown on aromatic compounds was performed.

Conclusions

Paraquat induced ferritin DPS-family DNA binding protein (DpsA), alkyl hydroperoxide reductase, fumarase C, catalase, glutathion peroxidase, oxygen peroxide resistance protein (Ohr) proteins. In addition, induction of diverse reducing enzymatic systems such as ferrihins and flavodoxins was observed. During hydrogen peroxide exposure, two alkyl hydroperoxide subunits and DpsA protein were induced. A transcriptional analysis indicated that during paraquat and hydrogen peroxide exposure, an up-regulation of *ahpC*, *katE*, *fumC* and *oxyR* genes was observed. During growth of strain LB400 on 3-hydroxyphenylacetate, the expression of *ahpC*, *sod*, *acn*, *fumC* and *oxyR* genes was up-regulated. The expression of *ahpC*, *sod*, and *oxyR* genes were up-regulated during growth on 4-hydroxyphenylacetate. In conclusion, this study illustrated diverse oxidative stress proteins from *B. xenovorans* LB400 that are active during oxidative stress and aromatic compounds degradation.

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Global regulatory networks

EXPERIMENTAL EVOLUTION WITH A GLOBAL REGULATOR MUTANT IN ESCHERICHIA COLI

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Background

The CsrA regulatory protein is the main actor of the carbon storage regulator (Csr) network, a well-conserved regulatory network in bacteria. It acts at the post-transcriptional level by binding specific sequences on target mRNAs, thereby modulating gene expression. This global regulator controls several physiological processes, including central carbon metabolism and social behavior. Insights into CsrA roles were mostly obtained with an E. coli K-12 csrA mutant, encoding a truncated form of CsrA that retains partial activity.

Objectives

This work aims at further characterizing the roles of CsrA by analyzing phenotypes presented by an E. coli strain deleted for the csrA gene.

Methods

The csrA gene was deleted by homologous recombination in the uropathogenic E. coli CFT073 strain. Experimental evolution was performed. Three independent evolved mutants were extensively analyzed by combining molecular techniques, microscopy and global approaches, including comparative proteomics and whole genome sequencing.

Conclusions

Deletion of csrA leads to severe growth defects and altered cell morphology, indicating that this gene, although not essential, is important for growth. Evolved mutants were rapidly selected (2-3 days of culture in LB medium), showing that deleting the csrA gene constitutes a strong selective pressure. Our data confirms that
csrA deletion strongly affects central metabolism and energy status. In addition, our data show that specific stress responses are constitutively induced in the deletion mutant, including phosphosugar, oxidative and envelope stress responses. This highlights the interconnection of multiple regulatory networks and their flexibility to compensate for genetic perturbations in *E. coli*.
TWO ISOFORMS OF YJJM TRANSCRIPTION FACTOR DIFFERING IN THE DNA-BINDING DOMAIN.

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Background

Ashwell and Entner-Doudoroff pathways provide an alternative to glycolysis being important for bacterial colonization and motility. There are several specific proteins (UxuR, ExuR, UidR and cAMP-CRP) sensing metabolic status of the cell and switching gene expression to the optimal pathway. YjjM was also predicted to be involved in this network.

Objectives

The goal of this study was to investigate yjjM transcription and modes of regulation.

Methods

Alignments and structural predictions were made using BLAST, TCoffee and Phyre 2. Promoters were mapped with PlatProm software, single round and 5'-RACE. Expression level was measured by beta-galactosidase assays and qRT-PCR. Proteins specifically bound to the promoter DNA were detected using pull-down assays and mass-spectrometry. YjjM products were identified by Western-blot.

Conclusions

YjjM structure is assumed to be very similar to other regulators of the GntR family, with oligomerization/effector binding C-terminal and DNA-binding N-terminal domain. Computational and experimental data indicated that yjjM may have two ORFs resulting in synthesis of two proteins variable in their DNA-binding domain by 36 aminoacids. As a basis for this heterogeneity, four possible promoters were mapped nearby the yjjM 5'-end, with two of them located within the main ORF. Gene expression analysis, Western-blotting and pull-down assays revealed that regulatory switch from one protein product to another may occur due to the complex interference between RNA-polymerase alpha subunits and transcription factors operating in the yjjM regulatory region. Thus in changing growth conditions, different promoters may tune-up DNA-binding properties of YjjM by switching the ratio of its isoforms.
THE INTEGRATED RESPONSE OF GENOME ORGANIZATION AND ACTIVITY TO ENVIRONMENTAL CUES

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Background
Loop formation is key to the global organization of genomes in organisms from all three domains of life. Genomes are organized dynamically and their re-modelling is implied in translating external signals into specific gene products. The bacterial chromatin protein H-NS is capable of forming bridges between DNA segments in vitro and thus a prime candidate for genomic loop formation in vivo. H-NS is also a global regulator of transcription.

Objectives
We have investigated how the DNA bridging activity of H-NS is fine-tuned by salt conditions and interaction with co-regulators Hha and YdgT.

Methods
We demonstrate, using a combination of biophysical (Tethered Particle Motion), biochemical (in vitro bridging assays) and computational (Molecular Dynamics simulations) techniques, that there is a delicate balance between two DNA binding modes (DNA stiffening and bridging), which can be shifted by environmental cues or H-NS interaction partners.

Conclusions
Our observations yield models for fine-tuning of genome organization and for the translation of changes in environmental conditions into transcriptional activity.
Background The thermoacidophilic crenarchaeon Sulfolobus acidocaldarius can utilize different carbon sources including maltose. In Archaea, sugar uptake is achieved by ABC transporters that are inducible by the corresponding sugars. How these transporters are regulated in Sulfolobales is an open question, since no regulatory proteins have been identified for any sugar ABC transporter so far. The regulation of the maltose uptake and utilization system is of special interest, since this system lies at the basis of the successful inducible expression system in S. acidocaldarius.

Objectives The aim of this study is to identify and characterize the regulator of the maltose gene cluster in S. acidocaldarius and use this knowledge to improve the S. acidocaldarius protein expression system.

Methods The maltose regulator (MalR) of S. acidocaldarius was identified based on its sequence similarity to proteins of the TrmB family of regulators. To confirm the role of MalR growth of a deletion mutant was monitored on different carbon sources. Quantitative RT-PCR was used to examine transcription levels of the maltose gene cluster in a maltose-induced or uninduced state. Electromobility shift assays showed that MalR binds the promoter of the maltose binding protein (malE). Improvement of the current protein expression system was achieved by cloning an additional copy of malR into the expression plasmid.

Conclusion The maltose regulator of Sulfolobus acidocaldarius was identified. It positively controls transcription of the mal operon harboring genes for maltose uptake and metabolism. With the gained knowledge about the maltose regulator we could improve the maltose inducible expression system in S. acidocaldarius.
FEMS-2100
High-throughput approaches

RAPID BACILLUS PUMILUS GROUP SPECIES DIFFERENTIATION USING MALDI-TOF-MS AND FTIR-ATR
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Background
Bacillus pumilus group members, which encompass the closely related Bacillus pumilus, Bacillus safensis, Bacillus altitudinis, Bacillus xiamenensis and the newly described Bacillus invictae, possess a high medical, biotechnological and industrial relevance, stressing the need for reliable, easy and rapid methodologies for their correct identification.

Objectives
In this study we demonstrated the suitability of Matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF-MS) and Fourier transform infrared spectroscopy with attenuated total reflectance (FTIR-ATR) to accurately and rapidly identify the five closely related species enclosed within B. pumilus group, which are difficult to differentiate by conventional and rDNA 16S sequencing methodologies.

Methods
MALDI-TOF-MS and FTIR-ATR were applied to discriminate a clonally diverse collection comprising B. pumilus (n=5), B. safensis (n=19), B. altitudinis (n=1), B. xiamenensis (n=1) and B. invictae (n=5). Snapshots of different protein composition were acquired with a MALDI-TOF/TOF mass spectrometer, operating in linear positive mode, within the ion range at m/z 2000–12000 and using α-cyano-4-hydroxycinnamic-acid as a matrix. Moreover, the whole-cell content spectra were acquired using a FTIR-ATR System with a PIKE accessory, between 4000 and 400 cm⁻¹, with a resolution of 4 cm⁻¹ and 32 scan co-additions. Both, mass and infrared spectra were analysed by multivariate data analysis.

Conclusions
The resulting unique protein profile and the differences gathered in the carbohydrates and phospholipids/DNA/RNA vibration regions obtained by these high throughput approaches, proved that MALDI-TOF-MS and FTIR-ATR are valuable tools for B. pumilus group species discrimination, and thus, should be promptly considered for
these species identification.
High-throughput approaches

ASSESSING THE IMPACT OF TRANSGENIC GRAPEVINE ROOTSTOCKS ON THE ENVIRONMENT USING NGS
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Background
Grapevine fanleaf virus (GFLV) is responsible for fanleaf degeneration, which is the most severe virus disease of grapevine worldwide. GFLV causes important economic losses by reducing grape yield, lowering fruit quality, and shortening the longevity of vines. Moreover, with the ban of nematicides and soil disinfectants, fallows over 8 years are the only mean to get rid of the disease in the soil. Amongst the new strategies developed to control fanleaf, genetic engineering to modify the rootstock relying on the concept of pathogen-derived resistance, is the most promising strategy to develop GFLV-resistant grapevines.

Objectives
These impact studies aimed to evaluate the potential environmental consequences of the presence of two transgenes (GFLV coat protein gene and nptII kanamycin resistance gene) expressed in GM rootstocks, the scion remaining wild-type.

Methods
This study was performed in the frame of a French project (ANR IMA-GMO), that will take advantage of next generation sequencing (454 and Illumina MiSeq) technologies and quantitative PCR, to analyze the potential impact of GM plant growth either via expression of transgenes or via possible transgene transfer events (horizontal gene transfer) to microbial populations from a reference soil (Rothamsted Appletree, UK) or a vineyard soil (Bergheim, F), irrespective of the cultivability of these microbes.

Conclusions
Preliminary results obtained from confined experiments show no significant major modifications in the structure of the bacterial populations, and no horizontal gene transfer. Our results further stressed the necessity to perform environmental impact evaluation in long-term experiments under real agronomical conditions, in open-field trials.
**Background**

Bacterial resistance to currently available antibiotics is becoming increasingly problematic. In the search for novel antimicrobial substances our strategy is to screen for substances which block essential bacterial pathogenicity mechanisms without killing the bacteria.

*Salmonella* is a major cause of food-borne diseases. It relies on two type III secretion systems (T3SS) for host cell invasion and intracellular survival inside the host cells. Selective disruption of these systems would block host cell invasion and leave the bacteria vulnerable to the host immune system.

T3SS are highly conserved across a broad range of Gram-negative bacteria. It is tempting to expect that compounds active against *Salmonella* T3SS will also affect other species such as *Yersinia*, *Shigella* and *Escherichia*.

**Objectives**

The aim of this project is to develop a screening assay compatible with high-throughput screening and use this assay to screen for anti-virulence compounds against *Salmonella* T3SS.

**Methods**

We adapted a standard secretion assay to 386-well ELISA format. Bacteria are grown directly in the plates in presence of small molecule compounds. Secreted effectors are adsorbed to the assay plate and subsequently quantified. Various methods for detection and quantification of secreted effectors were evaluated for their sensitivity and robustness.

**Conclusions**

We have developed a screening approach to search for anti-infective pathogenicity-blocking small molecule compounds, which do not affect bacterial viability. Using this screening assay, we screened multiple compound libraries for active compounds. Potential actives were thoroughly validated in additional assays. In depth testing of the most promising candidates is currently underway.
FEMS-0641
High-throughput approaches

INFLUENCE OF COLD AND OSMOTIC STRESS ON TRANSCRIPTION AND TRANSLATION IN ESCHERICHIA COLI O157:H7 SAKAI
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Background
The enteric pathogen Escherichia coli O157:H7 Sakai (EHEC) expresses different sets of genes dependent on growth conditions. Growth at environmental conditions or in food comprises other challenges than growth in the host. Commonly, transcription intensities are compared between different conditions to obtain information about specific gene regulations. Indeed, changes in transcription levels are informative and have resulted in interesting findings in the past. However, many examples of poor correlation between mRNA and protein level exist due to translational or posttranslational regulation.

Objectives
In the past, transcriptomes for defined sets of genes were derived by microarrays. Today, RNAseq allows investigation of complete genomes with wider dynamic range and higher sensitivity compared to the latter technique. In addition, translationally regulated genes can be identified by ribosomal footprinting, where a snap shot of ongoing translation is obtained (translatome).

Methods
Strand-specific transcriptomes and translatomes of EHEC were analyzed at two different conditions (control: BHI, 37°C; stress: BHI, 14°C, a_w 0.976) to examine which genes show altered expression and to distinguish transcriptional from translational regulation. For translatome sequencing ribosomal movement is stopped by chloramphenicol, unprotected mRNA is digested by RNases, ribosomes are isolated, and purified mRNA is sequenced.

Conclusions
Differences in transcription point, as expected, towards cold and osmotic stress inducible genes. However, the additional translation profiles obtained complete the picture. Examples of translationally regulated genes are presented which do not show changes at the mRNA level, but altered numbers of ribosomal footprints. The regulation might be achieved by, e.g., RNA thermometers.
High-throughput approaches

HIGH-THROUGHPUT SCREENING OF LACTOBACILLUS SPECIES FOR ANTI-MICROBIAL ACTIVITY COMBINED WITH A COMPARATIVE GENOMIC APPROACH

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Background

Lactobacilli are frequently isolated from fermented food-products and are used in starter and protective cultures to control fermentation and avoid spoilage.

Objectives

To identify genes involved in fermentation performance of lactobacilli using broad genotypic/phenotypic screenings with focus on antimicrobial activity.

Methods

A new high-throughput screening technique was developed to rapidly detect antimicrobial activity, using an agar diffusion assay in microtiter plates. All secondary effects like low pH, organic acids or peroxide production were controlled with buffering the media and catalase treatment. Heat and protease treatment were performed to identify bacteriocins. The approach allows screening of test conditions with 10'000 datasets per day, 5'000 antibacterial interactions per day and 3'000 antifungal/antiyeast interactions per day.

We screened 500 Lactobacilli strains from 22 species mainly derived from food environments. A total of 28 conditions were tested including peroxide, bile salt and heat resistance. Further antimicrobial activity was analyzed. We found 65 bacteriocin-like producing candidates and 25 candidates displaying antifungal or antiyeast activity. The antimicrobial activity was diverse and activity against all 12 tested Enterococcus strains, all 3 tested Listeria species and against Aspergillus, Candida and Kluyveromyces isolates was observed. In a next step pooled DNA samples of active an inactive strains were sequenced and comparative genomics performed to identify genes that are responsible for antimicrobial activity and for other tested phenotypes.

Conclusions
The high-throughput methods allow us to screen lactobacilli with minimal laboratory equipment at low costs within short time, bringing phenotypic analyses in pace with next generation sequencing.
UNRAVELING THE PATHWAYS OF AMYLOID TOXICITY IN A BACTERIAL MODEL SYSTEM: THE REPA-WH1 PRIONOID

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Background

Mitochondria are recurrently found to be central in protein amyloidosis. Thus to study mitochondrial physiology in response to protein amyloidosis is a hotspot in the amyloid research field. Since this is a quite challenging task, we have experimentally approached mitochondria in the bacterial model RepA-WH1 prionoid.

We had reported that RepA-WH1, a bacterial protein conformational switch (dWH1/mWH1) functional in plasmid replication, causes an amyloid proteinopathy in Escherichia coli hampering cell proliferation. RepA-WH1 aggregates are vertically transmissible but not infectious (horizontally transmissible), and enable conformational templating by cross-seeding in vitro and in vivo. It is thus the first bacterial prionoid. We have recently found that DnaK, the Hsp70 chaperone in E.coli, modulates RepA-WH1 amyloidogenesis in vivo.

Objectives

Study the pathways of amyloid toxicity of RepA-WH1 combining genomic, proteomic and physiological approaches

Methods

We have analyzed through “omics” the gene expression pattern of cells expressing RepA-WH1 (Affimetrix microarrays) and the fraction of E.coli proteome co-aggregated with RepA-WH1 amyloid inclusions. We have also studied membrane integrity, oxygen consumption rates, iron and ATP levels and sensibility to oxygen reactive species (ROS) in cells expressing RepA-WH1.

Conclusions

Our results indicate that cell damage is produced by a ROS-dependent mechanism, most likely involving Fenton chemistry. These findings support a role for free iron in amyloid cyto-toxicity, possibly on top of membrane damage by amyloid oligomers as it has been proposed for general amyloid induced cyto-toxicity in human amyloidosis.

The results presented here underline the power of the bacterial RepA-WH1 prionoid as a synthetic minimalist model system for amyloid proteinopathies.
HIGH-THROUGHPUT SEQUENCING OF BACTERIA ASSOCIATED WITH COMMERCIIALLY IMPORTANT PELAGIC FISH

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Background

As the global demand for fish is increasing, and many marine species are fully exploited, a sustained profit in the industry has to result from improved quality and better utilisation. In Norway, the total volume of captured pelagic fish exceeded 1.1 million metric tonnes in 2013, giving a first-hand value of 730 million USD (4.8 billion NOK), where Atlantic mackerel (Scomber scombrus), Atlantic herring (Clupea harengus), blue whiting (Micromesistius poutassou), and capelin (Mallotus villosus) were the most important species. FAO estimates that 10-25 % of all captured fish are lost post-harvest, where microorganisms are of great importance in deterioration of fish products. Currently, bacteria associated with fish are mostly identified by culture-dependent methods. New approaches are needed to understand the complexities of bacterial communities and their influence in fish products, as most bacteria are unculturable.

Objectives

To describe the bacterial community composition of gills, muscle tissue, skin, intestine, and stomach content of pelagic fish. In addition, examine water samples and zooplankton from the area of catch.

Methods

Samples were obtained during commercial catch of important pelagic fish. Bacterial DNA were extracted by DNeasy Blood and Tissue Kit, amplified using NEXTflex 16S Amplicon-Seq Kit, and sequenced by the Illumina platform technology, prior to data processing. Results will be discussed.

Conclusions

The information gathered in this study is important for a better understanding of the microbiology of pelagic fish species. This may again support a reduction of post-harvest losses and increase the profitability of the Norwegian pelagic fish industry.
SYNTHETIC BIOLOGY FOR THE DESIGN OF ANTIMICROBIAL PEPTIDES

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Background

The number of multi drug resistant pathogens is constantly growing and novel antibiotic substances are desperately needed in order to at least maintain the status quo. Ribosomally synthesized antimicrobial peptides are not yet exploited for human applications despite an indisputable potential. Among those, lantibiotics, a class of posttranslationally modified and thioether-ring bearing antimicrobial peptides, show desirable properties as high but specific antimicrobial activity and excellent stability.

Objectives

We follow a synthetic biology approach for the generation of novel lantibiotics employing the blueprint of natural lantibiotics. Based on their structural and functional features, natural lantibiotics are dissected into modular subunits. These subunits are then shuffled to generate thousands of novel, putative active, chimeric lantibiotics. The lantibiotic genes are subsequently produced by combinatorial DNA de novo synthesis. These libraries will then be screened for molecules with high antimicrobial activity.

Methods

To enable for high screening rates we developed a platform based on nL-sized reaction vessels (nL-reactors) that are used for peptide production and activity-screening in a single step and at rates of $10^5$ variants per day. Library cells are grown to microcolonies within nL-reactors along with a sensor strain. Cells secreting an active antimicrobial peptide will deactivate the sensor cells. Clearance of an nL-reactor from the sensor thus indicates the presence of a strain secreting a highly active peptide.

Conclusions
$10^4$ different chimeric variants derived from the module-shuffling of ten natural lantibiotics were analyzed. Several interesting antimicrobials have been isolated, demonstrating for the first time the generation of new-to-nature lantibiotics by module-shuffling.
FEMS-2320
High-throughput approaches

LACTOCOCCAL SURFACE PROPERTIES AND THEIR ROLE IN MICROBE-MATRIX INTERACTIONS IN DAIRY PRODUCTS

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Background
Microbial surface properties are highly variable and they are considered to be important for the interactions of the bacterial cell with the product matrix. Lactic acid bacteria are industrially used for the fermentation of milk to produce, e.g., yoghurt or cheese. In fresh milk, bacteria encounter particles such as casein micelles or fat droplets, while in fermented milk products, the proteins form a semi-solid casein matrix, in which fat globules and whey pockets are present. However, the extent to which the interaction of dairy starter cultures with the various matrix components is important for the result of fermentations is largely unknown.

Objectives
To which extent do bacterial cell surface properties influence interactions with the matrix in a fermented dairy product? Does it impact on starter functionality and what are the underlying mechanisms?

Methods
ZetaSizer was used to measure cell surface charge, microbial adhesion to Hydrocarbons (MATH) method - to measure surface hydrophobicity, and Flow Cytometry - to measure cell attachments to milk proteins.

Conclusions
Here, we analyzed the surface properties of 55 L. lactis strains and found big variations in bacterial cell-charge, surface hydrophobicity and cell attachment to milk proteins. As we also have the full genome sequences of these 55 strains, genotype-phenotype matching was used to identify mechanisms involved in the described surface properties. The identified properties and mechanisms and their relevance in relation to microbe-matrix interactions will be discussed.
IS THE PREVALENCE OF CYCLOSPORA CAYETANENSIS IN HIV INFECTION UNDERESTIMATED?

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Objectives: Cyclosporiasis can occur as a locally-acquired infection, among travelers or in patients with HIV/AIDS. In our country, the prevalence of Cyclospora cayetanensis are not clear, and researches about this microorganism in HIV infected individuals are limited. The aim of this study was to determine prevalence of Cyclospora cayetanensis in symptomatic and asymptomatic HIV infected individuals with conventional and molecular methods.

Methods: One-hundred stool samples of HIV infected individuals were included in this prospective clinical study. The stool samples were investigated by modified Ziehl-Neelsen staining and Real-Time Polymerase Chain Reaction (PCR) methods to determine the existence of Cyclospora cayetanensis. Permission to conduct this study was obtained from the local ethics committee of Istanbul University Istanbul Medical Faculty, Turkey.

Results: Cyclospora cayetanensis was found positive by Real-Time PCR in two samples (%2), and in one sample was found positive by modified Ziehl-Neelsen staining (%1).

Conclusions: As cyclosporiasis is an endemic disease in our country, Cyclospora cayetanensis should be considered in clinical cases, especially in immunocompromised or have long-term diarrhoea patients. In terms of cyclosporiasis, laboratory request would be appropriate in not only immunocompromised patients but also have a healthy immunity individuals. Also, patients with a history of overseas trips to endemic areas, a history of recurrent or long-term diarrhoea and abdominal pain complaints might be examined for Cyclospora cayetanensis. The use of Real-Time PCR seems logical, faster, efficient and more sensitive method for the laboratory diagnosis of Cyclospora cayetanensis according to microscopic methods in our study.
EVOLUTION AND PHYLOGENY OF PHYTOPLASMA ASSOCIATED WITH LEMON WITCHES’ BROOM AS A FIRST RECORD FROM EGYPT

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Background

Witches' broom disease (WB) is a serious disease of acid limes. The first report of the disease was in Oman more than 30 years ago and has since spread to the United Arab Emirates, Iran and India where the destruction of 98% of lime trees occurred and has become a major limiting factor for lime production.

Objectives

The application of nested-PCR assays, the use of conserved sequences and sequencing the new emerged strains allow to broadly detect the evolution and virulent strains in field.

Methods

Symptoms of the Phytoplasma Associated with (WB) infection were observed on limes for the first time in Egypt. The disease was biologically characterized. Cytopathological detection referred to Diene's stain was used to differentiate the phloem tissues of leaf sections from infected trees. Transmission Electron Microscopy (TEM) indicated the presence of phytoplasmas and were detected by the specific amplification of their 16S-23S rRNA gene. Polymerase chain reaction (PCR), utilizing the universal phytoplasma-specific primers (P1/P7 followed by R16F2n/R16R2). The amplified PCR fragment of 16S rRNA gene was successfully cloned and sequenced. The nucleotide sequences were published to the GenBank with accession number "KJ948653" and named as Lemon Witches' broom-Egyptian strain (LWB-Eg strain). Phylogenetic analysis was performed to identify the LWB-Eg strain through the alignment with the corresponding phytoplasma strains available in the GenBank.

Conclusions

This study discovered and identified the phytoplasma strain of LWB-Eg \textit{Candidatus Phytoplasma aurantifolia} as a natural evolved causal agent from Lime WB strain from Oman (EF186828, 16SrII) which induces severe Witches' broom epidemic disease on lemon in Egypt.
ENTEROCOCCUS FAECIUM GENOME DYNAMICS IN LONG-TERM PATIENT GUT COLONIZATION

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Background

Since the mid-1980s Enterococcus faecium, normally a harmless gut commensal, has emerged as an important nosocomial pathogen that has acquired multiple antibiotic resistances. Patients can be colonized with multidrug-resistant E. faecium during hospitalization. These bacteria can remain present in the intestinal tract for a long period of time after hospital discharge.

Objectives

Our aim is to analyze the genome dynamics of multi drug-resistant E. faecium isolates that colonized the gut of patients over a large timespan.

Methods

The genomes of 96 isolates obtained over 8 years from 5 different patients were sequenced. We generated a core genome-based phylogenetic tree with these genomes, together with 70 previously published E. faecium genomes of diverse origins. Furthermore, we investigated highly similar subsets of patient isolates to describe micro-evolution in closely related strains.

Conclusions

Whereas some patients were colonized for almost 2 years with a clonal population of E. faecium, other patients were colonized by a diverse population of E. faecium strains. Almost all isolates could be assigned to a hospital-associated sub-population of E. faecium. By studying the pan-genome of the strains in this patient isolate dataset, we identified gene gain and gene loss events. We also found evidence for recombination events, and for transmission of E. faecium strains between patients during hospitalization. This study highlights the different mechanisms that contribute to the genomic flexibility of clinical E. faecium strains. This may be a crucial factor in the ability of E. faecium to rapidly adapt to new ecological niches.
Background

_Corynebacterium glutamicum_ is a well-known amino acid producer microorganism. L-glutamic acid and L-lysine have become bulk products inside an opened worldwide market to threonine or tryptophan. Thus, the portfolio of _Corynebacterium_ has been enlarged by relevant bioproducts such as biodegradable polymers and ingredients or additives in food, feed, cosmetics and pharmaceuticals. This new compounds production has been eased by the huge knowledge accumulated along the last decades in _C. glutamicum_, as well as the introduction of the genome sequencing facilities and other omics methodologies.

Objectives

The release of more than twelve genome sequences of _C. glutamicum_ allows the location of their genes, putative pseudogenes, transposones or pro-phage sequences. These data have permitted the selection of putative dispensable genes or genome regions. However, the recent studies on small RNA (sRNA) or the description of Actinobacteria cryptic secondary metabolite biosynthetic gene clusters have forced to review the idea of 'dispensable regions'.

Methods

A meticulous RNAseq study of _C. glutamicum_ transcriptome combined with bioinformatic analyses have allowed the update of the genome organization and the identification of confirmed intergenic regions free of possible genetic elements (sRNA, terminators and extended operons).

Conclusions

Consequently, a pool of these intergenic regions suitable for the integration of heterologous or homologous genes has been defined in _C. glutamicum_. In addition, several genes that enhance the oxygen uptake, primary metabolism precursors supply or the translation activity have been used to test the best region candidates. A set of validated _C. glutamicum_ intergenic regions have been defined for genetic engineering applications.
FEMS-1565
Microbial genomics, evolution, phylogeny

SIMILARITIES AMONG PSYCHROTOLERANT BACILLUS CEREUS SENSU LATO GROUP MEMBERS SUGGEST THEIR COMMON EVOLUTIONARY HISTORY

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Background
The Bacillus cereus group comprises seven bacterial species with different ecological properties and impact on human activity. Besides virulence and heat-resistance of spores, these bacteria often show psychrotolerance, being a serious problem in food processing. Until now, cold adapted bacilli were noted among isolates representing different taxa of the entire group, however precise relationships between them remain unknown.

Objectives
We decided to investigate (i) the frequency of psychrotolerant B. cereus group members, (ii) assess their enterotoxicity, (iii) genetic diversity and (iv) relationships between cold adapted isolates in the context of their origin.

Methods
B. cereus group strains (N=400) were PCR screened for the presence of the psychrotolerant form of the cspA gene and cultivated at 7°C to identify cold-adapted bacteria. Enterotoxicity was assessed by PCR technique and immunochromatographic Duopath Cereus tests. Next, 16S rRNA gene sequencing and MLST typing were applied to establish diversity and genetic relationships among tested strains. Dendrograms were constructed using NJ method in Mega 5 software and index of association was calculated using START 2 software.

Conclusions
Psychrotolerant isolates were found among B. cereus, B. weihenstephanensis, B. mycoides and B. thuringiensis from range of habitats in different frequency. They form one homogenous group in opposition to polymorphic mesophilic isolates, probably due to the slower multiplication and stabilizing effect of natural selection. Similarities in sequences of chromosomal genes and diversity of plasmid-borne features suggest that all the cold-adapted strains may have common evolutionary history.

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GENOME COMPARISONS REVEAL THE EVOLUTION OF NITROGEN FIXATION IN BURKHOLDERIA

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Background

The genus *Burkholderia* includes various rhizobial species capable of establishing nitrogen-fixing symbioses with certain legumes. The ability to biologically fix atmospheric nitrogen is conferred by the bacterial partner and may be transferred horizontally between species. Previous work suggests a unique evolutionary origin for the nitrogen fixing abilities of South African rhizobial species of *Burkholderia*.

Objectives

In this study we investigated the evolutionary history of the nitrogen fixation loci in South African *Burkholderia* bacteria with rhizobial properties by making use of genome data.

Methods

The publically available genomes of 16 *Burkholderia* species were used in this study. These included 4 South African rhizobia, 8 rhizobial species from other parts of the world, as well as 4 diazotrophic members of the genus. We also determined the genomes sequences for an additional three rhizobial species of *Burkholderia* from South Africa. The nitrogen fixation locus of these bacteria were identified and characterized, after which synteny was investigated and phylogenetic histories determined.

Conclusions

Genome comparisons revealed three distinct structures for the nitrogen fixation locus among the 19 bacteria examined. One of these was represented by the South Africa species, while the other two were respectively represented by the diazotrophs and the rhizobial species from other parts of the world. Phylogenetic analyses suggested distinct evolutionary origins for the three types of nitrogen fixation locus. Overall, our
data indicated that horizontal gene transfer, gene rearrangement and/or gene replacement contributed to the differences observed in the architecture of the nitrogen fixation loci within the genus *Burkholderia*. 
COMPARATIVE ANALYSIS OF THE GENOMES OF THE BACTERIA PANTOE A ANANATIS AND PANTOE A AGGLOMERANS

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Background

Pantoea ananatis causes diverse plant diseases, including fruit rot of pineapple, leaf blight of eucalyptus, brown stalk rot of maize, center rot of onion, and sheath rot of rice. This bacterium is also known as an opportunist human pathogen. Pantoea agglomerans causes bacteria palea browning of rice.

Objectives

In the present study, the sequenced genomes of the closely related bacteria P. ananatis and P. agglomerans were compared with emphasis on genome organization and coding capacity.

Methods

Whole genome DNA sequencing of bacteria was performed using Illumina sequencing via the synthesis method on an Illumina HiSeq 2000 genome analyzer.

Conclusions

The genomes of P. ananatis strain PA13 from diseased grains of rice, and strain HY02 from onion center rot consist of a single circular chromosome consisting of ~4.58 Mb and a circular plasmid (~0.28 Mb). The genome of P. agglomerans strain SW2 consists of a single circular chromosome consisting of 3.95 Mb and three circular plasmids, Pagg_1p (0.42 Mb), Pagg_2p (0.16 Mb), and Pagg_3p (94 Kb). Short collinear regions, which are identical to DNA sequences in the Escherichia coli MG1655 chromosome, were widely dispersed along the Pantoea genomes. The genome information and the mutant library will allow a better understanding of how Pantoea causes diseases on their plant hosts and can be utilized to develop effective control methods against these important plant pathogens.
ANALYSIS OF METHYLOTROPHIC METABOLISM OF PARACOCCUS AMINOVORANS JCM 7685 (ALPHAPROTEOBACTERIA)

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Background
Paracoccus aminovorans JCM 7685 (Alphaproteobacteria) is a facultative methylotroph, which utilizes a wide range of C1 compounds, including methanol, methylated amines and N,N-dimethylformamide. Whole genome sequencing and sequence analysis revealed that the strain contains several genetic modules presumably involved in methylotrophy. Interestingly, many of these modules are unique among bacteria of the genus Paracoccus spp.

Objectives
The identified genes, predicted to be involved in C1 metabolism, show only moderate similarity to the corresponding genes of other well-known methylotrophs. Therefore, we conducted a set of experiments to confirm their functions and to analyze their regulation.

Methods
The mutant strains of P. aminovorans were constructed with application of transposon mutagenesis and gene replacement strategy. Proteins upregulated during methylotrophic growth were identified by the comparison of protein profiles of the bacteria grown on various C1 compounds (SDS-PAGE), followed by mass spectroscopy of selected protein bands. Activities of promoters were studied under methylotrophic and non-methylotrophic conditions using β-galactosidase assay.

Conclusions
Methylotrophic metabolism of P. aminovorans differs highly from that of the closely related, well-studied methylotroph P. denitrificans (type strain of the genus Paracoccus). The studied strain encodes not only more enzymes enabling oxidation of wider range of C1 compounds but has also completely different central metabolism of C1 subunit assimilation (the serine cycle in P. aminovorans vs. the Calvin cycle in P. denitrificans). Almost all genes activated during methylotrophic growth are encoded within large extrachromosomal replicons (chromids pAMV1 and pAMV3). Therefore it is highly probable that these genes might have been horizontally acquired, building mosaic methylotrophy pathways.

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CHARACTERISATION OF PROPIONIBACTERIUM FREUDENREICHII THROUGH COMPARATIVE GENOMICS

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Background

*Propionibacterium freudenreichii* is an industrially important bacterium recognised for its well-documented role in ripening of Swiss-type cheeses, production of active form of vitamin B₁₂, capability to produce considerable amounts of propionate as well as for its recently acknowledged probiotic potential. Moreover, this species is characterized by hardiness and ability to adapt to various conditions such as those encountered in dairy and cereal environments. To this date only one genome sequence of *P. freudenreichii* has been published, hindering the possibility to fully explore its potential. *P. freudenreichii* is characterized by high G+C content and regions of repeated sequences, which has been an obstacle in unraveling the genomes of this species using Next Generation Sequencing methods such as Illumina and 454.

Objectives

In this project we sequenced twenty-five genomes of Propionibacteria associated with various environments to identify pathways contributing to adaptation and to industrially important traits of *Propionibacterium freudenreichii*.

Methods

The PacBio RSII sequencing technology was used as the method of choice, since it enables genomic library preparation without PCR amplification, and produces long reads (~10kbp) allowing for complete coverage of regions with repeated sequences. Genomes were annotated and analysed with various bioinformatic tools.

Conclusions

Whole genome sequencing of *P. freudenreichii* is feasible with the PacBio RSII method and the produced genomes will help to elucidate the true potential of the species.
Background
Ornithine-lipids (OLs) are prokaryote-specific phosphorus-free membrane lipids. A function for OLs in stress conditions and in host-bacteria interactions, either symbiotic or pathogenic, has been shown in some bacteria. A variety of hydroxylations in different structural positions and its stress resistance implications have been reported. Recently, for the first time the presence of N-methylated OL derivatives had been described in the planctomycete Singulisphaera acidiphila. However, the gene/enzyme responsible for the methylation in S. acidiphila remained elusive.

Objectives
Identify and characterize the novel OL-N-methyltransferase OlsG from S. acidiphila.

Methods
We identified twelve candidate genes from S. acidiphila encoding enzymes possibly responsible for methylated OL synthesis. Recently, our group identified the OL synthase OlsF from S. proteamaculans that upon expression in E. coli caused OL formation, so the twelve candidate genes were expressed one by one in this OL-forming E. coli strain and we analyzed the lipid composition of [14C]acetate labeled cultures of the resulting strains by two-dimensional thin-layer chromatography, leading to the identification of the novel methyltransferase OlsG, also confirmed by mass-spectrometry.
In vitro characterization shows that OlsG is responsible for the threefold OL methylation, resulting in monomethyl-OL (MMOL), dimethyl-OL (DMOL), and trimethyl-OL (TMOL). Time-dependency of OL-N-methyltransferase activity was observed.

\[ \text{OlsG} \text{ contains 192 amino acids, is soluble, requires S-adenosylmethionine (SAM). Maximal N-methyltransferase activity was observed at pH=8 and in presence of} \]
Conclusions

OlsG is responsible for a new type of OL modification. OlsG is the first OL-N-methyltransferase described.
GENOMIC ANALYSIS OF SPHINGOPYXIS (FORMERLY SPHINGOMONAS) MACROGOLGTABIDA STRAIN TFA REVEALS NEW INSIGHTS FOR SPHINGOPYXIS GENUS

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Background

TFA is a gram negative bacterium able to grow on tetralin as the sole carbon and energy source. Based on a partial 16S sequence it was initially ascribed to the Sphingomonas genus. Despite the exhaustive study of the tetralin degradation pathway, other genomic and metabolic characteristics of TFA remain unknown.

Objectives

The main objective is to establish the TFA metabolic map using a genomic approach and further experimental validation. New metabolic features are being experimentally validated.

Methods

Contigs resulting from the sequencing process were assembled. Prodigal software was used to predict protein-encoding genes, which were functionally annotated by Sma3s. The final annotation was analyzed with Pathway Tools software to generate a preliminary metabolic network. This analysis was complemented with antibiotics and secondary metabolites clusters prediction with antiSMASH. CMG-biotools, BRIG and IslandViewer were employed in comparative genomic analysis.

Conclusions

TFA has been re-assigned to the Sphingopyxis genus based on reciprocal identity comparisons versus members of the Sphingomonadaceae family. Highly similar regions between TFA genome and other α-proteobacteria have been detected. Despite no plasmids have been detected, plasmid-related genes have been annotated, suggesting the importance of horizontal transfer in the construction of TFA genome. Comparative genomic predicts 914 TFA genes absent in other Sphingopyxis. A complete nar genes cluster encoding a membrane-bound respiratory nitrate reductase and genes for cobalamin biosynthesis seem to be exclusive characteristics of TFA.
Anaerobic growth reducing nitrate has not been described in *Sphingomonadaceae* family before. Preliminary metabolic reconstruction suggests that TFA might be able to degrade other aromatic compounds.
MULTIPLE PATHWAYS FOR AEROBIC AND ANAEROBIC RESPIRATION IN MELIORIBACTER ROSEUS BELONGING TO THE NOVEL PHYLUM IGNAVIBACTERIAE

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Background
Ignavibacteria are phylogenetically closely related to Chlorobi but essentially differ from them by their facultatively anaerobic mode of metabolism and obligate organotrophy. Melioribacter roseus is one of the two cultured representatives of Ignavibacteriae which grows either by fermentation of sugars and peptides, aerobic respiration or dissimilatory reduction of arsenate, nitrite or Fe(III). Genome analysis highlighted key determinants of respiratory metabolism in M. roseus and provided insights into its adaptation to fluctuating environmental conditions. Full set of proton-translocating membrane complexes and several putative terminal oxidoreductases were identified in the M. roseus genome.

Objectives
We aimed at experimental evaluation of the role of these enzymes in energy metabolism of the organism and deciphering the mechanisms for aerobic and anaerobic respiration, basing on profound genomic analysis and experimental data.

Methods
Inhibitor analysis and direct membrane potential measurements were used to reveal the activity of electron transfer chain in the organism. Phylogenetic analyses of the catalytic subunits of revealed oxidoreductases, transcriptomic and proteomic studies were used to confirm the function of predicted oxidoreductases in M. roseus metabolism.

Conclusions
Activity of electron transfer chain under aerobic or Fe(III) respiration was revealed. Key role of a novel cc(o/b)c₃-type cytochrome oxidase and ACIII complex in aerobic respiration as well as the involvement of two non-Arr molybdopterin oxidoreductases and putative porin-cytochrome complex in anaerobic respiration were identified. Our results provide the basis for the reconstruction of the evolution of aerobic and anaerobic respiratory pathways in Chlorobi-Bacteroidetes-Ignavibacteriae group and expand the knowledge base on the novel phylum Ignavibacteriae.
Background
Aquatic systems are in constant transformations for the accelerated artificial eutrophication process that means nutrients enrichment. The artificial Paranoá Lake, located in Brasilia (Federal District), yet in constant eutrophication process, supplies a higher degree of humidity to the local dried atmosphere. The Funil reservoir, an eutrophic environment located in Resende (Rio de Janeiro), was constructed to normalize the flow of the Paraiba do Sul River and to attenuate the impact of floods. Blooms of cyanobacteria are being reported for both aquatic ecosystems.

Objectives
The aim of this work is to obtain the complete genome of two cyanobacteria strains previously classified as: CYLP, non-toxic, isolated from Paranoá Lake; and CYRF, a toxic strain, isolated from Funil Reservoir; and to assess the fundamental genetic differences between the strains.

Methods
A next generation sequencing protocol was performed to run in Ion PGM™ platform. DNA fragments size of 400 bp from both CYLP and CYRF strains were prepared and sequencing runs were performed. Reads were downloaded from Ion Server and trimmed (quality/size) using CLC Genomics Workbench v7.5, resulting in 709Mb for CYLP and 867Mb for CYRF. The best hit species from BLASTN results was against Cylindrospermopsis raciborskii. So, reads were mapped against the reference sequence CS-505 (ACYA01000000). Reads were assembled to contigs using MIRA and the final assembly was performed using CONTIGuator.

Conclusions
Two strains with different toxicities isolated from two Brazilian aquatic ecosystems were successfully sequenced. The assembly generated about 50 contigs. Functional analysis is being performed.
Background
The microbial communities in the small intestine rely on their capacity to rapidly import and ferment available carbohydrates to survive in a complex and highly competitive ecosystem.

Objectives
Understanding how these communities function requires elucidating the role of its key players and the interactions among them and with their environment/host. Hence, we present the complete genome of Romboutsia ilealis CRIB¹, a natural inhabitant and key player of the small intestine of rats.

Methods
The complete genome of R. ilealis CRIB¹ was sequenced and annotated, followed by a whole-genome transcriptome analysis after growth on specific carbohydrates (glucose, L-fucose and fructo-oligosaccharide [scFos]).

Conclusions
R. ilealis CRIB¹ possesses a circular chromosome of 2,581,778 bp and a plasmid of 6,145 bp, carrying 2351 and eight predicted protein coding sequences, respectively. Analysis of the genome reveals only limited coding capacity to synthesize amino acids and vitamins. However, multiple and partially redundant pathways for the utilization of different relatively simple carbohydrates are present in the genome. The transcriptome analysis allows pinpointing the key components in the degradation of glucose, L-fucose and fructo-oligosaccharides. This reveals that R. ilealis CRIB¹ is adapted to a nutrient-rich environment where carbohydrates, amino acids and
vitamins are abundantly available. This work shows how genome mining and functional analyses with single microbes can provide an insight in the functional potential of new members of the intestinal microbiota, and will help us with understanding the microbial communities in us.
MLST TO RESOLVE DISCREPANCIES IN THE IDENTIFICATION OF
CRONOBACTER SPECIES
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Background

Differentiation of closely related species, such as those in the Cronobacter genus can be difficult. Previous characterization of Cronobacter isolates by phenotyping, ribotyping and 16S rRNA sequencing revealed discrepancies in the identification of certain strains (1).

Objectives

This study used multilocus sequence typing (MLST) to resolve these discrepancies in the identification of Cronobacter species.

Methods

MLST analysis was performed as previously described (2). Species identifications based on MLST were compared to previously reported phenotyping, ribotyping and 16S rRNA sequencing results (1).

Conclusions

MLST confirmed ribotyping and 16S rRNA sequencing results by identifying strain 05CHPL02 as C. dublinensis and strain 05CHPL53 as C. sakazakii. These strains were misidentified biochemically. Strains 05CHPL46 and 07HMPA87A were identified as C. malonaticus, confirming the phenotyping and ribotyping results. Both strains were misidentified as C. sakazakii by 16S rRNA sequencing, likely due to the close relationship of these two species. The identification of strain 05CHPL65 as C. sakazakii by phenotyping and ribotyping was confirmed by MLST, though it was misidentified by 16S rRNA sequencing. Finally, strain 05CHPL47 was identified as C. malonaticus by MLST, whereas the other methods identified it as C. sakazakii.

Phenotyping, ribotyping and 16S rRNA sequencing limited in their use for the identification of Cronobacter spp. DNA sequence-based methods, such as MLST, are much more reliable for the identification of Cronobacter isolates.
References
IS THE RIBOSOME TARGETED BY ADAPTIVE MUTATIONS?
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Background

RNA polymerase and ribosomes, composing the macromolecular synthesis machinery (MMSM), carry out the central processes of transcription and translation, but are usually seen as mechanical elements with no regulatory function. Extensive investigations of gene regulation and the high degree of evolutionary conservation of the cellular MMSM tend to support this view. However, under certain selective conditions the machinery itself may be targeted by adaptive mutations, which result in fitness increasing phenotypic changes. Here we investigate and characterise the role of ribosomal mutations in adaptive evolution.

Objectives

To elucidate the consequences of mutations in ribosomal protein genes in infecting bacterial isolates from human lungs.

Methods

Several mutations in ribosomal genes have been identified in the genome analysis of nearly 700 P. aeruginosa isolates from infected cystic fibrosis patients. Among these mutations we have repeatedly identified insertions, deletions and substitutions in specific ribosomal genes. Preliminary assays show that mutant strains have reduced growth rate and an altered antibiotic resistance pattern. The selection for mutations in ribosomal protein genes is partly explainable by the antibiotic treatment of the patient. But other mutations cannot be directly associated with antibiotic resistance. The bacterial phenotypes of the mutated strains will be investigated.

Conclusions

Clarification of the potential pleiotropic consequences of the specific mutations in ribosomal proteins is important for our understanding of biological evolution, and will have impacts on the design of new treatment strategies to combat microbial infections.
EXPRESSION ANALYSIS OF LACCASE MULTIGENE FAMILY IN TRAMETES HIRSUTA
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Background
Fungal Laccases (EC 1.10.3.2) are intensively studied due to their wide biotechnological application. However, the laccases industrial use is still limited in knowledge of enzymes biogenesis and its regulation. The regulation of laccase biosynthesis at the transcriptional level is suggested to occur via large number of specific responsive elements in the promoter regions of genes.

Objectives
The work is aimed to study the regulation of multigene laccase family expression in Basidiomycete Trametes hirsuta – an effective lignin degrader.

Methods
The full-length cDNA sequences for 7 laccases from T.hirsuta, using NGS techniques and RACE-PCR has been obtained. Analysis of laccases promoter regions (with MatInspector software) revealed different composition of cis-elements, including MRE, ACE, XRE, creA etc. Transcriptional dynamics for individual laccase genes were studied under application of different effectors (e.g. copper ions, different groups of xenobiotics) and confirmed with secretome analysis. The laccase complex activity was monitored during the cultivation under all conditions studied.

Conclusions
The major isozyme LacA was found to be expressed in all investigated media, and apparently represented a constitutive laccase form. The expression profiles of six other laccase isozymes and laccase complex activity were different and changed depending on effectors used as well as growth phase. Analysis of transcriptional patterns and modeling of the amino acid structures of laccase isozymes allow suggesting physiological roles and functions of individual laccases, as well as biogenesis regulation within this family.

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INTERACTION OF BOVINE HERPESVIRUS (BHV-1) AND BACILLUS SPP.

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Background
It is well known that some viruses can integrate to genome of eukaryotes. As for prokaryotes there is a special group of viruses – bacteriophages, that can interact with them. But there are almost no information about interaction of viruses pathogenic for animals and bacteria.

The aim of this work is to study possibility of integration of BHV-1 genome and bacillus.

Methods
Strains of bacillus were extracted from feces of cattle by standard microbiological procedure.
Screening for antigens of BHV-1 in bacilli was performed by reaction of agglutination (RA test) of bacteria with positive BHV-1 serum.
Screening for genome was done by routine and real-time PCR.
Test for induction of immunity was made by immunizing experimental group of mice with isolates of bacilli twice in 7 days interval. Control group 1 was immunized with vaccine strain of BHV-1, Control group 2 – with sterile saline. Specific antibodies were indicated in reaction of neutralization on MDBK cell culture.
For identification of bacilli automatic biotyper “Vitek” was used.

Results
From 70 isolates of bacilli 20 showed positive RA test. And 3 out of this 20 were positive in PCR for glycoproteins D and B genes of BHV-1.
Serums of experimental group of mice were positive for BHV-1 antibodies in dilution up to 1:40, control group 1 - up to 1:60, control group 2 - no antibodies.
Positive bacilli stains were identified as Bacillus lisheniformis.

Conclusions
Obtained results show possibility of interaction and realization of genome of pathogenic viruses in bacteria in natural conditions. Also, application of this bacteria may be possible for diagnostic or prophylactic range.
CONSORTIUM OF ENZYME-PRODUCING ACTINOMYCETES FROM FRUIT ORCHARD OF MUZAFFARPUR: MOLECULAR IDENTIFICATION BY 16S rRNA GENE SEQUENCE

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Background
Sucrose gradient method and International Streptomyces Project (ISP) media were adopted to isolate actinomycetes in order to constitute a consortium of actinomycetes from the fruit orchard of Muzaffarpur, which were capable to produce important industrial enzymes. Microscopic, cultural and biochemical tests, followed by features-comparison as per the Bergey's Manual of Systematic Bacteriology confirmed a total of 52 strains in placing their systematic affiliation in the present investigation. The 16S rRNA gene sequencing method confirmed these species belonging to Streptomyces, Nocardia, Nocardiopsis and Actinomyces. Based on the consensus nucleotide sequences of ten strains, SK 701 to SK 710, phylogenetic dendrograms were constructed by using neighbour-joining algorithm. Two of these strains, SK 702 and SK 705 were successfully screened to have high phytase activities (380 and 315 U ml⁻¹, respectively). The optimum extracellular enzyme production was noticed at 35 °C on 4th day of culture, when the bacterium entered into late exponential or stationary phase. Though the initial pH of media was 7, however, the phytase production was stable within 3.5 to 4.5 pH.

Objectives
The objective is to identify the actinomycetes from fruit orchard because of their capability to produce industrial enzymes and other beneficial compounds at optimum conditions.

Methods
ISP and PSM were used to screen phytate-degrading actinomycetes, which were finally phylogenetically identified by 16S rRNA gene sequencing.

Conclusions
Species of Streptomyces and Nocardiopsis with phytase producing ability were identified in Litchi orchard of Muzaffarpur for which PCR amplification of 643 base pairs of DNA was achieved.
PHYLOGENETIC HETEROGENEITY OF TUMORIGENIC BACTERIAL STRAINS RECOVERED FROM RASPBERRY IN SERBIA

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Background
Tumorigenic bacteria belonging to the genera Agrobacterium and Rhizobium are responsible for crown gall disease of various fruit species. During the last few years, high incidence of crown gall has been recorded in some young raspberry plantations throughout Serbia, causing considerable economic losses.

Objectives
The objective was to study phylogenetic structure and diversity of tumorigenic strains occurring on raspberry in Serbia.

Methods
A total of 14 strains were isolated from tumor tissue of diseased plants collected in six localities during 2011-2013. Tumorigenicity of strains was determined by PCR using primers specific for tumor-inducing (Ti) plasmid genes and pathogenicity assay on several test plants. PCR targeting the 23S rRNA gene, and 16S rRNA and recA gene sequence analysis were used to identify phylogenetic placement of strains studied. Genetic diversity of isolated strains was evaluated by using ERIC-PCR method.

Conclusions
Tumorigenic bacteria associated with crown gall of raspberry showed extensive genetic diversity. Out of 14 strains, 12 were identified as tumorigenic Rhizobium rhizogenes, one belonged to Agrobacterium tumefaciens genomic species G8, while the remaining strain formed separate phylogenetic lineage within A. tumefaciens species complex, different from all known genomic species. The fact that some geographically separated R. rhizogenes strains exhibited similar ERIC-PCR profiles, may indicate that they have a common origin and were probably disseminated by movement of infected plant material. This study was funded by the national III46008 and EU AREA No 316004 projects.
UNDERSTANDING THE ROLE OF PII PROTEIN IN ARTHROSPIRA PLATENSIS PCC 7345
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Background

PII proteins are known to play a crucial role in the regulation of cyanobacterial N metabolism, but its role and interacting partners in Arthrospira (Spirulina) platensis, a popular neutraceutical, nitrate-utilizing cyanobacterium, have not been characterized.

Objectives

The main aim of this study was to study the regulation of glnB (PII) gene by nitrate and its downstream metabolites and to screen for PII interacting proteins in A. platensis

Methods

We have cloned full length gene (glnB) in pET28a vector, expressed and purified PII protein by affinity chromatography for in vitro interaction studies. A. platensis cDNA yeast two hybrid library was constructed to screen for interacting partners of the PII.

Conclusions

Our studies have revealed transcriptional regulation of glnB (PII) by nitrate and its downstream metabolites. We have identified 28 colonies after library screening as potential interacting partners. Their characterization and confirmation by other approaches is underway.
PHYLOGENOMIC COMPARISON OF 16 O5 SHIGATOXIGENIC ESCHERICHIA COLI (STEC) ISOLATED FROM

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Background

In developed countries, \textit{Escherichia coli} producing Shiga toxins (STEC) cause enteritis and (bloody) diarrhoea in young calves and in humans, the haemolytic–uremic syndrome (HUS) in humans, and are also present in the intestines of healthy cattle. Besides the most frequent serotypes, STEC can belong to several subdominant serotypes, like O5:H- that is present in young calves and, though more rarely, in humans and healthy cattle.

Objectives

The aim of this study is to analyse, at genome level, the phylogeny of 12 calf (isolated at FVM-ULg and ARSIA) and 4 human (isolated at UZ-VUB) O5 STEC belonging to different pathotypes; the eae\textsuperscript{+}stx\textsuperscript{1+} (14 isolates), eae\textsuperscript{+}stx\textsuperscript{1+}stx\textsuperscript{2+} (1 isolate) or stx\textsuperscript{1+} (1 isolate) pathotypes.

Methods

Genomic DNA was extracted from a growing colony and sequenced on the Illumina MiSeq platform. The raw sequences were assembled into scaffolds using an in-house pipeline and subjected to an automated annotation pipeline. The contig data were used to determine the MLST and the pathotypes. A scheme of 20 housekeeping and virulence genes were concatenated and used to select the representative strain, which will be submitted to another run of sequencing and used as the reference for a phylogenomic analysis. The phylogenomic tree will allow to analysing whether the O5 STEC strains could be clustered according to the origin parameter (human or bovine).
Conclusions

Genomic sequencing has become a quick and efficient tool that can be used to replace numerous existing typing analysis methods by a unique analysis scheme.
BACTERIOPHAGE-MEDIATED, HIGH EFFICIENCY HORIZONTAL GENE TRANSFER IN BACTERIAL PATHOGENS AND BIOCONTROL AGENTS

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Background
Transduction is one of the main drivers of horizontal gene transfer in bacteria and contributes to the rapid evolution of prokaryotes. No transducing phages have yet been described for Serratia plymuthica, a biocontrol bacterium that produces bioactive molecules - including the antifungal macrolides, oocydins (1).

Objectives
The aim of this work was to isolate a generalized transducing phage(s) for Serratia plymuthica, and related strains.

Methods
New phages were isolated, scored for transduction capacity, phenotypically and
morphologically characterised, and genomically sequenced.

**Conclusions**

The new *Serratia* phage, ϕMAM1 (2,3), transduced chromosomal markers at high frequencies. We demonstrated transduction of the 77-kbp oocystin gene cluster and showed heterogenic transduction of plasmids. Morphological, genomic and phylogenetic analyses grouped ϕMAM1 within the suggested new genus, Viunalikevirus (3). We show that four additional viunalikeviruses, infecting plant and human pathogens, were highly efficient generalized transducing phages. We hypothesised that related phages are likely to perform efficient horizontal gene transfer. To test this hypothesis, we first isolated several new viunalikeviruses from the environment, ϕXF1, ϕXF3 and ϕXF4. All these phages transduced chromosomal markers and plasmids at high frequencies (4). We predict that all viunalikeviruses will perform efficient horizontal gene transfer in their respective hosts. These phages will have utility for functional genomics, bacterial engineering and synthetic biology studies, but may be inappropriate candidates for use in phage therapy.

**References:**

THE COMPLETE GENOME OF MYCOPLASMA SYNOVIAE WVU1853

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Background

*Mycoplasma synoviae* causes infectious synovitis, respiratory tract infections, and eggshell apex abnormalities in commercial poultry. Sequencing of the field isolate MS53 has provided a basis for research exploring the virulence and evolution of this organism; however, this isolate is attenuated and is not available to laboratories worldwide. Explorations into loci of interest reveal high levels of genetic diversity between MS53 and other isolates, emphasizing the need to solve the genomes of additional strains. Past efforts have been complicated by a large repeat-rich region encoding variable surface antigens (the *vlhA* locus).

Objectives

The objective of the work was to generate a complete genome sequence for the *M. synoviae* type strain WVU1853 and compare its features to MS53.

Methods

Sequence read were generated using a combination of Illumina MiSeq and Pacific Biosciences SMRT technologies. Genome assembly was carried out using CABOG (J. Craig Venter Institute), Newbler (454 Life Sciences), and Ray (Boisvert et al., 2012) assembly software programs.

Conclusions

The *M. synoviae* WVU1853 genome was sequenced and closed. It is larger than the MS53 genome, and many notable changes in the *vlhA* locus were apparent.
NIC-GENE CLUSTER’S MODULAR DESIGN WITHIN THE ARTHROBACTER GENUS

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Background

The pAO1 megaplasmid of *Arthrobacter nicotinovoras* shows low levels of sequence similarity with the *Arthrobacter* plasmids, but shares most of its *nic*-genes with three *Arthrobacter* draft genomes: M2012083 (GI:NZ_AKKK00000000), SJCon (GI: NZ_AOFD00000000) and AK-YN10 (GI: NZ_AVPD00000000).

Objectives

The objective of the current study is to make an evolutionary analysis of the *nic*-cluster taking into account the arrangement and collinearity of the *nic* ORF’s.

Methods

The three draft genomes of the above mentioned *Arthrobacter* strains were assembled based on the existing final *Arthrobacter* genomes using MAUVE, annotated with RAST and further aligned with Artemis. *Arthrobacter* sp. AK-YN10 (a gift from Dr. Atya Kapley, CSIR-NEERI, Nagpur, India) and pAO1 strains were grown on citrate medium supplemented with nicotine. Nicotine consumption in the medium was followed by HPLC.

Conclusions

The *nic*-gene cluster can be divided into five modules, each module encoding a precise step in the nicotine-pathway. For each module, a general rule can be observed: the pAO1 modules are the most complex, with a large number of genes, including transposases and insertion elements. The SJCon modules are the most simple, with a small number of ORF’s and large non-coding regions. The AK-YN10 strain is somewhere in the middle, but the five modules are spread through the genome. The test for the nicotine resistance of this bacteria has shown levels comparable with the pAO1 strain. Moreover AK-YN10 can grow on nicotine containing citrate medium without forming the characteristic nicotine-blue pigment.

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Background

*Delftia* sp. JD2 is an auxin/siderophore plant-growth-promoting bacterium that promotes growth and nodulation of alfalfa when co-inoculated with *Sinorhizobium meliloti*. Additionally, JD2 reduces Cr(VI) to Cr(III) and shows multi-resistance to several heavy-metals and antibiotics.

Objectives

The aims of this work were: 1) to perform a comparative genomic analysis using the draft genome of JD2 and, 2) to analyze the molecular elements involved in JD2 phenotype, such as chromium resistance, auxin- and siderophore-production, among others.

Methods

We had recently sequenced the genome of JD2 and its automated annotation has been finished. We are currently analyzing the potential pathways involved in chromate resistance, auxin production, polyhydroxy-alkanoates production and accumulation, horizontal gene transfer elements and others. Results from comparative genomic analysis and different approaches to inferring phylogeny will be shown during the meeting.

Conclusions

JD2 has a 6.76Mb genome. The annotation and comparative genomic analysis revealed some functional characteristic associated with its previously reported plant growth promoting effects on alfalfa. JD2 has numerous genes related to tryptophan synthesis and degradation, indol 3-acetic acid biosynthesis, siderophores production and secretion and TonB dependent transporters with various substrate specificities. Consistent with its environmental adaptability, JD2 contains a large number of genes involved in the transport and efflux of toxic compounds. Syntenic genes related to
chromate resistance \((chr)\) were identified. Interestingly, in adjacent positions to \(chr\) cluster some mobile elements (trasposases and integrases) were found. The results provide insights into the potential use of JD2 in heavy metal bio-remediation programs and biofertilization technologies.
SPECIES REASSIGNMENT OF ENTEROBACTER CLOACAE COMPLEX ISOLATES BY IN SILICO PHYLOGENETIC ANALYSIS

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Background

The taxonomic resolution of closely related species belonging to Enterobacter cloacae complex ("EcC", E. cloacae, E. hormaechei, E. asburiae, E. kobei and E. ludwigi) by conventional methodologies or MALDI-TOF MS is often unreliable, and sequencing of 341bp hsp60 has been proposed for their discrimination. Moreover, a multilocus sequence typing (MLST) assay targeting dnaA, fusA, gyrB, leuS, pyrG, rplB and rpoB genes has been suggested for epidemiological typing of E. cloacae, considered one of the most frequent species in the clinical setting.

Objectives

To assess by a phylogenetic approach, using different biomarkers, the distribution of EcC species in available databases.

Methods

A phylogenetic-based study was conducted to analyze the hsp60 (complete gene) and individual and concatenated sequences of dnaA, leuS and rpoB (partial genes) from 352 Enterobacter sp. isolates (including type and/or reference EcC strains), deposited in the Pathosystems Resource Integration Center (Patric) and MLST databases. Phylogenetic trees were constructed using Neighbour-Joining method supported by Tamura 3-parameter model and the reliability of internal branches assessed by bootstrap from 1000 re-samplings.

Conclusions

The topology of the trees (individual and concatenated) congruently revealed 5 main clusters corresponding to the five known EcC species. Interestingly, E. cloacae were minority (3%), whereas E. hormaechei (63%) and E. asburiae (15%) were overrepresented, highlighting the current misidentification of EcC species, with potential relevant clinical implications. Finally, cut-off values of ≤91.7% and ≤95.9% are proposed, respectively, for leuS and hsp60, the most discriminatory genes for EcC species determination, which could support their reliable identification.
GENETIC ANALYSIS OF THE LISTERIA PATHOGENICITY ISLAND 1 OF L. MONOCYTOGENES 1/2A AND 4B ISOLATES

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Background
Genetic analysis of the Listeria Pathogenicity Island 1 (LIPI-1) as well as several other genes has been applied in order to gain insights into the evolution of L. monocytogenes.

Objectives
The aim of the present study was to apply descriptive, phylogenetic, recombination and selection analyses on alignments of the LIPI-1 of 1/2a and 4b isolates of different origin in order to facilitate understanding of the evolution of this virulence gene cluster.

Methods
A total of 19 L. monocytogenes isolates (9 meat isolates, serotype 1/2a; 5 meat isolates, serotype 4b; 5 strawberry isolates, serotype 4b) were subjected to sequencing of their LIPI-1. Alignments of the complete island as well as intragenic and intergenic regions were used for descriptive, phylogenetic, recombination and selection analyses.

Conclusions
Descriptive analysis revealed extensive nucleotide diversity mostly in the intragenic regions. The actA gene of 1/2a and 4b meat isolates and the hly gene of the 4b strawberry isolates exhibited the higher diversity. On the contrary, limited diversity was observed in prfA and plcA genes of the 4b isolates and mpl gene of the 1/2a isolates. Phylogenetic analysis of the complete island resulted in two major clusters that were consistent with serotype assignment of the isolates. Moreover, within the 4b cluster, isolates were separated according to their isolation source. Selection analysis revealed that the island consisted of randomly evolving DNA with the exception of prfA gene of 1/2a isolates and actA gene of 4b meat isolates for which purifying selection was indicated. Finally, no statistically significant evidence for recombination has been observed.
EFFECT OF LEMONGRASS ESSENTIAL OIL ON LISTERIA MONOCYTOGENES GENE EXPRESSION
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Background
The transcriptomic response of L. monocytogenes to abiotic stresses has been the epicenter of intensive study over the last decade and the complexity of regulation of gene expression has been highlighted.

Objectives
The aim of the present study was to assess the transcriptomic response of L. monocytogenes isolates on the exposure to lemongrass essential oil.

Methods
Overnight culture of six strains previously isolated from a strawberry sample was spread over the surface of BHI agar and exposed to 5 and 10 μL of lemongrass essential oil that were applied on a Whatman paper placed on the lid. Three of them survived in the former case and one in the latter; in all cases biomass was collected and RNA was isolated. The expression of the key virulence genes sigB, plcA, plcB, hlyA, inlA, inlB, inlC, inlJ, lmo2470 and lmo2672, as well as accA, acpP and fapR involved in fatty acid biosynthesis/metabolism and murE and pbpB involved in peptidoglycan biosynthesis was assessed by RT-qPCR.

Conclusions
Downregulation of virulence genes was observed with the exception of sigB for which upregulation in two isolates was observed. On the contrary, significant differences between the isolates were observed regarding the expression of accA, acpP, fapR, murE and pbpB. The increased amount of lemongrass essential oil affected significantly the expression of accA, pbpB, sigB, hlyA, inlA, inlB, inlJ and lmo2672.
AN HOMOLOG OF THE FRZ PHOSPHOENOLPYRUVATE: CARBOHYDRATE PHOSPHOTRANSFERASE SYSTEM OF EXTRAINTESTINAL PATHOGENIC ESCHERICHIA COLI IS ENCODED ON A GENOMIC ISLAND IN SPECIFIC LINEAGES OF STREPTOCOCCUS AGALACTIAE

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Background

Streptococcus agalactiae and Escherichia coli have in common the capability to colonize and infect similar hosts, and to survive in the environment and in foods. In E. coli, the Frz system is involved in environmental sensing and regulation of the expression of adaptation and virulence genes. Thus, it is possible that S. agalactiae possesses a system similar to Frz to sense its environment and to adapt to environmental conditions.

Objectives

The presence of a system similar to Frz was searched in S. agalactiae strains representative of the genetic diversity of the species.

Methods

The genomes of 492 strains of S. agalactiae of human and animal origins were analyzed in silico and by PCR. The strains were characterized by multilocus sequence typing.

Conclusions

We identified an operon (named fru2) coding a system similar to Frz. fru2 encodes three subunits of a PTS transporter of the fructose-mannitol family, a transcriptional activator of PTSs of the MtlR family, an allulose-6 phosphate-3-epimerase, a transaldolase and a transketolase. fru2 is present in a genomic island. The showed that the presence of fru2 is closely linked to the phylogeny of S. agalactiae. Our results indicate that fru2 was acquired during the evolution of S. agalactiae before the divergence of CCs 1, 7, 10 and 283, and STs 130 and 288. As S. agalactiae strains of CCs 1 and 10 are frequently isolated from adults with invasive disease, we hypothesize that the S. agalactiae fru2 system senses the environment to allow the bacterium to adapt to new conditions encountered during the infection of adults.
COEVOLUTION OF HUMAN AND HELICOBACTER PYLORI GENOMES SHAPES THE RISK OF GASTRIC DISEASE

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Background
Helicobacter pylori is the principal cause of gastric cancer (GC). However, H. pylori prevalence generally does not predict cancer incidence.

Objectives
To determine if the coevolution between host and pathogen influences the risk of GC in two Colombian populations: Tumaco, a coastal population with low risk of GC, and Tuquerres, a mountain population with high risk of GC. Our hypothesis postulates an association between the severity of gastric lesions and the patterns of genomic variation in matched human and H. pylori samples.

Methods
292 patients were recruited from the two populations. Histological diagnostics were based on the Sydney classification. 252 strains of H. pylori were included for MLST analysis. Human DNA was genotyped using the Immunochip to characterize the human ancestry. The Admixture model of STRUCTURE assigns proportions of ancestry to each individual sample.

Conclusions
H. pylori isolates with an ancestral African cluster predominating in a low-risk (coastal population), and a European cluster in a high-risk (mountain population). The human ancestries of biopsied individuals also vary geographically, with mostly African ancestry in the coastal region, and mostly Amerindian ancestry in the mountain region. The interaction between host and pathogen ancestries completely accounted for the difference in the severity of gastric lesions in the two regions of Colombia. H. pylori ancestry was relatively benign in humans of African ancestry but was deleterious in individuals with Amerindian ancestry. Thus, coevolution probably shapes disease risk, and the disruption of coevolved human and H. pylori genomes can explain the high incidence of gastric disease in mountain population.
EXPLORING GENETIC DIVERSITY OF XANTHOMONAS ARBORICOLA PV. CORYLINA STRAINS AND DEVELOPMENT OF SPECIFIC MOLECULAR MARKERS
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Background: Xanthomonas arboricola pv. corylina (Xac) is the causal agent of bacterial blight of hazelnut. The most efficient distribution of the pathogen to new geographical areas occurs through contaminated plant material. Therefore, fast and reliable pathogen detection using highly specific molecular diagnostic tools is essential for the production of disease-free planting material.

Objectives: The aim of this work was to study genetic structure and diversity of Xac strains and develop molecular tools for specific and rapid detection of the pathogen.

Methods: In order to study genetic diversity forty Xac strains originating from Serbia were initially analyzed by rep-PCR using BOX, ERIC and REP primers. Obtained genetic profiles from all three PCR methods were transformed into a binary matrix, combined and used for phylogenetic analysis and construction of the dendrogram. Seventeen representative strains with different rep-PCR patterns were selected for further macrorestriction analysis using SpeI restriction enzyme and pulsed-field gel electrophoresis (PFGE). Specific primers were designed based on specific DNA fragment obtained by ERIC-PCR.

Conclusions: The strains were differentiated into eight clonal and highly related genetic groups by rep-PCR analysis. Unique PFGE patterns were obtained for each representative strain indicating more polymorphism between the strains, compared to rep-PCR. Designed PCR set of primers enabled detection of Xac strains from different geographic origin. However, they could not distinguish Xac from closely related Xanthomonas spp.
EXPLORING DIVERSITY AND EVOLUTION OF MICROBIAL GENOMES USING NEXT-GENERATION SEQUENCING

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Background

Genome organization and sequence is dynamic, undergoing changes as part of evolutionary process based on genomic structure and environmental conditions under which organisms grow. The whole genome sequences (WGS) are useful information to understand evolutionary aspects of any organism. We have used Next-generation sequencing (NGS) to survey and analyse genomic changes occurring in different species and isolates of bacteria.

Objectives

1. Identification and analysis of mutation hotspots in genome of $M. tuberculosis$.
2. Construction of phylogenetic trees from NGS datasets of microbes.
3. Identification of IS-elements from NGS datasets and understanding role of IS6110 in genome evolution of $M. tuberculosis$.

Methods

1. We have used Shewhart Control Chart to identify genomic mutation hotspots in $M. tuberculosis$.
2. We have developed a method which uses random sampling of genome to reduce computational complexity to construct phylogenetic tree. It is able to construct trees in absence of reference genome, which is important for non-model organisms. This was used for NGS datasets of $M. tuberculosis$, $E. coli$ and $V. cholerae$.
3. We have used an split-read approach to identify location of IS elements in microbial genome from NGS datasets. Distribution and copy number of IS6110 was identified from 1377 $M. tuberculosis$ strains representing all major global lineages.

Conclusions

Identification and analysis of hotspots enables to study regions responsible for diversification and phenotypic consequences such as, drug-resistance. Generation of phylogenetic trees directly from NGS reads enables study of evolutionary relationships among hundreds of closely related strains. Lineage specific IS6110 distributions were observed and gene truncation, possible promoter activity and IS mediated genomic rearrangements were studied in detail.
A POSSIBLE EVOLUTIONARY SCENARIO FOR THE LOSS OF OPGGH IN YERSINIA PESTIS.

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Background

The opgGH operon encodes the glycosyltransferases OpgH and OpgG, which synthesize osmoregulated periplasmic glucans (OPGs) and are required for motility, biofilm formation and virulence in various bacteria. OpgH also sequesters FtsZ, in order to control bacterial size as a function of nutrient availability. The role of opgGH in Yersinia pseudotuberculosis is unknown. Furthermore, opgGH was lost during the emergence of Yersinia pestis (the agent of plague) from Y. pseudotuberculosis (an enteropathogen). This loss is surprising, since Y. pestis is extremely virulent and flea-borne transmission of Y. pestis relies on biofilm formation.

Objectives

To determine the role of opgGH in Y. pseudotuberculosis and consider the role of opgGH’s loss in the emergence of plague.

Methods

Genetic approaches were used to generate (i) a ΔopgGH Y. pseudotuberculosis strain, and (ii) Y. pestis, ΔopgGH Escherichia coli and ΔopgGH Dickeya dadantii strains expressing Y. pseudotuberculosis opgGH. Biochemical assays (used to detect OPG synthesis), motility and biofilm assays, microscopy (used to measure bacterial size) and virulence testing (using endive leaves, mice and fleas) were used to determine the impacts of the mutation and the opgGH expression in the strains of interest. Animal studies were performed in compliance with the French regulations.

Conclusions

Our results suggest an evolutionary scenario in which an ancestral strain of Yersinia lost or did not acquire a cofactor that was important for OPG biosynthesis but kept opgGH to control its bacterial size in a nutrient-dependent manner. The opgGH gene was then lost presumably because the nutrient-dependent size-control mechanism was no longer necessary.
Background
The genus *Treponema* comprises several uncultivable human pathogens including *Treponema pallidum* subsp. *pallidum* (TPA), the causative agent of syphilis, *T. p.* subsp. *pertenue* (TPE, the causative agent of yaws), and *T. p.* subsp. *endemicum* (TEN, the causative agent of bejel).

Objectives
The goal of this study was to identify positively selected genes in comparisons within and between TPA, TPE and TEN strains.

Methods
Altogether, nine whole genome sequences comprising 4 TPA, 4 TPE and 1 TEN strains were used to identify positively selected genes. Several software packages were used including Mega4, PAML and TimeZone in all possible pairwise combinations.

Conclusions
A set of 25 treponemal genes were identified among more than 1000 tested TPA, TPE and TEN orthologous genes. Eleven of these genes were found to be positively selected in TPA-TPE comparisons but not within TPA or TPE strains. In contrast, five genes including *bamA* and *mcp-2* were positively selected also within TPA and TPE strains. Two genes (TP0559, TP0966) appear to be positively selected within TPA but not within TPE strains suggesting their role in syphilis pathogenesis. Gene TP0304 is positively selected within TPE but not within TPA. Two genes (*tprF,I*) were found positively selected within TPA and TPE strains but not in comparison between TPA and TPE strains. Except for *tprC* gene where positive selection was identified between TPA and TPE comparisons, TEN orthologs resembled TPE orthologs. Altogether, positively selected genes encode for Tpr proteins, protein antigens (e.g. TP0326, BamA), putative lipoproteins and outer membrane proteins.
Background
The saprophyte L. monocytogenes is well established in the environment.

Objectives
Persistent L. monocytogenes and Listeria spp. isolates (characterized by PFGE and MLST typing) are hypothesized to be better adapted to the food processing environment (FPE).

Methods
The whole genome of 15 L. monocytogenes and Listeria spp. genotypes established in the same food processing environment was sequenced applying the Illumina sequencing technique.

Conclusions
The L. monocytogenes core-genome was highly syntenic. Differences were detected in mobile elements of the accessory genome. Eight L. monocytogenes and two L. innocua strains indicated the presence of 15-90 kbp plasmids. The long-term L. monocytogenes persistent strains, assigned to genetic lineage I [sequence type (ST) 5] and lineage II (ST204), harbored a plasmid comparable to plm80. When comparing L. monocytogenes ST37 at the beginning and the end of isolation, the loss of a distinct phage region (~33 kbp) was noticeable. The β-lactamase encoding transposon Tn552, was detected in the plasmids of L. monocytogenes ST204 (n=1) and ST5 strains (n=6). The transposon Tn554, suspected for benzalkonium resistance, was found in L. monocytogenes ST9, one L. innocua, and L. seeligeri. The L. monocytogenes stress-survival islet (SSI-1) was present in ST5, ST204, and ST9.

The results indicate mutations in the L. monocytogenes ST5, ST37 phage and/or plasmid genome suspected for better adaption and survival in the particular FPE. Gene regions for better adaption to disinfectants and acidic stress (Tn554, SSI-1) were found in long term persistent L. monocytogenes strains.
AN IMPROVED CLASSIFICATION OF DYP-TYPE PEROXIDASE FAMILY ACCELERATES THE RESEARCH OF THE FAMILY

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Background
So far, the range of DyP-type peroxidase family was obscure. Therefore, the name of DyP-type peroxidase family and its superfamily tended to be confused.

Objectives
To solve this problem, we focus on the relationship between tertiary and primary structures of DyP-type peroxidases associated with their function. From the viewpoint, we propose the range of DyP-type peroxidase family reasonably.

Methods
Dye-decolorizing peroxidase from the basidiomycete Bjerkandera adusta Dec 1 (DyP) is a heme peroxidase. This name reflects its ability to degrade several anthraquinone dyes. The substrate specificity, the amino acid sequence, and the tertiary structure of DyP are different from those of the other heme peroxidase families. Therefore, many proteins showing the similar amino acid sequences to that of DyP are called DyP-type peroxidase which is a new family of heme peroxidase identified in 2007. Although all structures of this family show a similar structure fold, this family includes many proteins whose amino acid sequence identity to DyP is lower than 15%. Moreover, their catalytic efficiency ($k_{cat}/K_m$) is a few orders of magnitude less than that of DyP. For the activity, some of them have been reported different from peroxidase activity (dechelatase activity).

Conclusions
Here, we report the differences and similarities of structure and function among this family and propose the reasonable improved classification of DyP-type peroxidase family, that is, class P, I and V. This classification is rather sophisticated and accelerates the logical study for DyP-type peroxidase family.
APPLICATION OF GENOMICS TO PROKARYOTIC SYSTEMATICS: THE GENUS RHODOCOCCUS AS A CASE STUDY

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Background

Prokaryotic systematics provides the framework for all research in microbiology. Just as 16S rRNA gene sequence analysis revolutionised prokaryotic systematics a generation ago, genome sequence analyses now have the power to be a similarly disruptive technology. Case studies are needed to highlight the power of the new methodological approaches.

Objectives

To apply whole genome sequence analyses to reappraise the taxonomy of genus Rhodococcus.

Methods

We have determined genome sequences for 15 strains belonging to the genus Rhodococcus. Using this data and publicly available sequences, we have analysed genome sequences of 83 strains of 20 species within the genus Rhodococcus using phylogenomic, genetic and functional distance-based approaches.

Conclusions

Based on our analyses the rhodococci were resolved into six robust lineages that were as distant from each other as they were from other genera including representatives of Nocardia and Gordonia. These analyses also revealed a widespread problem of strain misidentification using classical approaches. The lineage containing Rhodococcus rhodochrous, the type species of the genus, represents the genus Rhodococcus sensu stricto. However, the others can potentially be designated as novel genera. Rhodococcus equi formed a distinct lineage with Rhodococcus defluvii consistent with our recent proposal that the former should be reclassified as “Prescottella equi”. These data provide a case study demonstrating that genome sequencing is a cost effective and powerful tool for resolving complex taxonomic questions.
THE PLASMID COMPLEMENT OF LACTOCOCCUS LACTIS NCDO712

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Background

The lactic acid bacterium Lactococcus lactis NCDO712 was isolated from a dairy starter culture in the 1950s and it is the ancestor of the prophage and plasmid cured model strain MG1363. L. lactis NCDO712 was described to contain 5 plasmids (~50 kb, 13.6 kb, 8 kb, 3.8 kb, 2.7 kb) of which so far only two have been sequenced.

Objectives

The characterization of \textit{L. lactis} NCDO172 will help establishing it as a relevant model strain because of its functional proximity to commercially used cultures. In combination with the genetically highly accessible derivative MG1363, it will form a useful couple to study industrially relevant traits of \textit{L. lactis}.

Methods

Total DNA of \textit{L. lactis} NCDO712 was sequenced using Illumina and PacBio. Additional PCR reactions and Southern hybridizations were done to complete the assembly. This resulted in 5 circular plasmids and one linear plasmid contig and allowed a SNP comparison with MG1363.

Conclusions

We here sequenced the complete plasmid complement of strain NCDO712, which resulted in the identification of 6 plasmids (55.4kb, 8.6kb, 3.6kb, 2.1kb, 15.5kb, 51.7kb) - one more than initially described. Next to an extracellular protease gene and the genes necessary for lactose utilization, which are on the already sequenced plasmid pLP712, the other plasmid-encoded genes include those of a variety of IS-located transposases, two restriction modification systems (specificity subunits) and stress related genes. The sequencing of the total DNA of strain NCDO712 also allowed identifying of some chromosomical differences between the strains NCDO712 and MG1363.

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Background

Acremonium chrysogenum is the industrial fungal producer of the β-lactam antibiotic cephalosporin C, which is an important drug due to its broad activity against Gram-positive and Gram-negative bacteria. The source material for the modern chemically modified cephalosporin C derivatives is still mainly produced by A. chrysogenum, which marks its importance for biotechnology and medicine.

Objectives

Therefore it is of high interest to gain a deeper insight into the biosynthetic processes involved in cephalosporin C production and their regulation. Already established molecular tools like homologous recombination and the FLP/FRT marker recycling system are essential for the work with this fungus. A further crucial step for effective scientific research with A. chrysogenum is to disclose its genomic sequence.

Methods

DNA sequencing, sequence assembly, annotation

Conclusions

After quality trimming and assembly we were able to obtain a genome sequence with an overall length of 28.6 Mb distributed on 542 scaffolds. Furthermore 8901 protein coding genes were annotated using the protein data from closely related species and the UniProtKB/Swiss-Prot database. The potential to produce a variety of secondary metabolites is underlined by 42 putative secondary metabolite cluster found in the genome sequence. The available genomic sequence and annotation of A. chrysogenum represents a versatile platform for future functional genomic approaches (1).
IN SILICO STRATEGIES FOR THE IDENTIFICATION AND CLASSIFICATION OF BACTERIAL AUTOTRANSPORTER PROTEINS

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Background

Autotransporters are the representatives of the type V secretion system proteins that are assembled into the outer membrane of bacterial cells, and are the most prevalent of the seven types of secretion systems in gram-negative bacteria.

Objectives

The huge data from genome sequencing studies provides a resource of autotransporter proteins as well as pose a challenge in the identification and classification of the autotransporter sequences. We interrogate the bioinformatics strategies to address questions that increase our understanding of sequence-structure-function relationships in autotransporter proteins.

Methods

By using the properties of the whole sequence we propose a new classification system for autotransporter proteins has been proposed. A Support Vector Machine model was subsequently constructed for the identification and classification of new FAEs into the pre-assigned clusters.

Conclusions

This study centers on the descriptor-based classification and structural analysis of experimentally verified and putative autotransporter proteins; nevertheless, the framework presented here is applicable to every poorly characterized enzyme family.
TEMPERATURE-INDUCED MUTAGENESIS AND MORTALITY IN CUPRIAVIDUS METALLIDURANS STRAINS
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Background
Cupriavidus metallidurans is a β-proteobacterium which can be found in industrialized and anthropogenic environments. It was previously observed that growth of C. metallidurans CH34 at 37°C in rich media leads to high mortality (TIMM).

Objectives
The aim of this study is: (i) to scrutinize TIMM-inducing media and (ii) to investigate this phenomenon at the genetic and molecular level.

Methods
First, TIMM-induction was analyzed with different growth media. Next, the TIMM-phenotype was studied using a time-lapse microscope. Finally, whole-genome sequence analyses were performed for two derivatives with inherited resistance to all tested TIMM-inducing media.

Conclusions
The individual amino acids phenylalanine and cysteine and high ammonium concentrations could activate TIMM at 37°C in C. metallidurans CH34. Moreover, a mineral medium with low salt concentrations, i.e. Schatz-medium, also induced a high mortality at 37°C. These observations led to the hypothesis that different TIMM-inducing triggers are responsible for the phenotype. First, uptake of specific amino acids can lead to an toxic imbalance. Second, the TIMM-phenotype could be linked with osmotic potential and membrane integrity as characteristics like cell elongation, reproduction halting and cell lysis were induced. Interestingly, counteractivating agents were found as induction was inhibited by increasing the osmotic potential of Schatz-medium or by adding sorbitol to the different TIMM-inducing media. Both observations provided support for the hypothesis that membrane integrity plays a crucial role in the TIMM-phenotype. A common missense mutation was identified in a gene coding for an alcohol dehydrogenase. To examine the effect of this mutation complementation studies are currently being performed.
AN UNUSUAL STRAIN WHICH PROBABLY BE AN ANCESTOR IN THE EVOLUTIONARY PROCESS OF YERSINIA

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Background

We had isolated an unusual Y. enterocolitica strain from Rattus norvegicus, the brown rat. The strain did not utilize urea where API 20E identified Escherichia coli; however it grew well at 25°C where E. coli does not. We deduced the strain was Y. enterocolitica using sequence alignment of the 16S rRNA gene.

Objectives

To analysis the discrepancy with other Y. enterocolitica and comprehend its location in the evolutionary process of Yersinia by sequencing the whole genome of LC20.

Methods

The LC20 genome was sequenced using a whole-genome shotgun sequencing strategy and Illumina Hiseq sequencing technology.

Conclusions

Whole-genome comparison confirmed LC20 was Y. enterocolitica, the whole genome was collinear with Y. enterocolitica 8081. LC20 carried a great number of predicted CDSs and IS elements. Two plasmids shared low genetic homology with pYV from the Yersinia genus, one was an ancestral Yersinia plasmid and the other was novel encoding a number of transposases. The core-based and pan-based phylogenetic trees showed LC20 was classified into Y. enterocolitica cluster but on a distinct branch, which was closer to Y. pestis and Y. pseudotuberculosis. Some pathogenic and non-pathogenic Y. enterocolitica specific genes coexist in LC20. The urease gene did not exist on the genome consistent with API 20. LC20 is sub-classified into Y. enterocolitica based at the level of the whole genome,
though its bacterial identification coding system of biochemical metabolism showed it was *E. coli*. The strain may be a progenitor of *Y. enterocolitica* and represent even an ancestor in the evolutionary process of *Yersinia* genus.
CHARACTERIZATION OF A NOVEL SULFATE-REDUCING BACTERIUM POSSESSING DESULFOVIRIDIN AND VIBROID MORPHOLOGY, BELONGING TO THE FAMILY DESULFOMICROBIACEAE

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Background
The order Desulfovibrionales is organized in four families; Desulfovibrionaceae, Desulfonatronumaceae, Desulfohalobiaceae and Desulfomicrobiaceae. The genus Desulfomicrobium is a sole member of family Desulfomicrobiaceae at the time of writing. The genus have been characterized as rod- or ellipsoidal-shaped morphology and absence of desulfoviridin. In this study, a novel strain Pf12Bᵀ related to Desulfomicrobium species was isolated in pure culture.

Objectives
The aim of our study is characterization of the strain Pf12Bᵀ.

Methods
Strain Pf12Bᵀ was obtained from an enrichment culture established under sulfate-reducing conditions at 45°C with an inoculum of sediment from a brackish meromictic lake. Isolation of the strain was carried out by repeated agar shake dilution method.

Conclusions
Cells were vibroid, and motile. The closest relative of the strain was Desulfomicrobium baculatum with 16S rRNA gene sequence similarity of 91%. Phylogenetic analysis based on the 16S rRNA gene sequence revealed that the strain Pf12Bᵀ belonged to the family Desulfomicrobiaceae, in the order Desulfovibrionales, Deltaproteobacteria. This result was also supported by the phylogenetic analysis based on DsrA amino acid sequence. Sulfate, sulfite, thiosulfate were used for electron acceptors. Formate and fumarate were used for substrate in the presence of sulfate. Yeast extract enhanced growth of the strain. The isolate grew at temperatures between 13-50°C with an optimum temperature range of 42-45°C. The strain was desulfoviridin positive. Major cellular fatty acids of the isolate were C₁₆:₀ and C₁₈:₀. It was suggested that the strain Pf12Bᵀ is a representative of novel species of novel genus within the family Desulfomicrobiaceae.
CRYSTAL STRUCTURE OF ANABAENA SP. DYP-TYPE PEROXIDASE GIVES AN INSIGHT INTO SUBCLASSIFICATION OF DYP-TYPE PEROXIDASE FAMILY

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Background
Dye-decolorizing peroxidase (DyP)-type peroxidase is a new family of heme peroxidase, distributing from bacteria to fungi. This name derives from the degradation ability of anthraquinone dyes. DyP-type peroxidases are known to show the similar structure. On the other hand, this family includes many proteins which show very low amino acid sequence similarity, resulting in inaccurate sequence alignment. This family has been subdivided into four classes (A, B, C, and D) based on the amino acid sequences. Recently, we reassessed the sequence alignment of DyP-type peroxidases using structure-based sequence alignment. The generated alignment reflected the differences hidden in the similar structures. Based on this result, we proposed the new subclass into three classes (P, I, and V). Here, classes P, I, and V mean former classes B, A, and C with D, respectively. The structures of proteins of former classes C and D are almost same.

Objectives
Anabaena sp. DyP-type peroxidase (AnaPX) has been classified into different class (C or D) by different researchers. Therefore, it is expected that structure-based sequence alignment including the structure of AnaPX gives an insight into the subclass, especially, class V.

Methods
We determined the structure of AnaPX by X-ray crystallography and generated phylogenetic tree using MATRAS program (T. Kawabata, Nucleic Acids Res. 31, 3367–3369 (2003)) as multiple three-dimensional alignment tool.

Conclusions
AnaPX showed the almost same structure as another structures belonging to class V and belonged to class V branch. This result supports our proposition that the border between former classes C and D is vague.
GENOMIC EPIDEMIOLOGY OF MULTIPLE ACINETOBACTER BAUMANNII OUTBREAKS IN A VETERINARY INTENSIVE CARE UNIT

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Background
Acinetobacter baumannii is an emerging pathogen primarily found in health-care associated settings. Due to its potential to acquire and control antimicrobial resistance and its ability to survive for extended periods in the environment, outbreaks are a threat and challenge for health care. Molecular epidemiology of A. baumannii is difficult because typing methods often lack resolution to identify transmission chains. Whole genome sequencing (WGS) forms an alternative method to investigate whether repeated outbreaks are independent events or are linked.

Objectives
Investigation into the relatedness of two outbreaks of A. baumannii in the companion animal intensive care unit (caICU) at Utrecht University in 2012 and 2014.

Methods
Multi Locus Sequence Typing (MLST) was performed for both outbreaks. All outbreak strains, three European Control strains (EC-I, -II and -III) and four environmental strains were sequenced on an Illumina Miseq platform. Genomes were assembled using SPAdes, horizontal gene transfer was detected using ClonalFrame, time-measured phylogeny was reconstructed using BEAST.

Conclusions
MLST of the isolates revealed that all animal isolates from both outbreaks in the caICU belong to ST2. Isolates from the outbreaks had similar resistance phenotypes, suggesting that the events are linked. Time-measured phylogeny on WGS data of the clinical isolates, reference isolates and public genome sequences revealed however, that the outbreaks are two independent events and that isolates from both outbreaks diverged at least 20 years before. Our study shows the benefit of WGS in outbreak management in a clinical setting for pathogens for which traditional molecular typing methods lack resolution.
IDENTIFICATION AND CHARACTERIZATION OF PATHOGENIC TYPES OF ESCHERICHIA COLI ISOLATES FROM A COHORT OF 1 TO 24 MONTH-OLD CHILDREN

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Background
Colonization by Enterobacterias takes place in the first hours of life. This event is influenced not only by the surrounding environment, but also by bacterial flora of those in contact with the newborn, and the host conditions. The kind of bacterial flora acquired in this stage of life could influences the development of the newborn in a decisive form.

Objectives
To study Enterobacterial colonization process in a cohort of newborns through the identification of the presence of Escherichia coli enteropathogenic strains.

Methods
Coproscopic cultures of a cohort of newborns were carried out for a period of 24 months where bacterial genus and species were identified through conventional techniques. Enteropathogenic E. coli strains were re-identified using Enterotest and were later characterized by Multiplex PCR to identify different types of diarrheogenic E. coli and the presence of characteristics genes of each type.

Results. We identified diarrheogenic strains in the samples of the children from the first month of life. Majority of the strains were enteroaggregative (EAEC) type followed by Enterodifuse (DAEC) type which occupied the second place. We did not find association between the types of maternal diarrheogenic E. coli and that of the children. Interestingly, the identification of the diarrheogenic types did not have relationship with the presence of diarrhea in the subjects all along the period.

Conclusions
The colonization by Enteropathogens seems to be a "normal" event in the studied community without association with diarrheic episodes.
INITIAL IDENTIFICATION AND CHARACTERIZATION OF STRAINS OF KLEBSIELLA PNEUMONIAE ISOLATES FROM A COHORT OF CHILDREN OF 1 - 24 MONTHS OLD

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Background
Bacterial colonization takes place in the first hours of the birth of a baby. Until now, there is a consensus on the fact that newborns are contaminated by the maternal flora, principally when birth is by normal delivery. The Enterobacterias rapidly colonize the gastrointestinal tract, adapt and remain for prolonged periods, usually throughout the life of the individual.

Objectives
To follow-up the process of bacterial colonization and adaptation using Klebsiella pneumoniae as a model.

Methods
64 strains of K. pneumoniae isolated from 16 pairs of mother-child during a period of 24 months were studied. Bacterial identification was carried out through conventional techniques while bacterial identity was confirmed by Enterotest system. Also, antibiotic sensitivity tests were performed. Bacterial strains were genotypically characterized using RAPD-PCR and PFGE. PAST Program was employed for the identification of similar groups while the presence of 4 genes that codify bacterial adhesins were identified by means of PCR.

Results. We identified a series of K. pneumoniae clones in the children that are not related with the maternal strains. Two colonization patterns by K. pneumoniae were found. However, we did not find an association of these patterns with the way of birth (normal or cesarean delivery). It was found that along the period studied, there was population exchange of K. pneumoniae in the children studied.

Conclusions
In the case K. pneumoniae, the colonization of the children is not apparently from their mother and the way of their delivery does not influence in the colonization patterns.
NATURALLY OCCURRING VARIANTS OF INVASION FACTORS AND POLYHOSTALITY OF LISTERIA MONOCYTOGENES

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Background

The Gram-positive bacterium *Listeria monocytogenes* causes a serious disease in humans and a wide range of animals. Cell invasion is a critical step of listerial infection. Bacterial proteins of internalin family InlA and InlB are critical for invasion of normally non-phagocyting cells. The previously found correlation between InlA and InlB sequences and a host organism suggested that some naturally occurring variants of invasion factors might facilitate infection of certain hosts for the strains carrying these variants thus providing *L. monocytogenes* polyhostality.

Objectives

To test the suggestion that naturally occurring variants of *L. monocytogenes* invasion factors provide higher tropism to certain hosts.

Methods

Isogenic recombinant strains were obtained by cloning of *inlB* alleles found in human and animal isolates into the *L. monocytogenes* EGDe\Delta inlB strain. To exclude regulatory effects, *inlB* s were put under the control of the EGDe \( P_{inlAB} \) promoter and their secretion was driven by a heterologous signaling peptide. Human cells Hep-2 and Hek293 and murine cells C26 were used for the invasion assay.

Totally 8 *inlB* alleles were cloned from human and animal isolates into the same genetic background. The VIMHA004 and the VIMPHk23 *inlB* alleles provided the highest invasion of human and murine cells, respectively. The VIMHA004 and VIMPHk23 variants differed by substitutions Q72H in the N-terminal CAP-domain, N73S, V91I, I138L, P164L in the receptor-binding LRR-domain, T251M in the IR-domain.

Conclusions
*L. monocytogenes* InlB variants carrying substitutions in the receptor-binding domain LRR and the CAP and IR-flanking domains demonstrated different efficiency of invasion of human and murine cells.
Background

*E. coli* O157:H7 is one of the most feared zoonotic pathogens that can cause life threatening human disease such as the haemolytic uremic syndrome (HUS). Food sources of animal origin especially cattle, are the main cause of infection but many genotypes exist within this serotype of which their contribution to the clinical outcome of infection in humans is not well known.

Objectives

1- Determine the frequencies of distribution of different genotypes among *E. coli* O157:H7 strains recovered from food and human clinical sources using a combination of molecular subtyping methods and assess whether specific genotypes are overrepresented in human clinical strains. 2- Determine the potential correlation of genotype clusters with severe clinical symptoms.

Methods

A diverse collection of non-clonal human (*n*=100) and food (*n*=70) isolates from Belgium was characterized by conducting different molecular assays that have previously demonstrated non random distribution of *E. coli* O157:H7 genotypes among clinical and non-clinical isolates: lineage-specific polymorphism assay, Shiga-toxin-encoding bacteriophage insertion site assay, clade typing, *Tir* SNP assay, and variant analysis of Shiga toxin 2 and antiterminator *Q* genes. Genetic clustering of all data was performed using MCMC model.

Conclusions
Our results supported the epidemiological evidence of existence of clinically significant microbial genetic markers that are overrepresented among *E. coli* O157:H7 clinical isolates and which are important determinants of human infection risk. Furthermore, genetic clusters 1 and 3 were observed to be more frequently isolated from bloody diarrhea and haemolytic uremic syndrome, respectively. Our results may contribute to monitoring of genotypes with high virulence/transmission potential.
DEVELOPMENT OF MICROSATELLITE DNA MARKERS FOR LACHANCEA THERMOTOLERANS

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Background

*Lachancea thermotolerans* is an important component of the grape/wine ecosystem and one of the first non-*Saccharomyces* yeasts recently commercialised as starter culture in winemaking. Therefore, there is an emerging need for a robust, rapid, and cost effective typing system to be used in ecological and technological studies.

Objectives

The complete genome sequence of the *Lachancea thermotolerans* was examined for development of variable number tandem repeat (VNTR) markers suitable in detecting genetic polymorphisms and discriminating among genotypes.

Methods

The Tandem Repeats Finder program was applied to detect microsatellites within the *L. thermotolerans* CBS 6340 genome (www.genolevures.org). After an initial evaluation of potential VNTR markers using standard PCR and agarose gel electrophoresis, 10 markers were retained for further evaluation. Forward primers were labeled with different fluorescent dyes and PCR products were run in an ABI 3730 XLs DNA Analyser.

Conclusions

Ten polymorphic VNTR markers were analyzed using 48 *L. thermotolerans* isolates from geographical distant vineyards in Greece. These markers have been carefully selected to be widespread in different chromosomes, so as to be appropriate for phylogenetic analysis. They show a high degree of polymorphism with multiple alleles easily scored, some of which even on agarose gels. Thus, they are quite promising for investigating within and among population variability.

This work is co-financed by the European Regional Development Fund (ERDF) of the EU and by National Resources under the Operational Program Competitiveness and Entrepreneurship (EPAN II), Action “COOPERATION 2011”.
GENETIC INSIGHTS INTO VEGETATIVE COMPATIBILITY GROUPING IN VERTICILLIUM: CHALLENGING THE MYTHS

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Background
Verticillium dahliae is a notorious phytopathogenic ascomycete that causes vascular wilt in hundreds of plants worldwide. Classification of isolates into Vegetative Compatibility Groups (VCGs) has been widely used over the past 3 decades as a convenient way to describe population structures and predict pathogenicity potential. This method is based on the general assumption of genetic isolation between VCGs; heterokaryon formation is thus considered possible only between members of the same VCG.

Objectives
To test the traditional hypothesis of inter-VCG genetic isolation in V. dahliae through genetic investigations and phylogenetic analyses.

Methods
(i) Complementation tests using nitrate-nonutilizing (nit) and other mutant types of V. dahliae isolates, either by the standard procedure performed on minimal medium agar or by a novel high-throughput method in liquid microcultures. (ii) Microscopic investigations of self- and non-self heterokaryon formation between isolates labeled with nuclear fluorescent proteins. (iii) Phylogenetic analysis of a V. dahliae collection comprising members of all VCGs, using a highly polymorphic sub-region of the ribosomal IGS region.

Conclusions
True heterokaryotic reactions can be forced between V. dahliae members of different VCGs by using appropriate mutant types and assay conditions. These heterokaryons are frequently unstable, with uneven distributions of genetically dissimilar nuclei tending to segregate into novel homokaryotic mycelia. Most importantly, inter-VCG heterokaryon formation implies the absence of strict genetic barriers between V. dahliae VCGs and/or subgroups. These findings are consistent with the results of phylogenetic analyses of VCGs, which demonstrate a lack of correlation between the VCG classification system and intra-specific genetic groups.
GENETIC POPULATIONS ANALYSIS OF XYLELLA FASTIDIOSA FROM CITRUS AND COFFEE PLANTS SYMPATRICALLY GROWN IN SAO PAULO STATE, BRAZIL

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Background
In Brazil Xylella fastidiosa (Xf) bacteria are responsible by diseases in economically important crops like sweet orange causing the Citrus Variegated Chlorosis and coffee causing the Coffee Leaf Scorch. Population-structure-based studies focusing on the systemic-colonizer, vector-depended, multi-host plant pathogenic bacterium Xf has provided remarkable information about their ecology and biology, specially for the citrus infecting Xf populations. In contrast, no information is currently available about the genetic structure of Xf populations in coffee and uncertain information are known about the cross infection in citrus and coffee plants by Xf.

Objectives
To determine the genetic structure of Xf populations infecting coffee trees both the genotypic and genetic diversity. Also, assay using cross infection between coffee and citrus strains were realized. Finally, we also like to understand how both hosts and agricultural management has shaped the ecology of both Xf populations.

Methods
We sampled symptomatic coffee plants from geographic regions sympatric to citrus plantations in Sao Paulo State, Brazil, and isolated hundreds isolates from each region. This isolates were genotyped using 14 genomic microsatellite markers.

Conclusions
The coffee-associated Xf populations had higher gene diversity and allelic richness than citrus, typical of an evolutionarily older population. Even sharing the same vectors and geographic area Xf from coffee and citrus are genetically and biologically different populations with no admixture of strains. Xf from coffee were geographically structured similarly as the sympatric populations from citrus. Cross infection were successful only in coffee plants inoculuted with Xf from citrus, but no colonization were observed after 13 months.
L-ASPARTIC ACID CONSUMPTION AS A BASE FOR AMINO ACIDS AND THEIR DERIVATES PRODUCTION BY CORYNEBACTERIUM GLUTAMICUM

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Background
Corynebacteria are Gram-positive Actinobacteria inhabitants of diverse ecological niches. The most industrial-interesting species is *Corynebacterium glutamicum*, which is involved in amino acids production as pharmaceutical products and mainly as food/feed enhancers. This bacterium produces per year more than 2.93 million tons of L-glutamic acid and 1.95 of L-lysine. However, the coryneform bacteria entered the 21st century with a refocusing of their industrial point of view. Thus, relevant bioproducts such as chemicals and biodegradable polymers, as well as ingredients or additives in food, feed, cosmetics and pharmaceuticals have extended the portfolio of *Corynebacterium* [e.g.: cadaverine].

Objectives
The aspartate pathway is responsible for the biosynthesis of a high number of metabolites of biotechnological importance such as L-lysine, L-threonine, L-methionine and L-isoleucine, as well as other close related compounds: the amino acid L-valine, the vitamin D-pantothenate or the biogenic amine cadaverine (1,5-diaminopentane). The last one is the proof-of-concept compound of the ERA-IB 3\(^{rd}\) joint call supported project named SCILS (http://www.era-ib.net/scils).

Methods

*C. glutamicum* is able to use different alternative nitrogen compounds as L-glutamine or creatinine, but little is known about the possibility of L-aspartic acid utilization. The 2D-DIGE analysis results, as well as a microarrays study of an aspartate-consumption defective mutant, have shown that *C. glutamicum* is able to use L-aspartic acid as nitrogen source.

Conclusions
The bioconversion to different amino acids, which can be the precursor of industrial relevant biomolecules (e.g.: L-lysine decarboxylation yields cadaverine), is the main explanation of the aspartate use.
A BLUEPRINT OF SULFUR METABOLISM FROM THE PROTEOME OF HYDROGEN-PRODUCER THERMOCOCCUS ONNURINEUS NA1

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Background
The hyperthermophilic archaeon, Thermococcus onnurineus NA1, has been shown to produce biohydrogen (H₂) when using CO, formate, or starch as a growth substrate. This strain can also utilize elemental sulfur (S⁰) as a terminal electron acceptor for heterotrophic growth and reduce it to H₂S.

Objectives
To gain insight into sulfur metabolism, the proteome of T. onnurineus NA1 cells sampled under sulfur culture conditions was quantified and compared with those under formate, CO and starch culture conditions.

Methods
Using label-free nano-UPLC-MS²-based comparative proteomic analysis, we found that approximately 38.4 % of the total identified proteome (589 proteins) were significantly up-regulated (≥1.5 fold) under sulfur culture conditions.

Conclusions
Many of these proteins are functionally associated with carbon fixation, Fe-S cluster biogenesis, ATP synthesis, sulfur reduction, protein glycosylation, protein translocation, and formate oxidation. Our results also revealed markedly lower expression levels of enzymes involved in the sulfur assimilation pathway, as well as cysteine desulfurase under sulfur culture condition. The present results provide a first global atlas of the proteome changes triggered by sulfur. This study will contribute to a better understanding of the mechanisms and physiology of T. onnurineus NA1 cells in response to sulphur.
PROTEOGENOMICS OF THE NUTRITIONALLY VERSATILE, MARINE, SULFATE-REDUCING BACTERIUM DESULFOCOCCUS MULTIVORANS

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Background

Sulfate-reducing bacteria play a major role in marine carbon and sulfur cycling. Members of the metabolically versatile deltaproteobacterial Desulfococcus/Desulfosarcina-cluster are abundant in marine sediments and therefore of ecophysiological importance. A metabolically versatile representative of this cluster is Desulfococcus multivorans that is able to completely oxidize a variety of organic acids, including fatty acids up to C₁₄, as well as aromatic compounds under anoxic conditions.

Objectives

Proteomics-enhanced genome annotation of D. multivorans was based on cells adapted to 17 different substrate conditions and aimed at metabolic reconstruction and deciphering the regulatory network.

Methods

Manual revising of automatically predicted and annotated protein-coding sequences (CDS) was combined with proteomic data (2D-DIGE, shotgun, membrane protein-enriched fraction) of substrate-adapted cells to allow for improved functional prediction.

Conclusions

The closed genome sequence consists of a 4.45 Mb circular chromosome with 3946 predicted CDS. The metabolic versatility of D. multivorans is reflected by the high number (269) of encoded proteins involved in organic carbon catabolism, 156 of which were detected. Out of the 324 genomically predicted transport proteins 104 could be detected in the membrane protein-enriched fraction. The presence of >200
genes encoding proteins involved in signal transduction should facilitate from the basis of a regulatory network for adapting to changing environmental conditions. Accordingly, out of 1247 identified proteins, only 220 were present under all analyzed substrate conditions, unravelling a high level of substrate-dependent regulation in *D. multivorans*. Overall, the study underpins the value of proteomic analysis to enhance functional genomic predictions, providing new perspectives on our metabolic understanding of sulfate-reducing bacteria.
Background
Fungi are known as active degraders of lignocellulosic materials in nature, soil, compost and forest topsoil. The method for degradation depends on the fungus genus with different results. The key for their degradation capacity resides in the fact that the extracellular enzymes have different capacities more or less active according to the lignocellulosic substrate. These enzymatic extracts are used in biotechnological processes in the industry.

Objectives

Penicillium chrysogenum B13 strain (CTM2012-32026 project) is an overproducer of feruloyl esterase (FAE) able to hydrolyze the ester bonds between the hydroxycinnamic acids and the plant cell wall polysaccharides. As a result of FAE activity, the polysaccharides are more accessible, allowing complete digestion by the rest of the enzymes involved in the plant cell wall degradation.

Methods

P. chrysogenum B13 strain was grown in presence of sugar beet pulp and the enzymatic extract secreted to the culture medium was obtained at the time of greatest FAE activity. Two-dimensional gel electrophoresis of proteins was performed and subsequently proteins were identified by means of a MALDI-TOF mass spectrophotometer system.

Conclusions

The analysis of the results showed that the FAE activity is increased by 9.27% with respect to the wild strain P. chrysogenum Wisconsin 54-1255 and that 42.22% of the secreted proteins are
enzymes associated with the degradation of plant cell wall, compared to only 11.59% when the culture medium does not contain plant substrate.

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IDENTIFICATION OF REDOX PARTNER OF HELICOBACTER PYLORI HP0377 PROTEIN.
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Background
Helicobacter pylori is the common human pathogen colonizing the gastric epithelium of humans with severe consequences. HP0377 is a Dsb thiol oxidoreductase which acts as CcmG protein, involved in the maturation of apocytochrome c. CcmG proteins are kept in the reduced form by an integral membrane protein DsbD or its shortened analogue, CcdA. Both proteins catalyze the transfer of electrons from cytoplasmic thioredoxin across the inner membrane to the periplasm. H. pylori does not encode DsbD, however it encodes homologue of CcdA (HP0265).

Objectives
The aim of this study was to establish whether HP0265 is responsible for HP0377 rereducing.

Methods
To inactivate hp0265, the allelic exchange strategy was used. A recombinant plasmid pUWM2019 (based on a vector non-replicating in Helicobacter cells) constructed by a two step PCR method, was used for mutagenesis. It contains the hp0265 gene disrupted by insertion of a kanamycin resistance cassette into the gene coding sequence. We then tested whether deletion of hp0265 affects the redox state of HP0377 using AMS trapping strategy This agent can only modify covalently free thiols, resulting in a molecular mass increase of about 490 Da.

Conclusions
We found that H. pylori HP0377 is rereduced by HP0265. Obtaining the hp0265 mutant was dependent on presence of the reducing agent, DTT. Moreover, the growth of the hp0265 mutated strain without the presence of DTT was much slower than the growth of wt strain, which indicates the importance of HP0265 and its reducing activity for bacterial survival.
A METAPROTEOMIC APPROACH FOR ANALYSIS OF MICROBIAL COMMUNITY STRUCTURE AND FUNCTION FOR IMPROVEMENT OF BIOGAS PLANT PERFORMANCE

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Background

To avoid downtime in operation and to further optimize operation of biogas plant (BGP) profound understanding of the composition and function of microbial community converting the biomass to methane is required.

Objectives

Our main goal is to understand how process conditions determine the process performance, the taxonomic and functional composition of the microbial community and vice versa. For such investigations, the quantification of the composition of the active community is essential. In particular, the presence of key enzymes of metabolic pathway correlates well with the community activity. Thus, the present study investigated the metaproteome of 40 industrial-scale BGPs.

Methods

The applied metaproteome workflow involved protein extraction using liquid phenol, tryptic digestion, peptide separation by liquid chromatography coupled to tandem mass spectrometry (Velos Orbitrap Elite), and data analysis with the MetaProteomeAnalyzer software [1].

Conclusions

For each BGP, about 500 proteins, covering the main metabolic pathways of the biogas process, namely hydrolysis, fermentation, acetogenesis and methanogenesis, were identified. The microbial communities detected by metaproteome analysis of the BGPs clustered according process temperature (mesophilic and thermophilic) [2,3], and substrate composition.

Application of metaproteomics allowed the acquisition of profound knowledge about community structure and function of microbial communities and contribute to our understanding of the conversion of biomass into methane. This knowledge could be used to improve monitoring and control of BGP, and to support development of new BGP designs.
IDENTIFICATION OF PEPTIDASES GENERATING CONSTITUTIVE
COLLAGENOLYTIC ACTIVITY OF GEOBACILLUS THERMOLEOVORANS
DSM15325
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Background
While the main biological importance of collagen degradation for prokaryotic
pathogens is tightly connected with virulence, thermophilic bacteria use
collagenolysis when exploit collagen for nutrition. To date the view that collagen
catabolism by thermophilic bacteria is a result of a cascade of synergistically active
inducible peptidases is prevailing. Usually this model of collagen degradation omits
the thermophilic bacteria constitutively produced collagenolytic peptidases. The
constitutive collagenolytic activity of thermophilic bacteria was never deliberately
analyzed and the understanding of constitutive peptidases importance for collagen
hydrolysis in vivo is obscure, what reduces the overall fundamental understanding of
collagenolysis biological roles for prokaryotes.

Objectives
Detect hydrolytic activity of Geobacillus thermoleovorans DSM15325 constitutively
produced collagenolytic peptidases. Identify detected hydrolases by mass
spectrometry.

Methods
G. thermoleovorans DSM15325 was cultivated in modified M9 medium up to late
exponential growth phase when the total secretome proteins were precipitated and
analyzed by SDS-PAGE. The evaluation of collagenolytic peptidases activity was
performed by gelatin zymography and azocollysis. Detected peptidases were excised
from acrylamide gel and subjected to mass spectrometry analysis. Obtained enzyme
sequences were analyzed in silico.

Conclusions
The Bacillolysin (protein ID AEV20496.1), Pz-peptidase A (protein ID BAD99433.1)
and B (protein ID BAD99434.1) were identified as peptidases generating constitutive
collagenolytic activity of G. thermoleovorans DSM15325.
Background

Bacteria from genus *Cronobacter* are opportunistic pathogens associated with rare but severe infections mainly among neonates. Based on the epidemiological studies and molecular typing methods such as MLST is obvious that *C. sakazakii*, *C. malonicus* and *C. turicensis* are prevalent in neonatal infections and particularly sequence types ST4, ST7 and ST19 are considered as an attribute of virulence strains.

Objectives

The aim of this study was to map membrane of virulent strains being as the gateway to the host organism and to contribute to the knowledge of virulence features of this genus.

Methods

We applied several techniques for membrane proteins isolation such as the whole cell membrane isolation by phenol extraction method, the membrane sub-fractionation into outer membrane, periplasm and inner membrane, and the immunoprecipitation using polyclonal antibodies against whole *Cronobacter sakazakii* cells. The isolated proteins were consequently identified by mass spectrometry and peptide mass fingerprinting method.

Conclusions

Membrane proteins play a pivotal role in bacterial pathogenesis thus a complex study of membrane proteome of pathogens is invaluable. Here we present membrane proteomes report of the strains considered as pathogenic focusing on virulence traits determination.
ANALYSES OF ACTIVATED SLUDGE METAPROTEOMES OF A WASTEWATER TREATMENT PLANT DURING WINTER AND SUMMER PERIOD

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Background
Molecular biological approaches (i.e. community fingerprinting, metagenome sequencing) revealed that only a small portion of the microbial community typically found in wastewater treatment plants (WWTP) can be cultivated in the laboratory. In order to understand its function without isolating single strains, metaproteomics analysis has been applied to activated sludge samples.

Objectives
The impact of changes in temperature (winter and summer period) on metaproteomes of activated sludge samples derived from a full-scale WWTP was investigated to enable a detailed description on functional level.

Methods
For reduction of sample complexity prior to MS analysis, different fractionation techniques including reversed phase liquid chromatography (RP-LC), sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE), and liquid isoelectric focusing were combined and compared regarding effort and quality of resulting protein identifications. The derived spectra were identified using the UniProtKB/Swiss-Prot protein database. Additionally, identified KEGG ontologies (Kyoto Encyclopedia of Genes and Genomes) and Enzyme Commission numbers were used to plot proteins hits into pathway maps of the central carbon and nitrogen metabolism.

Conclusions
As expected, the high resolving but most time consuming three-dimensional approach yielded the highest amount of protein identifications. The results revealed taxonomic differences in the abundance of specific classes, such as Beta- and Gammaproteobacteria, in activated sludge collected during winter and summer period. Fractions of eukaryotic proteins decreased from winter to summer period, indicating a better degradation performance during summer period due to increasing process temperature. Additionally, more proteins associated to the nitrogen
metabolism, like hydroxylamine dehydrogenase or ammonia monooxygenase, were identified in summer period.
CALCIUM-TRYPTOPHAN INTERACTIONS IN CALCIUM-BINDING PROTEINS

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Background
Calcium ion is the important structural element in bacterial cells and is also involved in cell signaling. Also, the calcium binding may enable protein folding and enzymatic activity. Aromatic amino-acids in proteins can stabilize the native protein structure via pi-pi interactions and may also play a role in calcium binding by cation-pi interactions. Terbium (Tb³⁺) is the phosphorescent lanthanide that may be used as a calcium binding analog. Tb³⁺ increases the quantum yield as it binds to calcium binding sites (in proteins, DPA), which enables spectroscopic ion binding measurements. In our previous studies of FrpC protein of Neisseria meningitidis we found the calcium binding as a driving force for FrpC folding. The two Trp residues of FrpC also play an important role as stabilizing factor in calcium binding and enzymatic activity, the autocatalytic cleavage.

Objectives
In order to estimate the importance of calcium-Trp interactions in proteins, we analyzed PDB crystal structures possessing both Trp residues and bound calcium.

Methods
In all calcium-containing-protein-structures in PDB (n=6403) relative Ca/Trp positions were analyzed (392000 cases). FrpC studies: Steady-state fluorescence spectroscopy.

Conclusions
We confirmed that in calcium-binding enzymatic domain of FrpC, calcium is bound in the close proximity to Trp residues that supports the calcium-Trp interaction. Using PDB database search, we found that such calcium-Trp position is rather rare among protein structures and that calcium is most frequently positioned toward nitrogen hetero-atom in indol.
QUANTITATIVE SHOTGUN PROTEOMIC APPROACH FOR THE STUDY OF S.COELICOLOR M145 AND S.LIVIDANS TK24, TWO PHYLOGENETICALLY CLOSELY RELATED STRAINS WITH VERY DIFFERENT ABILITIES TO PRODUCE ANTIBIOTICS

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Background
Streptomyces are filamentous Gram positive soil bacteria of industrial interest due to their ability produce two-thirds of all known antibiotics as well as other so-called "secondary metabolites" useful to human health or agriculture. A reverse correlation between the content in TriAcylGlycerol (TAG) and the ability to produce antibiotics was recently demonstrated. In the presence of glucose, the lipid content of S. coelicolor is less than 10% its dry cell weight (DCW) indicating that its TAG content is very low and abundantly produces antibiotics whereas the total lipid content of S. lividans is up to 30% its DCW and does not produce antibiotics. Interestingly, in the presence of glycerol, the two strains have the same lipid content (about 30% of their DCW).

Objectives
In order to understand the molecular basis of these phenotypical differences between these two Streptomyces species, we developed a novel gel-free shotgun proteomic approach.

Methods
This strategy involves a simple protein extraction followed by proteolysis in presence of trypsin and LysC and LC-MS/MS analysis by Q-exactive MS (Thermo Fisher). We performed a three-factor analysis of variance that included: strain factor (S.lividans/S.coelicolor); carbon source factor (glucose/glycerol), and time factor (36h/48h/72h).

Conclusions
More than 2500 proteins were identified corresponding approximately to 30% of the genome. This study showed different pattern of expression of genes of central carbon and lipids metabolism as well as antibiotics biosynthesis, upon growth on glucose or glycerol. These results revealed some potentially interesting genes to be genetically modified for further improvement of antibiotics and/or TAG production by Streptomyces species.
THE SMALL PROTEIN SCO2038 CONTROLS STREPTOMYCES COELICOLOR DIFFERENTIATION BY MODULATING TRYPTOPHAN BIOSYNTHESIS

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Background

In Streptomyces coelicolor amino acid metabolism is an important clue of the morphological and physiological differentiation program and, differently from other bacteria, the expression of amino acid biosynthetic genes is not subjected to end-product negative regulation. In some amino acid biosynthetic gene clusters, such as tryptophan, histidine and proline, small orfs (about 100-300 nucleotides) were identified. These small orfs, such as sco2038, encode proteins whose cellular role have to be elucidated to highlight possible novel and crucial molecular mechanisms controlling amino acid synthesis and, thus, differentiation program.

Objectives

The aims of this work are:

1. the understanding of the effects exerted by tryptophan on primary metabolism, morphological differentiation and antibiotic production;

2. the study and characterization of the SCO2038 function as modulator of tryptophan biosynthesis.

Methods

- Differential proteomic analysis based on 2D-DIGE and MS procedures.
- SEM analysis.
- Generation and characterization of sco2038 mutants
- Identification of potential SCO2038 interaction partners by pull down assay coupled with MS identification and Bacterial Adenylate Cyclase Two Hybrid System.
- qRT-PCR analysis.

Conclusions
The obtained results revealed that tryptophan controls the expression of metabolic and regulatory proteins and promotes aerial mycelium formation, spores production and actinorhodin antibiotic biosynthesis. Moreover, the small orf sco2038, encodes a 7 KDa protein playing a key role in modulating tryptophan biosynthesis and thus, morphological differentiation. In the light of these results we propose to rename sco2038 as *trpM*, the gene encoding the tryptophan biosynthesis Modulator TrpM.
SLP PROFILES IN ENDEMIC AND NON-ENDEMIC CLOSTRIDIUM DIFFICILE PCR-RIBOTYPES

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Background

*Clostridium difficile* is divided into approximately 400 PCR-ribotypes, some of them are widespread and associated with severe disease. *C. difficile* PCR-ribotypes 014/020 and 002 are the most common among endemic PCR-ribotypes.

Many bacteria, including *C. difficile*, express surface layer protein (Slp). This protein forms surface-exposed proteinaceous layer which mediates interactions with the external environment, i.e adhesion to host cells as in case of *C. difficile*. Two *Slp* proteins with molecular weights (MWs) of 32-38 kDa (low MW) and 42-48 kDa (high MW) are expressed and vary in size in different *C. difficile* strains. Slp typing is described, but not widely used.

Objectives

The aim of the study was to compare slp profiles of two endemic PCR-ribotypes (014/020 and 002) and selection of non-endemic PCR-ribotypes.

Methods

Eighty-eight *C. difficile* isolates from 21 different PCR-ribotypes were included. Extraction of Slp proteins was performed by using low-pH glycine solution and further analyzed by SDS-PAGE.

Conclusions

Five different slp profiles (1-5) were identified. Slp profiles 1 and 2 were the most frequent and present in the majority of PCR-ribotypes. Endemic and non-endemic
PCR-ribotypes shared 3 out of 5 slp profiles. Two remaining slp profiles were present only in one strain each.

Our results indicate that slp profile does not contribute significantly to *C. difficile* colonization properties and endemicity.
Background

The aim of the project is to develop biomarkers to help distinguish between the inner and the outer membrane of *Escherichia coli*. The fluorescent proteins mCherry, *E. coli* Flavin binding Fluorescent Protein and superfolder GFP, were used as biomarkers. These were directed to the inner or outer membrane using signal sequences from lipoproteins or by fusion with membrane proteins. The fusion partners were the outer membrane protein A (OmpA), the artificial TAT-lipobox signal sequence for transport to and lipid anchoring in the outer membrane, the major coat protein of phage pf3 that is inserted into the inner membrane, and mistic, a integral inner membrane protein from *Bacillus subtilis*.

Objectives

The aim of the project is to develop a bacterial strain (*E.coli* BL21 DE3) with optimized expression levels of the fluorescent markers for co-localization studies of bacterial membrane proteins.

Methods

The methods used in the project include molecular biology (cloning, PCR), microbiological (culturing, recombinant protein expression) and biochemical (cell fractionation, membrane isolation) techniques. In addition, the project will also use analytical methods such as fluorometry and fluorescence microscopy.

Conclusions

The fluorescent marking of the two membranes makes membrane fractions easier to detect after membrane separation. It also help to assess the quality of the separation procedure.
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CHARACTERISATION OF THE BSA TYPE THREE SYSTEM SECRETOME OF BURKHOLDERIA PSEUDOMALLEI USING HYPER-SECRETING MUTANTS
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Background
Many Gram-negative bacteria utilise Type III secretion systems (T3SS’s) to deliver effector proteins into target host cells where they hijack cellular processes for their own benefit. T3SS’s form an injectosome with proteins spanning both membranes of the bacterium and an external needle through which effector proteins are delivered into the target cell. B. pseudomallei, the causative agent of the tropical disease melioidosis in humans and animals, possesses three distinct T3SS’s, of which the Bsa system has been shown to play a role in invasion of non-phagocytic cells, escape from the endocytic compartment and virulence in murine models of melioidosis. To date few proteins have been proven to be secreted by the Bsa apparatus and, by analogy with other pathogens that deploy T3SS’s, it is likely that there are many other effectors awaiting discovery

Objectives
To characterise the total secretome of B. pseudomallei in standard laboratory media and the repertoire of Bsa-secreted effector proteins.

Methods
We have determined the effector secretome of the Bsa T3SS using hyper-secreting mutants of coupled with iTRAQ, a gel free quantitative proteomics technique.

Conclusions
Our study provides one of the most comprehensive core secretomes of B. pseudomallei described to date and identified 26 putative Bsa-dependent secreted proteins that may be considered candidate effectors.
THE EXCRETORY/SECRETORY PROTEOME OF TRICHOMONAS VAGINALIS INDUCED BY NEUTROPHIL

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Background
The protist parasite *Trichomonas vaginalis* causes one of the most common non-viral sexually transmitted disease in humans, trichomoniasis. Previous studies indicate that neutrophils, leukotriene B₄ (LTB₄) and interleukin-8 (IL-8) are found in the vaginal discharges and vaginal smears of patients infected with *T. vaginalis*.

Objectives
IL-8 is often associated with inflammation, also known as neutrophil major chemotactic factor, causing them to migrate toward the site of infection by *T. vaginalis* and start innate immune response to defense. Previous studies focused on the profiling of chemoattractants from neutrophils stimulated by *T. vaginalis* or *T. vaginalis* ESPs, but very little is known about the cell-cell interaction between *T. vaginalis* and neutrophil. We hypothesize that both *T. vaginalis* ESPs and direct contact with neutrophil can induce the release of chemoattractants in the vagina.

Methods
In this study, we confirmed that both *T. vaginalis* and ESPs can induce neutrophil to release IL-8. We also used 2-Dimensional SDS-PAGE to investigate the differentially expressed ESPs from *T. vaginalis* before and after contact with neutrophil. A total of 16 differentially expressed protein spots were identified by MALDI-TOF-MS, and discussed the possible roles of identified proteins in the biology of host-parasite interactions.

Conclusions
We established a representative 2-DE map of ESPs of *T. vaginalis* induced by neutrophil and identified differentially expressed protein that may promote cytoadherence and long-term colonization while ESPs from neutrophil may be related to implicate in the first line of host defense.
Background

Mycoparasitic species of the filamentous fungus *Trichoderma* are commercially applied as biological control agents against fungal plant pathogens. The mycoparasitic interaction of *Trichoderma* with the host fungus comprises specific recognition events leading to activation of "molecular weapons" in the mycoparasite which are involved in host attack and lysis and result in utilization of the host fungus as a nutrient source for the mycoparasite.

Investigations on the underlying intracellular signal transduction pathways of *Trichoderma atroviride* revealed the involvement of a Mitogen-activated protein kinase (MAPK) which is essential for triggering the mycoparasitic Response. Mutants missing the *tmk1* gene show reduced mycoparasitic activity against host fungi.

Objectives

The aim of the presented study was the identification of target proteins being regulated by the Tmk1 MAPK pathway upon host recognition. To this end, *T. atroviride* wild-type and the delta-*tmk1* mutant were co-cultivated with the host fungus *Rhizoctonia solani* and *Trichoderma* mycelia were harvested from the confrontation zone upon direct interaction between the two fungi and from respective un-induced (self-confrontation instead of confrontation with the host) controls.

Methods

Two-dimensional Difference Gel Electrophoresis (2-D DIGE) was applied for comparatively analysing the proteomes of the four samples with the aim to find mycoparasitism-relevant targets of the Tmk1 MAPK.
Conclusions

Bioinformatic analyses revealed 60 proteins being regulated in the wild-type response (WT vs. *R. solani* minus WT vs. WT) and 126 proteins being host-induced Tmk1 targets (WT vs. *R. solani* minus delta-tmk1 vs. *R. solani*), of which 30 and 60 proteins, respectively, were identified by mass spectrometry.
PROTEOMICS ANALYSIS OF THE B. SUBTILIS INNER SPORE MEMBRANE

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Background

The endospore is the dormant form of Bacillus subtilis and many other Firmicutes. By sporulation these spore formers can survive very harsh physical and chemical conditions. Yet they need to go through germination to return to their growing form. The spore inner membrane (IM) has been shown to play an essential role in triggering the initiation of germination; however, its protein composition was not comprehensively characterized yet.

Objectives

To isolate IM from B. subtilis spores and to characterize the B. subtilis spore IM proteome.

Methods

The spores were chemically decoated and enzymatically deprived of the cortex. Subsequently the core was mechanically disrupted and the IM fraction was collected by differential centrifugation. Tryptic peptides, acquired from in-gel digestion, were analyzed using Ion trap LC-MS/MS, which led to identification of the spore IM proteins.

Conclusions

We have adapted the spore IM isolation protocol to proteomics studies. Over 900 proteins could be identified from the B. subtilis spore IM preparations, in which ca. one-third were predicted to be membrane proteins. In addition to the previously known IM proteins, a number of novel IM proteins were identified which are likely to provide new insights into the IM protein composition, its functions and in particular into the spore germination machinery.
Background

The presence of plasmids carrying class 1 integrons in a collection of psychrotolerant enterobacteria isolated from Fildes Peninsula was previously established. In these plasmids, we could identify a DNA region with three ORFs (intI1, dfrA14 and hyp, encoding a hypothetical protein) highly similar to a fragment originally identified in plasmid pKOX105 from Klebsiella oxytoca (HM_126016).

Objectives

DNA sequence analysis of plasmids purified from some of these enterobacterial isolates.

Methods

Plasmid DNA from isolates CDTR5, CN11 and HP19 were purified with Gene Plasmid Midprep Elute® HP (Sigma-Aldrich Co, MO) and sequenced in a Ion 314TM Chip on a Ion PGM™ System (Life Technologies,) at the IIBCE. The reads were assembled using CLC Genomics Workbench v6.5 (CLC bio). Annotation of pHP19 contigs and comparison with plasmids pECL_A and pKOX105 were done using RAST server (http://rast.nmpdr.org/rast.cgi).

Conclusions

Sequence analysis of pHP19 showed a region of 7181 nts highly similar (99% identity by Blastn) with a region of pKOX105 (54641pb). This contig includes the intI1 gene and other ORFs encoding a DNA-cytosine methyltransferase and a probable membrane protein.

The sequences of other plasmidic regions were similar to those of Enterobacter cloacae ATCC 13047 plasmid pECL_A (IncFII) (199562pb) (NC_014107), including genes for replication, stability and conjugation. Other cluster genes related with mercury, arsenic, tellurite, nickel and copper resistance identified were identical to those found in pECL_A. Thus this plasmid is likely a mosaic of portions of plasmids
that have been characterized in mesophilic enterobacteria, showing that HGT between these bacteria and the Antarctic microbiota is rather common.
Background

The intracellular nucleotide c-di-GMP has recently emerged as one of the central elements of the signal transduction network linking perception of environmental or intracellular cues to specific alterations in cellular function. In response to different input signals, the opposing activities of diguanylatecyclases (DGCs) and phosphodiesterases (PDEs) control the cellular concentration of c-di-GMP. Inside the cell, c-di-GMP binds a currently unknown number of effector molecules that subsequently interact with cognate target components to produce a variety of output phenotypes (e.g. transition from planktonic to biofilm). Given the multiplicity of c-di-GMP related genes in most bacterial genomes and their scattered distribution within the genomes, the elements that belong to the same c-di-GMP control module are difficult to uncover.

Recently, an integrative conjugative element -ICEAcaTY.2- has been characterized in the extreme acidophile Acidithiobacillus caldus ATCC 51756. ICEAcaTY.2 is an actively excising element of widespread occurrence in a collection of At. caldus strains, that behaves as a cohesive heritable unit. Bioinformatic analysis of ICE.2-type elements has revealed that c-di-GMP synthesis/degradation functions and putative c-di-GMP binding effectors and targets co-occur within the element.

Objectives
To gain insight into the participation of c-di-GMP genes encoded by ICEAcaTY.2 as a functional control module.

Methods
To test this hypothesis the expression of the c-di-GMP genes encoded by ICEAcaTY.2 was quantified by real-time PCR under different growth conditions (e.g. planktonic, attached, stressed).

Conclusions
Transcriptional behavior of key c-di-GMP module components in attached cells and upon DNA damaging conditions support the hypothesis and further suggest that this control module could be critical for biofilm development and DNA exchange.
ENVIRONMENTAL ISOLATES OF V. PARAHAE~MOLYTICUS CARRY SXT/R391 INTEGRATIVE CONJUGATIVE ELEMENTS

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Background

SXT/R391 Integrative conjugative elements (ICEs) are self-transmissible mobile genetic elements able to confer adaptive features to bacterial hosts, including Vibrionaceae. They share a conserved genetic scaffold encoding their own conjugation, integration, excision, and regulatory machinery, interspersed with variable DNA clusters located in conserved hotspots.

Objectives

In silico analysis of 96 V. parahaemolyticus genomes was performed to identify Integrative Conjugative Elements (ICEs) of the SXT/R391 family.

Methods

Four novel ICEs were detected in environmental non-pandemic V. parahaemolyticus strains isolated in China, Spain, and Malaysia (2002-2008). ICE sequences were annotated using the RAST annotation pipeline and comparative analysis was performed with the Artemis Comparative Tool, BLASTN and BLAST-PSI. Given their unique genetic contents they were named: ICE VpaSpa1 (~111 kb), ICE VpaMal1 (~70 kb), ICE VpaChn4 (~66 kb), ICE VpaChn5 (~86 kb). We identified gene sequences from other ICEs as well as unique genetic features such as a set of heat-shock proteins and chaperones likely to respond to environmental stress (ICE VpaSpa1); genes with transpeptidase domains belonging to the beta-lactamase TEM family (ICE VpaSpa1); putative type 1 restriction/modification systems (ICE VpaChn5, ICE VpaSpa1); type III restriction-modification systems (ICE VpaMal1); ars operon encoding an efflux system mediating arsenic resistance (ICE VpaMal1, ICE VpaChn4); and an antibiotic resistance cluster carrying floR, strAB and sul2 genes (ICE VpaChn5).

Conclusions

V. parahaemolyticus ICEs exhibit significant genetic polymorphisms, sign that the structure of each ICE has been shaped by frequent recombination, with genetic
content likely dependent on the specific locale in the environment from which it was originally isolated.
OBTENTION OF HYPERACTIVE INTEGRASES BY EVOLVING SYNONYMOUS GENES

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Background

Integrons are gene recruitment platforms that allow for rapid bacterial evolution, playing a major role in the acquisition of antimicrobial resistance genes. The integrase catalyzes the reaction between attC and attI sites. However, its activity is finely tuned to preferentially process attI x attC and attC x attC reactions, as they allow for the acquisition and rearrangement of cassettes, rather than the attI x attI reaction. Being a less frequent phenomenon, the structural basis of attI recognition has yet remained elusive.

Objectives

We have conducted directed evolution experiments on the integrase to obtain hyperactive mutants for the attI x attI reaction. In order to explore a broader evolutionary landscape of the integrase we have recoded the protein into two alternative, yet synonymous, alleles of the intI1 gene.

Methods

In a first round of experiments we obtained among the three alleles 8 mutations conferring an increase in recombination rates. We reinserted all mutations into the three wt alleles, obtaining a 100-fold increase in recombination efficiency. Re-evolving these alleles yielded new mutants with recombination rates for attI x attI comparable to those of the wild type protein for the attI x attC reaction. Along these experiments we obtained landscape-specific mutations and the re-evolution of a codon to a residue beyond its evolutionary landscape in any of the three starting alleles.

Conclusions

We have successfully explored an enlarged evolutionary landscape of the integrase allowing us to obtain hyperactive molecules, while shedding light on the structural features that are important for the recognition of the attI site.
Background

Horizontal gene transfer is an important mechanism in Pseudomonas aeruginosa for adapting to a wide range of environments and, in particular, for acquiring virulence determinants which are mostly carried on pathogenicity islands. One of the largest pathogenicity island of P. aeruginosa is PAPI-1, a 108-kb island which is known to be transferred from a donor to a recipient strain through a type IVb pilus. However, the PAPI-1 acquisition mechanism has not been elucidated to date.

Objectives

We aimed at investigating the mechanism of PAPI-1 acquisition in recipient cells.

Methods

To determine the receptor for conjugative type IVb pilus, a standard PAPI-1 transfer assay described by Carter et al. (2010) was carried out using PA14TnC2 as a donor and 38 PAO1 mutants with an altered lipopolysaccharide (LPS) biosynthesis pathway as recipients. The loss of receptor for conjugative type IVb pilus in PAO1 mutants can result in a significant decrease of PAPI-1 transfer. The conjugative pilus receptor was then confirmed with a competition assay by adding outer-membrane and LPS preparations into the standard mating assay.

Conclusions

Transfer efficiency of PAPI-1 to the AlgC mutant producing a truncated LPS core was reduced by three orders of magnitude compared to that of wild-type PAO1, indicating that LPS acts as a receptor for conjugative type IVb pilus. The homopolymer of D-rhamnose, or rhamnan structure was confirmed as a specific receptor for conjugative type IVb pilus. Further investigation will define if P. aeruginosa that have acquired
PAPI-1 specify a surface exclusion mechanism by LPS modification preventing further recognition by conjugative pilus.
Background

Horizontal gene transfer (HGT) and mobile genetic elements (MGEs) are major generators of genome plasticity and diversity. Certain of these elements can exist in the *Legionella pneumophila* (*Lp*) genome in an excised or integrated form and encode for type IV secretion systems (T4SS) and/or conjugation systems. Intriguingly, each of these elements encodes a homologue of CsrA (carbon storage regulator). CsrA is a global regulator of virulence in *Lp*. The Lvh-region is one of these MGEs encoding a T4ASS and a CsrA homologue.

Objectives

Deciphering the mechanisms of mobility of the Lvh

Methods

Distribution of the Lvh was analyzed by comparative genomics of 50 *Lp* strains, the mobility and regulation of transfer were analyzed by mutagenesis, conjugation experiments, Q-PCR, RNA-Seq and whole genome sequencing of the transconjugants.

Conclusions

The Lvh mobility is mediated by a phage-like integrase. It is transferable at a rate of $10^{-4}$ and inserts specifically in the tmRNA gene in all analysed transconjugants. Using a Δ*dotA* strain we show that the Dot/Icm T4SS is implicated in the conjugation of the lvh-region, as the transfer rate drops to $10^{-7}$. Conjugation experiments in a Δ*ihf* mutant (Integration host factor), a nucleoid-associated protein revealed its role in the mobility of the Lvh. Most interestingly the putative RNA-binding protein LvrC (CsrA homologue) affects the mobility of Lvh by conjugation in specific growth conditions. Thus, The mobility of the predicted self-transmissible Lvh-genomic island is regulated by a complex network of global (IHF) and specific regulatory proteins (LvrC) and depends largely on the Dot/Icm T4SS.
Background

Three ancient Acinetobacter lwoffii strains were aseptically isolated from East-Siberian permafrost sediments and carbon-14-dated to 15,000-30,000 years and 2-3 million years before present.

Objectives

Investigate the structure and properties of plasmids from ancient Acinetobacter lwoffii strains, and the presence of various heavy metal and antibiotic resistance genes on those plasmids.

Methods

Complete genomic sequences (average coverage >30) of three strains were obtained using pyrosequencing method. 13 plasmid sequences were assembled and manually analysed using bioinformatics software and public databases and verified with PCR amplification of regions where ambiguous results occurred.

Conclusions

Each strain carried at least two plasmids with their sizes varied between 6 kb and 287 kb. Three largest plasmids designated pKLH208 (287 kb), pKLH211 (190 kb) and pKLH220 (130 kb) belonged to ED23-35, ED45-23 and VS15 strains respectively. Each of these plasmids encoded similar replication and segregation systems and 12 to 18 IS-elements of IS1, IS4, IS5, IS6 and IS66 families with almost twice as many IS-elements with interrupted trp genes, suggesting frequent transposition processes. Apart from two different mercury resistance operons found only on pKLH208 and pKLH211, large plasmids encoded operons conferring resistance to cobalt, zinc, cadmium, chrome and arsenical compounds, several copies of copper and iron intake regulating operons, urea degrading operons, aadA (streptomycin/spectinomycin) and tetA-tetR (tetracycline) antibiotic resistance genes. We found that 11 of 13 identified plasmids, including two large plasmids pKLH208 and pKLH220, encoded either
mobilisation or conjugation and type IV secretion systems and are potentially capable of transferring horizontally.
COMPARISON OF INTEGRON ATTC SITES: THEIR STRUCTURAL FEATURES DIFFERENTIALLY INFLUENCE THE FREQUENCY AND ORIENTATION OF CASSETTE INTEGRATION

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Background

The integron is a bacterial recombination system for capturing and rearranging gene cassettes through site-specific recombination, which plays a role in the rise of antibiotic multiresistance in Gram-negative bacteria. The distinctive property of integron recombination sites flanking the cassettes (attC) is that they have almost no sequence conservation. Instead, their single-stranded DNA possesses conserved structural features, and attC sites are recombined as a folded form of the bottom DNA strand. Contrary to the bottom strand, the recombination of the top strand would integrate the cassette in the opposite orientation, preventing the expression of its gene from the promoter provided by the integron platform.

Objectives
In order to understand the basis of this specificity, we tested the recombination of several attC sites, and selected 3 of them for further analysis.

**Methods**
For these sites, we mutated their structural features and quantified the changes in frequency and orientation of cassette integration for the corresponding bottom and top strands.

**Conclusions**

We observed that some of the structural features, such as the extrahelical bases (EHBs), play similar roles for all of the tested sites, but the extent to which they influence the frequency and the orientation of cassette integration differs among attC sites. Yet other structural features, such as the unpaired central spacer (UCS), have very different, sometimes even opposing effects depending on the attC site. Our study suggests that the structural requirements for attC site recognition are rather relaxed, and that the control of cassette integration might be governed by more complex mechanisms than previously assumed.
POSSIBLE ROLE OF A MOBILE GENETIC ELEMENT IN TRANSLATIONAL MODULATION

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Background
A consortium of microorganisms adapted to grow in acidic environments participate in the bioleaching of minerals. The facultative aerobic bacterium Acidithiobacillus ferrooxidans is a Gram negative, chemolithothrophic, autotroph that utilizes either Fe²⁺ or sulfur reduced compounds as electron donors.

Objectives
We are interested in the role of non-coding RNAs (sRNAs and tRNAs) and RNAses encoded in a mobile genetic element in translation control.

Methods

By means of bioinformatic analyses 37 genes encoding tRNAs were detected in the mobile element ICEAfe1 from A. ferrooxidans strain ATCC 23270. Encoded tRNAs are aminoacylated in vivo in A. ferrooxidans, although they are transcribed at low levels.

Two strategies were used for their identification of sRNAs: 1. Bioinformatic prediction of transcription units in intergenic regions (IGR) of A. ferrooxidans genome and comparison to RNA deep sequencing data. 2. The identification of sRNAs that bind to the RNA chaperone Hfq were identified by RNA deep sequencing. Currently we are working in the validation of target mRNAs of the identified sRNAs. At least two snRNAs encoded in the ICEAfe1 were validated.

Four type II toxin-antitoxins systems specific for strain ATCC 23270 were identified in the in the ICEAfe1. All four toxins encoded an RNAs that is counteracted by the
antitoxin

Conclusions
Different genes encoding possible factors that modulate translation have been identified in the integrative conjugative element ICEAfe1 from *Acidithiobacillus ferrooxidans*. Whether these systems effectively modulate translation in this organisms is currently under investigation.
A NEW SHUTTLE-EXPRESSION SYSTEM TO GENERATE POLYHISTIDINE-TAGGED FUSION PROTEINS IN STAPHYLOCOCCUS AUREUS AND ESCHERICHIA COLI

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Background

A better understanding of the metabolism of S. aureus is needed. However, only few tools are available for production of tagged fusion proteins.

Objectives

To develop S. aureus-E. coli shuttle vectors for the expression of target genes and rgs-his₆ codon fusions.

Methods

The strong constitutive promoter of S. aureus type 1 capsule biosynthetic gene 1A (P-cap) upstream of a multiple cloning site (MCS) and codons for rgs-his₆ were synthesized and inserted into pBUS1 background. The MCS contained a BcgI site to generate C-terminal RGS-His₆ translational fusions of cloned genes through a simple cut-religation method. Using PCR-based techniques, two vectors were constructed that contain the P-cap-MCS-rgs-his₆ sequence (pBUS1-P-cap-HC and pTSSCm-P-cap) and two promoter-less variants suitable for cloning genes including their own promoter (pBUS1-HC and pTSSCm). The plasmids contained the E. coli origin ColE1, the terminator sequence rrmB(T1)₅ and the tetracycline resistance marker tet(L) for S. aureus and E. coli. Gram-positive replicon was improved through either complementation of the minimum pAMα1 replicon from pBUS1 with the single strand origin oriL from pUB110 or substitution with a pT181-family replicon. Plasmid stability, copy numbers and recombinant protein synthesis was analyzed in S. aureus and E. coli.

Conclusions

The new shuttle vectors displayed increased copy numbers and segregational stability in S. aureus. Feasibility of rgs-his₆ codon-fusion, gene expression and protein purification were demonstrated in S. aureus and E. coli using the macrolide, lincosamide, streptogramin B resistance gene erm(44). The new His-tag expression
system represents a helpful tool for the direct analysis of target gene function in staphylococci.
THE PRESENCE OF TEMPERATE BACTERIOPHAGES IN UROPATHOGENIC ESCHERICHIA COLI STRAINS

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Background

Horizontal gene transfer vectors, including bacteriophages, promote the spread of a broad spectrum of genes, among them some encoding medically relevant traits such as virulence factors. In Escherichia coli (E. coli), temperate bacteriophages have been shown to promote the spread of shiga toxin genes across shigatoxigenic strains of the species¹.

Objectives

Here we aim to assess the presence of lysogenic bacteriophages in uropathogenic E. coli (UPEC) strains, and to investigate a possible correlation with the virulence factor profiles of the examined strains.

Methods

Twenty UPEC strains were screened for the presence of 15 different virulence-associated traits (fimH, papC, papGII, sfaDE, focIc, afaBC, draBC, hlyA, cnf1, usp, ibeA, iucD, iroCD, iroN, fyuA and ireA) by PCR amplification of target genes, and were subsequently assigned virulence scores by counting possession of these genes. Additionally, strains were screened for the presence of lysogenic bacteriophages by induction with UV light. Data were analysed using Student’s t-test.

Conclusions

We have found that out of the 20 screened strains, nine possessed UV-inducible lysogenic bacteriophages. Furthermore, data analysis has shown that the strains harbouring UV-inducible lysogenic bacteriophages also possessed a significantly higher virulence score than the strains lacking UV-inducible bacteriophages (p < 0.01). The results of this preliminary study imply that temperate bacteriophages might have a potential role in virulence factor transfer across UPEC strains.

Reference
Background
Transposable elements (TEs) are components of nearly all bacterial genomes. They promote structural changes in DNA, which may result in different phenotypes. We focused on the elements of the Tn3 family – the most abundant group of bacterial transposons.

Objectives
Our studies shed a new light on Tn3 family, which is much more diverse than previously thought. We showed that Tn3 elements comprise (i) a composite transposon generated by two copies of a non-composite cryptic transposon, as well as (ii) simple non-autonomous elements (TIMEs - Tn3-derived inverted-repeat miniature elements) and (iii) their derivatives. We have provided for the first time evidence that some of the non-autonomous transposons may contain, in their core regions, functional plasmid replication and stabilizations systems, which can be transferred among replicons co-residing in a bacterial cell. The transposition events may therefore significantly influence structures of bacterial plasmids leading to generation of molecules exhibiting novel properties. Bioinformatic analyses of the nucleotide sequences of several bacterial genomes revealed the presence of large elements originating from the Tn3-family non-composite transposons, which acquired additional genetic information, thus forming a kind of transposable genomic islands.

Methods
Standard methods of molecular biology were used.

Conclusions
The obtained results indicate the powerful role of the Tn3-family TEs in the mobilization to transposition of segments of genomic DNA. Such shuffling of genetic information, may significantly influence the structures of bacterial genomes. Huge diversity of the identified elements allows to draw conclusions concerning the high dynamics of evolution of this group of TEs.
Background

A composite transposon is a transposon consisting of two insertion sequences (ISs) flanking a DNA segment which often carries antibiotic resistance (AR) genes. If both IS elements are the same then it is possible that one copy of the IS element plus the intervening DNA segment can excise as a circular form. This circular molecule has been called an “Unconventional Circularizable Structure” (UCS) or “Translocatable Unit” (TU). IS1216 composite transposons have been shown to carry various AR genes. In addition UCSs have also been found containing IS1216.

Objectives

To screen for IS1216 composite transposons and IS1216 containing UCSs in human oral metagenomic DNA.

Methods

PCR primers were designed to amplify in the outward direction from IS1216. The IS1216 PCR products were then cloned and analysed.

As our PCR strategy would also detect UCSs, PCR primers were designed to amplify outward from the intervening DNA region to determine if each IS1216 amplicons was derived from a UCS circular structure.

Conclusions

We found four different IS1216 composite transposons and confirmed the presence of two novel UCSs in the oral metagenomic DNA. One of them contains an antiseptic resistance gene and one contains a universal stress response protein encoding gene.

This is the first report of a PCR strategy to amplify the DNA segment on the composite transposons in metagenomic DNA, and is also the first time that UCSs have been found in metagenomic DNA. This PCR strategy can be used to identify
novel AR genes associated with mobile genetic elements in metagenomes from any environment.
DETECTION AND CHARACTERISATION OF INTEGRON GENE CASSETTES IN HUMAN ORAL METAGENOMIC DNA.

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Background

An integron is a genetic element, which is able to capture gene cassettes and direct the expression of open reading frames (ORFs) contained within them. They are involved in the dissemination of antibiotic resistance genes (ARGs) in many pathogens. The oral cavity is a reservoir of ARGs; however, these have not been shown previously to be present on an integron cassette.

Objectives

To investigate the presence of integrons and associated gene cassettes in the human oral cavity by using a PCR-based approach.

Methods

Different primer sets were tested on oral metagenomic DNA obtained from both the UK and Bangladesh, including novel primers based on the Treponema denticola integrase and attC sequences, and previously published primers targeting various conserved regions on different integrons. The resulting metagenomic amplicon libraries were then sequenced and analysed.

Conclusions

A diverse array of ORFs presents on gene cassettes and associated with integron integrases have been identified. The cassettes contain ORFs predicted to confer antibiotic resistance and adaption to environmental stresses including a putative chloramphenicol acetyltransferase, bleomycin binding protein, multidrug transporter, cof-like hydrolase, competence and motility related proteins.

This is the first study confirming the presence of integron gene cassettes in oral metagenomic DNA. The predicted proteins are likely to carry out a multitude of functions however the function of the majority is as yet unknown. We hypothesise that the oral cavity may be a rich source of diverse integron cassettes because it is such a variable physicochemical and stressful environment for its bacterial inhabitants.
Background
We address the central role of a functionally related group of transposable elements called “HuH elements” in structuring and shaping bacterial genomes. We have identified and characterised a widespread class of IS (IS200/IS605 family) fundamentally different from classical TEs: they use obligatory single strand DNA intermediates and have ends with subterminal imperfect palindromes (IP) which are recognised and bound by their transposases. These transposases are members of a larger “HuH” endonuclease superfamily. The protein binds the subterminal IP located some distance from the cleavage sites. Remarkably, cleavage sites are not recognised directly by the protein but by short “guide” sequences 5’ to the IP foot. Recognition involves a network of canonical and non-canonical base interactions similar to those found in RNA structures.

Objectives
We have demonstrated the importance of the lagging strand template for activity of some members and our in silico genomic analysis suggests that all IS200/IS605 family members have evolved a mode of transposition that exploits ssDNA at the replication fork.

Methods
Some members of this group appear to have been domesticated to perform important roles in the prokaryotic cell: such as homing endonucleases in some group I prokaryotic introns and as enzymes responsible for proliferation of short intergenic multicopy palindromic regulatory sequences called REPs.

Conclusions
The results provide an overview of the role of “HuH” transposases and their associated mobile elements in shaping the prokaryotic genome.
FUNCTIONAL CHARACTERIZATION OF AN INOSITOL METABOLISM ENCODING GENE CLUSTER CONTAINED IN ICEEFM1 OF ENTEROCOCCUS FAECIUM

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Background
Whole genome sequencing revealed the presence of an integrative conjugative element, ICEEfm1 in a clinical Enterococcus faecium (Efm) isolate (E1504) with an insertion of a cluster of 10 genes putatively involved in inositol metabolism.

Objectives
Functional characterization of the inositol metabolism (iol) gene cluster.

Methods
A cre-lox recombination system was used to construct a markerless iolD deletion mutant in strain E1504. The phenotype was determined by growth on minimal medium with inositol as carbon source (M-ino). The genetic organization of the gene cluster was determined by operon and promoter mapping and expression was determined after growth in rich and minimal medium in the presence and absence of myo-inositol. An in vivo mouse gut-colonization model was used to compare colonization abilities of E1504-wildtype and iolD mutant.

Results. The iolD deletion resulted in abolished growth in M-ino. The gene cluster is organized as an operon of 10 genes with its own promoter. Expression was only observed in the presence of myo-inositol and absence of other carbon sources. No difference was observed in gut-colonization abilities between wild-type and mutant.

Conclusions
The 10 gene insertion in ICEEfm1 confers Efm the ability to grow when only inositol is available as carbon source. Due to the availability of additional other sugars the iol gene cluster did not provide a selective advantage in the in vivo mouse model. Possibly the gene cluster was acquired in an environment where carbon sources were limited and only inositol was present like e.g. plants and soil.
Background
Bacterial viruses (phages) are ubiquitous and have a strong impact on microbial population dynamics. Temperate phages may become prophages upon cell infection, producing lysogenic hosts and thus contributing to the evolution of bacterial gene repertoires.

Objectives
We identified and studied 2246 prophages among the genomes of 1196 bacterial species to pinpoint the genetic and life-history traits associated with lysogeny and how they shape the distribution of prophages among bacteria.

Methods
Surprisingly, whereas lysogens have larger genomes and are more often pathogens, they are not different in terms of cell volume or defense systems, like CRISPR-Cas. Importantly, growth-related trade-offs seem to be key to the decision of lysogeny. We propose that lysogeny is more adaptive when hosts show large variations in the resources they can provide. Accordingly, bacteria with low minimal doubling times, which are associated with strong variations in growth rates and cell volume, show remarkably high rates of lysogeny. This fits theory suggesting that lysogeny is a strategy of slow vertical reproduction in waiting for future more efficient horizontal reproduction. This strategy has important consequences for bacterial evolution: we find that lysogens have much more diverse gene repertoires, after controlling for other factors, showing the key role of lysogeny in bacterial adaptation.

Conclusions
Our work shows a complex dependency of bacterial genetic diversification on host-parasite interactions and the life-history traits of both bacteria and phage. It allows establishing coarse-grained predictions of the abundance and distribution of prophages and explains the paucity of prophages in certain bacterial clades.
CONTRIBUTION OF INSERTION SEQUENCE ELEMENTS TO THE GENOME PLASTICITY OF CUPRIAVIDUS METALLIDURANS AE126

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Background
Mobile genetic elements play a significant role in bacterial evolution. Insertion Sequence (IS) elements are of specific interest as they constitute an importing driving force for genome plasticity. *Cupriavidus metallidurans* AE126 (type strain CH34 cured of pMOL30) harbors 19 distinct IS elements from nine IS families, reaching a total of 52 intact IS copies. *C. metallidurans* AE126 displays in the presence of toxic zinc concentrations spontaneous mutations resulting in an increased zinc resistance.

Objectives
The objective of this study is: (i) to determine the contribution of IS elements to genome adaptation of AE126 after zinc stress and (ii) to examine the putative stress-specific inducible transposition of IS elements.

Methods
Zinc resistant AE126 derivatives were screened by PCR and mutations were identified by sequencing. Promoter transcription of IS elements was analyzed with the promoter probe vector pGLR1.

Conclusions
All derivatives had a base substitution/indel (45%) or an insertion of an IS element (55%) in the regulatory locus of *cnr* (cobalt-nickel resistance). Seven IS elements were identified to be transposed into the regulatory locus of *cnr*. Three elements were found more often with respect to their copy number and transcription of their promoters were induced by supplementing zinc or cadmium to the growth medium but not nickel or cobalt. Some AE126 derivatives, e.g. with an *ISRme5* element inserted in the same orientation as *cnrYXH*, are pre-adapted as they evolved to ameliorate resistance to other stress challenges. Pre-adaptation could putatively be explained by the increased transcription of the mutated *cnr*-operon and concomitant co-transcription of *ISRme5*. 
Objective To investigate the distribution and characteristics of vibrio seventh pandemic island-I (VSP-I) in Vibrio cholerae. Method All V. cholerae genome sequences in GenBank were downloaded and screened for VSP-I island using VSP-I (VC0175~VC0185) of strain N16961 as the reference sequence. MUMmer software was used to analyze single nucleotide polymorphism (SNP); OrthMCL software was used for homologous gene analysis; finally the BLAST software was used to search sequence variation and functional annotation. Conclusions VSP-I widely exists in the genomes of all the seventh epidemic V. cholerae O1 strains and its sequence is highly conserved; Among the 105 V. cholerae O1 strains, the VSP-I sequences of 94 strains were completely identical, the VSP-I sequences of 11 strains had 1~14 SNP loci, respectively. The VSP-I exists also in the genomes of two non-O1/O139 strains. However, their VSP-I sequences contain some insertion and deletion (INDEL) comparing to that of O1 strains. VSP-I had 9 conserved genes totally. Among them, VC0183 existed in all strains, but its length of the non-O1/O139 strains was different to the O1 strains. Five (VC0181~VC0185) of 9 genes were involved in its transfer and integration. And VC0178 encoding potato glycoprotein analogues and VC0180 encoding protein hydrolase may be related to virulence. As a specific genetic cluster within the seventh cholera pandemic strains, VSP-I has been transferred to the non-O1/O139 strains, and variation of its sequence has been occurred as well.
Background

The antimicrobial activity of silver is well documented, however its mechanism of action is not well understood. It is unclear whether the effects of silver nanoparticles (Ag-NPs) and Ag⁺ ions are similar or have different antimicrobial mechanisms.

Objectives

The objective was to compare the damages caused by Ag-NPs and silver nitrate (AgNO₃) on gram-positive and gram-negative bacteria by transmission electron microscopy.

Methods

Ag-NPs were synthesized by reduction of AgNO₃ with sodium citrate. *Pseudomonas aeruginosa* ATCC 15442 and *Staphylococcus aureus* ATCC 6538 were treated with Ag-NPs or AgNO₃ at 37 °C for 2 h. The cells were fixed with 2.5% glutaraldehyde for 24 h and post-fixed in 1% osmium tetroxide for 2 h at 25 °C, washed once in 0.1 M PBS and dehydrated by sequential transfer in a graded series of ethanol solutions. Fine sections were cut with diamond blades using ultramicrometer and double stained with uranyl acetate and lead citrate saturated and observed using the transmission electron microscope with EDX detector (FEI Tecnai-G2-20, 2006) available in microscopy center of Federal University of Minas Gerais.

Conclusions

The Ag-NPs affected severely *P. aeruginosa*, forming a condensation region in the center of the cell and a clear zone around. According to the EDX spectrum, the condensation region is composed mainly of silver, sulfur and phosphorus. However, these elements were not detected in the clear area. Damages to *S. aureus* were similar to those caused on *P. aeruginosa*, however, milder form. Damages caused by AgNO₃ were similar to those caused by Ag-NPs.
LIPID PEROXIDATION IS NOT RELATED TO THE ANTIMICROBIAL MECHANISM OF SILVER NANOPARTICLES AGAINST PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS

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Background

The antimicrobial activity of silver nanoparticles (Ag-NPs) may be related to the formation of reactive oxygen species (ROS). A common feature among the different ROS is the ability to cause some oxidative damage in cells. The lipids are the main targets during oxidative stress.

Objectives

The objective was to determine whether the Ag-NPs cause oxidative stress in the lipids of membranes of Pseudomonas aeruginosa or Staphylococcus aureus.

Methods

The Ag-NPs were synthesized according Monteiro et al. (2009), with some modifications. The lipid peroxidation analysis was performed according to Becerra et al. (2006), with some adjustments. Samples containing 50 µg/ml Ag-NPs or AgNO₃ were inoculated with 1.0 x 10⁸ CFU/ml of P. aeruginosa or S. aureus and incubated at 37°C for 1 h. Then was added 1 ml of 35% trichloroacetic acid (w/v). After 20 min, the samples were treated with 1 ml of 0.5% thiobarbituric acid (w/v) and heated at 80°C for 30 min, then cooled and centrifuged. The absorbance was determined at 535 nm. The controls with water and 2,2'-azobios (2-methyl-propionamide) dihydrochloride (AAPH) were made. A standard curve of malondialdehyde was obtained by hydrolysis of 1′1′3′3′-tetraetoxipropano.

Conclusions

The concentration of malonaldehyde, a product of lipid peroxidation, in P. aeruginosa and S. aureus cells treated with Ag-NPs and AgNO₃ was not different (p > 0.05) from cells treated with water, however, was lower from the cells treated with AAPH, a ROS generator. This indicates that lipid peroxidation is not involved in the antimicrobial activity of Ag-NPC and AgNO₃.
FEMS-2156
New antimicrobial mechanisms

ASSESSMENT OF E. COLI BIOFILM GROWTH ON DIFFERENTLY TERMINATED DIAMOND THIN FILMS
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Background
Diamond is a material with interesting properties, such as chemical inertness, hardness, high transparency and good biocompatibility with eukaryotic cells. These qualities make this material very promising for various bio-medical applications such as biosensors, articular implant coatings, tissue-engineering and regenerative medicine. Several published works also indicate that diamond thin films, due to their tailored surface properties, could possess anti-adhesive or even antimicrobial properties.

Objectives
The objectives of our work is to develop and standardize the methods for diamond thin film deposition on glass substrate, to develop and optimize the methods for testing of antibacterial properties of diamond surfaces, to assess these properties and to compare the influence of diamond surface terminations on their possible antibacterial character.

Methods
We used diamond films CVD depositeed by microwave plasma CVD process. Escherichia coli was chosen as the model bacterium. Bacterial biofilm was cultivated on diamond-coated glass in CDC biofilm reactor. Fluorescence microscopy and ATP assay was used to quantify bacterial growth on diamond surfaces.

Conclusions
We managed to deposit diamond thin films of standard properties on glass over large areas (1x3”). We successfully developed the continuous cultivation method for biofilm growth and testing of antibacterial properties of diamond thin films under different conditions. We compared the antibacterial potential of diamond films with different surface terminations (H, O and F).
This work was supported by funding from Czech Science Foundation (GACR 15-01687S).
FEMS-1662
New antimicrobial mechanisms

SUSCEPTIBILITY OF FOOD CONTAMINATING MICROBIAL COMMUNITIES TO ATMOSPHERIC COLD PLASMA AND IMPLICATIONS FOR RESISTANCE-PROOF ANTI-MICROBIAL TREATMENT

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Background

Atmospheric cold plasma (ACP) has received increasing attention as a novel anti-microbial treatment, which may be less prone to resistances. Containing ions, free electrons, UV light and reactive species, plasma can cause damage to microbial cell walls and/or intra-cellular components.

Objectives

To determine the anti-microbial efficacy of ACP on food-contaminating microorganisms and their ability to develop resistance to plasma treatment.

Methods

The inactivation of a range of common microbial food contaminants using di-electric barrier discharge ACP was investigated and cultures of S.aureus and E.coli were subjected to repeated sub-lethal plasma treatment. The background microflora of plasma-treated meat products was analysed over 2-4 week storage and community profiling was performed using PCR-amplification and denaturing gradient gel electrophoresis (DGGE) of 16s RNA.

Conclusions

Microorganisms displayed varying susceptibility to ACP, with C.jejuni and B.thermosphacta showing rapid inactivation whereas E.coli, L.monocytogenes and S.aureus required longer treatment times. Repeated selection of sub-lethally treated colonies did not increase resistance in short-term adaptation studies. The background microflora of meat products was reduced up to 1log over 2-4 week storage following treatment and PCR-DGGE revealed differences in plasma tolerance between microbial populations. In an era of growing anti-microbial resistances, resistant-proof alternative treatments are gaining significance. ACP has shown anti-microbial effects on a variety of microorganisms and acts through a range of physico-chemical effects that may prevent the development of resistances. Such novel anti-microbial strategies have implications not only in the food industry but also the medical sector but will require long-term adaptation studies to ensure resistance-proof.
Background
Healthcare textiles are known to be able to transfer pathogens.

Objectives
To provide new antimicrobial textiles, we functionalized decontaminating photocatalytic textiles by the innovative LBL (Layer-by-Layer) technique. Photocatalysis deals with the activation of TiO$_2$ semiconductor particles by UVA, which produces highly Reactive Oxygen Species responsible for the microorganisms inactivation.

Methods
The LBL coating is based on successive application of positively charged layers and negatively charged layers. To improve the adhesion of the first layer on the fibers, the textile was first submitted to a hydrolysis using KOH solution. Negatively-charged TiO$_2$ particles with a polyethylenimine (PEI) polycation layer were alternatively spraying onto textiles. Another functionalization was also performed by adding an interfacial layer of PEI before building bilayers of alternatively positively-charged TiO$_2$ particles with a poly(styrene sulfonate) (PSS) polyanion. These sequence were repeated 1-5-10 times until (PEI/TiO$_2$)$_n$ or PEI/(PSS/TiO$_2$)$_n$ bilayers were achieved. The antibacterial properties of functionalized textiles against *E.coli* were determined after 15-30-60min of solar light radiation (226W/m$^2$ of visible light).

Conclusions
Whereas no antibacterial effect was observed for the textiles pretreated by KOH and kept in the dark, we respectively obtained a 2log reduction for the textiles functionalized with only one layer of PEI and a 6log reduction with addition of a single layer of TiO$_2$ (PEI/TiO$_2$)$_1$. For the PEI/(PSS/TiO$_2$)$_n$ textiles, antibacterial activity is function to solar light duration and also depends of the number of bilayers. The best antibacterial activity (4log reductions) was obtained for PEI/(PSS/TiO$_2$)$_{10}$ under 60min of solar light radiation.
Background

Free nitrous acid (FNA) has recently been demonstrated as an antimicrobial agent to a range of microorganisms. However, the antimicrobial mechanism of FNA is largely unknown.

Objectives

Consequently, we aim to gain a systematic understanding of the bacteriostatic/bactericidal effects of FNA. This was examined on Pseudomonas aeruginosa PAO1, an organism with intrinsic resistance to various antibiotics and disinfectants.

Methods

P. aeruginosa was grown under anaerobic denitrifying conditions and when FNA was added (0.05 mg N/L) growth temporarily stopped. From cultures either exposed or not exposed to added FNA, RNA was extracted and transcripts were detected by HiSeq-Illumina sequencing. Transcripts were quantified and those exhibiting ≥2.5 or ≤-2.5 fold abundance change were designated as differentially expressed in response to FNA.

Conclusions

In response to FNA 177 genes showed increased transcription while 471 genes exhibited deceased transcript levels. Genes coding ribosome proteins showed significantly decreased transcript levels, implicating that protein biosynthesis was severely inhibited by FNA. Respiration was likely inhibited, as lowered substrate utilization coincided with decreased transcript levels of genes coding for denitrification. In contrast, genes encoding nitric oxide reductase were up-regulated, which possibly functioned to remove toxic NO derived from the added nitrite. In the response, genes of the tricarboxylic acid cycle were less expressed, while enzymes of the glyoxylate shunt had increased levels. Evidently the cells would generate less reducing power through these cycles. This first investigation elucidating the
mechanisms of FNA effect on *P. aeruginosa*, contributes underpinning knowledge for potential application of this novel antimicrobial agent.
Background

*Pseudomonas aeruginosa* is a gram-negative pathogen that causes severe infections in immunocompromised patients and is responsible for 10 – 15% of the hospital-acquired infections worldwide. Due to the emergence of multidrug-resistant *P. aeruginosa* strains, there is an urgent need for the development of new antibiotics with activity against *P. aeruginosa*. We recently identified a new antibacterial compound, SPI003, which inhibits growth of various gram-positive and gram-negative pathogens, including *P. aeruginosa*.

Objectives

In this study, we sought to unravel the mode of action of SPI003, using *P. aeruginosa* as a model organism.

Methods

In order to unravel the mode of action of SPI003, different approaches were used. First, a library of *P. aeruginosa* mutants was screened to identify mutants that are sensitive or resistant to SPI003. Second, spontaneous resistant mutants were generated by plating *P. aeruginosa* cells on agar plates supplemented with SPI003 at a final concentration of 5 x MIC. Third, a transcriptome analysis was carried out on exponentially-growing cells of *P. aeruginosa* treated with the compound. Finally, the membrane permeability of *P. aeruginosa* was examined after exposure to SPI003 using established assays.

Conclusions

Our study shows that SPI003 exhibits a fast membrane-damaging activity, which is likely to be the primary cause of its antibacterial activity.
INVESTIGATION OF ATMOSPHERIC COLD PLASMA AGAINST ESCHERICHIA COLI MUTANTS

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Background

Atmospheric cold plasma (ACP) is effective against a wide range of pathogenic microorganisms with promising application in food and medical industry. Generation of reactive oxygen species (ROS) plays a crucial role when air/oxygen containing gases are used. As a mixed group of oxidative reactive species generated by ACP, there are limited publications about the cellular regulation system under plasma pressure.

Objectives

To investigate the effect of the plasma stress on the microbial inactivation, intracellular reactive oxygen species and genetic regulation.

Methods

*Escherichia coli* BW 25113 and its isogenic mutants in *soxR*, *soxS*, *oxyR*, *rpoS* and *dnaK* genes were suspended in phosphate buffered saline (PBS) at $10^8$ CFU/ml. Samples were then treated with high voltage plasma (80 kVRMS) in a sealed package for 1, 3 and 5 min following 0, 1 and 24 h post-treatment storage. Intracellular ROS were measured using fluorescence spectrophotometer (Biotek synergy, 480/530nm) with the probe 2', 7'- dichlorofluorescein diacetate immediately after storage.

Conclusions

Without post-treatment storage time, the absence of RpoS and OxyR led to higher reduction after treatment. However, DnaK showed its repairing function correlated to cell survival with 24 h, while it has no effect on reducing intracellular ROS level. SoxS became important with both 1 and 24 h storage, which is not observed in SoxR. It implied their different characteristics as subunits. Overall, the cell response against plasma generated oxidative stress could be divided to short term and long term, which are dominated by oxidative stress deletion genes and damage repairing genes.
NEW ENVIRONMENTAL-FRIENDLY COMPOUNDS TO COMBAT CITRUS CANKER

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Background
The culture of citrus fruits is constantly under the threat of plant diseases. One of them is citrus canker, which is caused by the bacterial plant pathogen Xanthomonas citri subsp. citri (Xac). Current modes of prevention of Xac infection and spreading include spraying trees with copper containing bactericides and eradication of infected and healthy trees in a certain radius around. Despite these efforts, about 300 Million tons of fruit per year were lost during the last decade.

Objectives
In addition to being inefficient, copper containing bactericides negatively impact soil quality since copper accumulates in the environment. Therefore, more sustainable ways to combat citrus canker are urgently needed. Recently, a class of plant-derived semi-synthetic compounds, the alkyl gallates, has shown promising activity against Xac [1] and the bacterial cell division protein FtsZ has been found to be their target (unpublished).

Methods
Here, we present in vitro results on the activity of the alkyl gallates against FtsZ from different bacteria. We also discuss the ideal alkyl chain length of a lead gallate candidate as a platform for chemical synthesis of new compounds with enhanced antibacterial activity.

Conclusions
Our initial results show that the development of novel environmental-friendly antibacterials to combat Xac is feasible. The efficacy of these compounds in the field will be evaluated and their toxicity to plants and animals will be determined. We expect that the compounds can be used as a sustainable alternative to prevent citrus canker and they will contribute to control the spread of Xac in citrus plants.
Background
Rapid development of antibiotic resistance of bacteria necessitates the search for new natural compounds inhibiting the growth and functioning of pathogenic microflora.

Objectives
To study the antibacterial mechanisms of warnerin, a new peptide from lantibiotic family that demonstrate the sequence of events causing the death of attacked bacterial cells.

Methods
Outcomes of warnerin attack of Staphylococcus epidermidis 33 GISK (Moscow) were analyzed using the combination of different methods such as polarography (respiration), spectrography (intracellular potassium loss), fluorescence assay of membrane potential dynamics and hydroxyl radicals' formation, electron and atomic-force microscopy of cell structure damages

Conclusions
Peptide introduction into the bacterial incubation medium results in the increase in their oxygen consumption followed by rapid cellular release of potassium ions and sharp decline of intracellular ATP concentration. Apparently, the disturbance of ATP-linked electron transport along the respiratory chain favors their accumulation on iron-sulfur centres. This results in increase of free Fe$^{2+}$ ions and as a consequence, to the elevation of hydroxyl radical concentrations. Peptide inhibition of ATP formation is accompanied by marked proton accumulation on the outer side of bacterial membrane, particularly in its adjacent layers of peptidoglycane. This facilitates the autolytic hydrolase extrusion from the binding sites and to uncontrolled total bacteriolysis that is revealed by microscopic data. Results of investigation evidence for the pluripotent antibacterial action of warnerin that point to expressed prospects for its practical use.

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FUNCTIONALIZATION OF PROTOPORPHYRIN IX WITH ANTIMICROBIAL PEPTIDES TO ENHANCE PHOTOANTIMICROBIAL CHEMOTHERAPY (PACT).


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Background
Photodynamic antimicrobial chemotherapy (PACT) combines activities of light, photosensitizing compound (PS) and oxygen, to eradicate microorganisms through the oxidation-related mechanisms. To obtain more specific photodynamic action, PS can be functionalized with various types of adducts. Presented approach combines activity of a photosensitizer and antimicrobial peptides (AMPs).

Objectives
Protoporphyrin IX (PPIX) covalently conjugated with antimicrobial peptides: PEX or CAM, was analyzed for its spectral properties and antimicrobial action. Bacterial strains used in the study were *Staphylococcus aureus* Newman (MSSA) and COL (MRSA).

Methods
Peptides were synthesized using the methodology Fmoc/tBu on polystyrene resin and purified by HPLC. MIC values were determined by the broth dilution method in BHI medium in the darkness. Absorbance spectra screening was performed using the EnVision™ microplate reader (PerkinElmer) within 350-700 nm range. PACT was conducted using the LED lamp (SecureMedia, Poland) emitting incoherent red light ($\lambda_{max}$ 627 nm). Viability of bacteria was calculated counting colony forming units (CFU).

Conclusions
Combination of pure compounds showed no influence on PPIX spectral properties and AMPs activity by means of MIC values and showed synergistic effect of PPIX and CAM in PACT experiments. PS-AMP conjugates reveal weaker antimicrobial activity by means of MIC values, as compared to pure peptides. Conjugation of peptides to PPIX changed its spectral properties, visualized by hypsochromic shift of absorbance spectra, hyperchromic effect in Soret band and hypochromic effect in Q bands. These observation implied weaker/no photodynamic activity induced by red light, which was confirmed experimentally, however indicated efficient activity induced by blue light.
SILVER NANOPARTICLES INHIBIT MOTILITY, PYOVERDINE PRODUCTION AND BIOFILM FORMATION BY PSEUDOMONAS AERUGINOSA

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Background
Antimicrobial efficacy of silver nanoparticles (AgNPs) was proved and broad applications were proposed so far. This study shows the potential mechanism of AgNPs action towards opportunistic pathogen Pseudomonas aeruginosa.

Objectives
Study of the AgNPs effect on selected P. aeruginosa virulence factors.

Methods

P. aeruginosa PAO1 was used in this study. Silver nanoparticles were obtained from Nano-Tech (Warsaw, Poland). Minimal inhibitory concentration of silver nanoparticles that inhibit biofilm formation (MBIC) was estimated by standard two-fold dilution method. Swimming motility was examined on 0.3% agar plates and swarming motility on 0.6% agar plates supplemented with 0.05–0.5xMBIC AgNPs. Twitching motility across the glass surface was analysed after staining with crystal violet. Pyoverdine production was estimated in 48-hour PAO1 biofilms grown in microtitre plates. Biofilm formation was studied after crystal violet staining and cell survival was determined in Live/Dead Assay (Promega). Statistical analysis of the results in comparison to control sample without AgNPs was performed.

Conclusions

Strong antibacterial effect of AgNPs was observed. MBIC was 4 microgram mL⁻¹. There was significant decrease from 22% to 40% in swimming and from 22% to 29% in swarming motility in samples containing 0.1–0.5xMBIC of AgNPs in comparison to P. aeruginosa PAO1 incubated without nanoparticles. No influence of AgNPs on twitching motility was observed. Pyoverdine production was significantly inhibited by AgNPs in concentration of 0.5 MBIC. The inhibition of biofilm formation by 44% was also observed. Hence, it was shown that silver nanoparticles affect several virulence factors of P. aeruginosa including swarming, swimming motility, pyoverdine production and biofilm formation.
ANTIBIOTIC ACTIVITY OF MICROBIAL STRAINS ISOLATED FROM MYTILUS EDULIS, KIEL FJORD, BALTIC SEA

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Background

Microorganisms associated with eukaryotic hosts are involved in various physiological functions, including chemical defense and immune responses. Production of bioactive molecules such as antibiotics and antifouling substances by associated bacteria, is one of the mechanisms to protect the host against pathogens, predation and surface colonization.

Objectives

In this study, extracts from 10 bacterial strains isolated from the blue mussel, Mytilus edulis (Kiel Fjord, Baltic Sea, Germany), were screened for their antimicrobial activity against two panels of test microorganisms: a standard panel (Escherichia coli, Bacillus subtilis and Candida albicans) and environmental panel composed of strains isolated from the mussels (Pseudomonas veronii, Bacillus pumilus and Shewanella baltica).

Methods

To identify isolated strains, PCR amplification of 16S rRNA was carried out and bioactivity tests were modified according to Schneemann et al. (2010) to evaluate the interaction of isolated strains with the test microorganisms.

Conclusions

Inhibiting effects varied considerably among the test microorganisms, with P. veronii showing the most pronounced antibacterial activity against both the standard and the environmental panels. Moreover, the Gram-positive bacteria, and particularly B. pumilus, were generally more susceptible than Gram-negative strains, whereas no strain was found to inhibit growth of E. coli. These findings suggest that bacteria associated with the blue mussels may be involved in preventing pathogen invasions and that these hosts represent a promising source for the isolation of
antibiotic-active bacteria.
FEMS-1088
New antimicrobial mechanisms

CHALCONE AND ITS HYDROXYLATED ANALOGUES: SYNTHESIS, ANTI-
CRYPTOCOCCUS AND CYTOTOXIC ACTIVITIES
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Background Cryptococcosis is a potentially fatal systemic mycosis, which is caused by Cryptococcus neoformans (Cn) and C. gattii (Cg). This disease is an opportunistic infection for AIDS. Cryptococcus is able to cause three types of infections, cutaneous cryptococcosis, pulmonary cryptococcosis and cryptococcal meningitis. The treatment recommended is the intravenous amphotericin B with flucytosine administered orally. This drug combination causes severe side effects and yeast-resistance, encouraging the search for innovative anti-Cryptococcus agents. Objectives The objectives of this work were synthesis of chalcone and hydroxychalcones, determination of antifungal activity against Cn and Cg strains, evaluation of cytotoxicity against mammalian cell lines, and evaluation of the inhibitory cell adhesion activity of Cn onto pneumocytes. Methods The chalcone and hydroxychalcones were synthesized by Claisen-Schmidt condensations. The anti-Cryptococcus activity was evaluated by microdilution test performed in 96-well plates, using CLSI-M27-A2 document. This assay demonstrated the values of Minimum Inhibitory Concentration (MIC) and Minimum Fungicidal Concentration (MFC) to five strains; Cn-ATCC90012, and four clinical isolates CnS, CnI, CnR and Cg118. The cytotoxicity was evaluated by MTT colorimetric assay, using five mammalian cell lines, A549, HepG2, MRC-5, U87-MG and NOK. Conclusions The hydroxychalcones exhibited potent antifungal activity, demonstrating MIC and MFC values ranging from 1.95 to 15.6µg/mL, which were related to high inhibition of Cn adhesion onto pneumocytes. In MTT assay, at 100µg/mL, these compounds were not toxic for all cell lines. We conclude that hydroxychalcones are antifungal hits, with relative selectivity for yeast, and their mode of action is related to inhibition of yeast adhesion onto host-cells.
FEMS-1859
New antimicrobial mechanisms

IN VIVO CLUSTER FORMATION OF NISIN AND LIPID II IS CORRELATED WITH MEMBRANE DEPOLARIZATION
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Background

Nisin and related lantibiotics can kill bacteria in two ways: by pore formation, which leads to membrane depolarization, or by sequestering Lipid II, which interferes with cell wall synthesis. Literature shows that during sequestration, nisin and Lipid II cluster into non-physiological domains, which are visible with fluorescence microscopy. Recently we observed that a reported ‘non-pore’ forming nisin mutant clusters LipidII, but also depolarizes the membrane (Lages et al., 2013, Environm. Microbiol. 15, 3272).

Objectives
To further investigate the relation between LipidII clustering and membrane depolarization.

Methods
Microscopy, fluorescent assays to detect membrane integrity.

Conclusions
We used in vivo assays to investigate how and when nisin forms pores, depolarizes the membrane, and segregates Lipid II into domains. By comparing the activity of pore-forming and pore-deficient mutants of nisin we show that clustering of nisin and Lipid II only occurs when (i) membrane integrity is reduced by pore formation and (ii) when the membrane is completely depolarized. Furthermore, pore-forming variants of nisin were also able to kill L-forms – which grow without a cell wall – whereas non-pore forming lantibiotics only kill bacteria with a cell wall. Our results have important implications for the mechanism-of-action of nisin. Because cluster formation never occurred with non-pore-forming lantibiotics, we suggest that lantibiotics either kill by pore-formation, or by occlusion of Lipid II for incorporation into the cell wall.
Background

Combinatory effect between conventional antimicrobial drugs and phytochemical products can be a new strategy to increase effectiveness of antibiotics.

Objectives

We aimed to verify synergism between essential oils compounds from plants and antimicrobial drugs for methicillin-resistant Staphylococcus aureus (MRSA).

Methods

Five essential oils compounds (citronellol, geraniol, eugenol, terpineol and cinnamadheyde) were tested against 10 strains of S. aureus MRSA. Mueller-Hinton agar (MHA) plus 0.5% Tween 80 was mixed with each compound (¼ of Minimal Inhibitory Concentration (MIC90%)). Nine antimicrobial drugs discs (oxacillin (1 µg), gentamicin (10 µg), erythromycin (15 µg), sulfazotrim (25 µg), vancomycin (30 µg), penicillin G (10 U) levofloxacin (5 µg), tetracycline (30 µg) and linezolid (30 µg)) were laid on culture medium inoculated with S. aureus. Assays were performed in duplicate. Mann-Whitney test was used as statistical. Time kill curves were performed between EOC and AD for positive cases. Treatments were incubated with 10⁵ CFU/mL of MRSA strain in MH broth. Media without EOC and/or AD was used as control. Aliquots were removed at different time intervals between 0 and 24h. Viable colonies were counted after incubation (24 h/ 37ºC) in order to obtain CFU/mL.

Conclusions

The present study showed a promising way of treatment for diseases caused by MRSA as well as against other bacterial species, because time kill curves for EOC an AD demonstrated additive activity. Thereby, these compounds can be used in combination with conventional antimicrobials.
SENSITIVITY OF MDR ACINETOBACTER BAUMANNII WOUND ISOLATES TO RUMEX SANGUINEUS, RUMEX CRISPUS AND URTICA DIOICA EXTRACTS

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Background
Acinetobacter baumannii is a pleomorphic, non-motile bacterium and etiological agent of various infections in immunocompromised patients. It has been characterized as a novel, drug resistant and rapidly emerging clinical pathogen with increasing prevalence. Regarding recent multi-drug resistant A. baumannii strains emergence, it is urgent to examine alternative antimicrobial agents, including various plant extracts.

Objectives
The aim of this study was to determine in vitro antimicrobial activity of Rumex crispus, Rumex sanguineus and Urtica dioica extracts against multi-drug resistant A. baumannii wound isolates.

Methods
The broth microdilution assay was used to determine minimal inhibitory concentrations (MIC) and minimal bactericidal concentrations (MBC) of 8 plant extracts obtained from different parts of the examined plants. For antibacterial activity testing 20 multiple resistant A. baumannii wound isolates and 2 reference strains (ATCC BAA-747 and ATCC 19606) were used.

Conclusions
Urtica dioica possessed no effect against A. baumannii, while Rumex sanguineus and Rumex crispus extracts showed significant antibacterial activity against all examined isolates. Bacteriostatic effect was obtained with concentrations 1 - 4 mg mL⁻¹ with median value 2 mg mL⁻¹, while bactericidal activity was achieved with a slightly higher concentrations of Rumex extracts (1 to 8 mg mL⁻¹ with median 4 mg mL⁻¹). The obtained result confirmed earlier detected antibacterial effect of R. crispus extracts against A. baumannii, while anti-A. baumannii activity of R. sanguineus extracts was detected for the first time. Accordingly, the plants from the genus Rumex are valuable source of bioactive compounds against this bacterium and antimicrobial activity of dominant extract components should be further examined.
CHARACTERIZING THE STAPHYLOCCUS AUREUS ANTI-VIRULENCE CANDIDATE SOLONAMIDE B AND IT’S LACTAM ANALOGUES

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Background
Staphylococcus aureus infections are becoming increasingly difficult to treat due to antibiotic resistance. Of particular concern are the community-associated methicillin-resistant S. aureus (CA-MRSA) strains such as USA300. Virulence of CA-MRSA is regulated by the agr quorum-sensing system, which in response to agr-encoded autoinducing peptides confines exotoxin production to the stationary growth phase with concomitant repression of surface-expressed adhesins. In need of novel treatment options, anti-virulence therapy, targeting virulence factors rather than viability, has been proposed. Solonamide B was recently isolated from a marine bacterium as a putative anti-virulence compound that markedly reduced expression of the toxins α-hemolysin and the phenol-soluble modulins.

Objectives
To further strengthen the anti-virulence candidacy of Solonamide B and synthesized analogues.

Methods
Chemically synthesized solonamides and lactam analogues were validated for interference with agr virulence gene regulation, both quantitatively and qualitatively, using reporter fusion strains. The compounds’ influence on biofilm formation and fibronectin binding of S. aureus were assessed as examples of complications related to agr negative strains; and lastly, toxicity of the compounds and the immune modulating capacity of solonamide-treated S. aureus were assessed on PBMCs, T-cell populations and DCs via flow cytometry.

Conclusions
We find that structural differences between solonamide analogues interfere differentially with agr, and that they do not display toxicity or influence immune cell activity and integrity. Importantly, our results also dismiss the major concern that application of compounds mimicking an agr-negative state may lead to adverse
interactions with host factors and immune cells.
Background

Preparation of liposome surface modified by various materials such as RNA, DNA or Proteins has been developed for several decades.

Objectives

We would like to investigate targeting ability of lysosomal membrane proteins related to killing bacteria through modifying the liposome surface to check the possibility of as antimicrobial agents whereby we tried to find specific membrane proteins.

Methods

Firstly, we used mixtures of lipid, cholesterol and lysosomal membrane protein isolated from lysosomes extracted from S. cerevisiae treated by H₂O₂ and NH₄Cl to modify liposome surface for antimicrobial activity after making liposomes prepared with hydration method. Liposomes were modified by lysosomal membrane proteins with two different methods in order to confirm the targeting ability.

Conclusions

The results showed that the antimicrobial activity was not shown without modification of liposome surface but it was better cell mortality in type 1 than type 2 method. Based on these results, we analyzed whole lysosomal membrane proteins by 2-DE assay to find specific proteins associated with antimicrobial activity and constructed recombinant S. cerevisiae. And then, we found the genes (HSP70) related to antimicrobial activity and prepared liposomes modified by lysosomal membrane proteins extracted from lysosomes isolated from recombinant S. cerevisiae tagged with GFP. The result showed that the antimicrobial activity of liposomes modified with Type1 method was better than control. Therefore, lysosomal membrane proteins for targeting bacteria will play an important a role.

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DOUBLE LEADING: LANTIBIOTIC - MICROCIN HYBRIDS
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Background
Lantibiotics and microcins form different groups of posttranslationally modified bacterial peptides mainly produced by Gram-positive and Gram-negative bacteria, respectively. Most of these peptides exhibit a potent antimicrobial activity, more active even than some antibiotics. Maturation of a peptide begins with a propeptide which is guided throughout specific modification events due to leader peptide (an amino acid sequence recognized by particular modification biomodule). The leader peptide is proteolitically removed in the last steps of peptide modification making the modified peptide active. Posttranslational modifications endorse lantibiotics and microcins with high target-specific activities and stability against proteolysis.

Objectives
To design plug-and-play biomodules for production of novel peptides containing both, lantibiotic- and microcin-specific posttranslational modifications.

Methods
We used a synthetic biology approach to design biomodules. We utilized peptide modification systems cloned under different controlled promoters to assure gradual and temporal activity of designed biomodules. Codon optimized synthetic genes of designed model peptides to be modified were employed together with modification machineries.

Conclusions
This is a unique case where posttranslational modifications from two different peptide classes are fused together into a single peptide chain in vivo. Here, we also show the possibilities of employing chimeric leader peptides for substrate recognition and procession by particular posttranslational modification machineries.
SILVER NANOPARTICLES WITH ANTIMICROBIAL ACTIVITY FOR BACTERIA GRAM POSITIVE, NEGATIVE AND YEAST

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Background
Silver and its compounds have strong antimicrobial activities for bacteria, fungus, and viruses. Recent advances in researches made on metal nanoparticles appear to revive the use of silver nanoparticles (SNPs) for antimicrobial applications. It has been shown that SNPs prepared with a variety of synthetic methods have effective antimicrobial activity.

Objectives
In this work, silver nanoparticles (Ag NP) with 5% silver chloride (NaNoRa²), were used to evaluate the antimicrobial activity against various bacteria Gram-positive, Gram-negative and yeasts.

Methods
Silver nanoparticles (NanoRa²) were synthesized with silver chloride to 5%, by the Sol-Gel method. Silver nanoparticles were characterized by SEM (Scanning Electron Microscope). Silver nanoparticles (NanoRa²) were tested for different Gram-positive bacteria as Enterococcus faecalis, Staphylococcus aureus, Staphylococcus epidermidis, and Gram-negative bacteria such as Klebsiella pneumoniae, Escherichia coli, Pseudomonas aeruginosa, Proteus mirabilis and vulgaris, as well as the yeast Candida albicans. The antibacterial activity of the silver nanoparticles was evaluated following the guidelines for the Kirby-Bauer diffusion method according to the CSLI.

Conclusions
Nanoparticles have a particle size of 100nm. The spectrophotometric profile showed characteristic IR peaks, links titanium dioxide that have the NanoRa². The silver nanoparticles were able to inhibit the growth of Gram-positive, Gram-negative and yeasts from 10 ppm and 50 ppm. In the case of Staphylococcus aureus methicillin resistant as well as Candida albicans, the silver nanoparticles at a concentration of 50 ppm, larger zones of inhibition were showed.
Background

Candida spp., major opportunistic pathogens, infects immunocompromised people and is responsible for life-threatening infections. The individuals commonly affected include those with oral candidiasis of HIV-positive patients, cancer patients undergoing radiotherapy, and xerostoma patients. Candida infections may further result in severe systemic infection and convert multiple organ failure (1). Moreover, Candida spp. can develop resistance to multiple antifungal drugs by over-expression of drug resistance-related targets.

Objectives

In this study, we developed a patented anti-fungal peptide P-113 to overcome the emerging issue of drug-resistance in Candida infections. We also modified P-113 by tandemly repeating sequences to improve the candidacidal efficiency.

Methods

The P-113, derived from human histatin-5 protein, retains a full antibacterial activity and exhibits a wide spectrum of activity in vitro against both bacteria and fungi (2). Several drug- (azoles-) resistant clinical strains including C. albicans, C. krusei, C. glabrata, and C. tropicalis were isolated from oral candidiasis patients. We investigated the susceptibility of these drug-resistant strains to P-113 by candidacidal assay (3).

Conclusions

The results showed that P-113 was able to kill these drug-resistant strains, suggesting a promising candidacidal activity of P-113 toward drug-resistant Candida spp. To improve the candidacidal activity, we designed P-113-derived dimeric and trimeric peptides which contain 2- and 3-time tandem repeats. The two tandem repeats improved the candidacidal potency of the peptides, and the dimer and trimer
have IC\textsubscript{50} values against \textit{C. albicans} of 0.5 and 0.4 \textmu M, respectively, compared with the monomer P-113 (IC\textsubscript{50} = 2.3 \textmu M). P-113 and its derivatives show a potential for the development of novel therapeutics against \textit{Candida} infections.
SYNERGISM OF FLORFENICOL IN COMBINATION WITH ANALOG THIAMPHENICOL AGAINST STAPHYLOCOCCUS HYICUS, STREPTOCOCCUS SUIS, AND PASTEURELLA MULTOCIDA

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Background
Florfenicol (FF) and thiamphenicol (TAP) are amphenicol antibiotics commonly used in veterinary and aquiculture practice with broad-spectrum antibacterial activity. Previous study found that FF and TAP exhibited synergistic effect against Staphylococcus aureus (including methicillin-resistant) clinical isolated strains.

Objectives
To further investigate the spectrum of synergistic activities of FF and TAP combination.

Methods
The antibacterial activities of FF and TAP combination against a total of 69 strains of Salmonella enterica, Pasteurella multocida, Pseudomonas aeruginosa, Riemerella anatipestifer, Streptococcus suis, and S. hyicus were evaluated by checkerboard assay and time-kill study. For protective effects of combination therapy on chicken, FF and TAP alone or in combination were administered intramuscularly 30 mins prior to the intramuscular infection of P. multocida.

Conclusions
The checkerboard assay showed additive effect (0.5≤ fractional inhibitory concentration (FIC) index ≤1) of FF/TAP combination in 75% of P. multocida, 53% of S. suis, 50% of S. hyicus, 33% of P. aeruginosa and 11% of S. enterica isolates that also include FF and TAP-resistant strains. In addition, 8 strains of P. multocida, S. suis, and S. hyicus were further demonstrated in the time-kill study confirming synergism between these two drugs. It also appeared that the combination enhanced the bacteriostatic effect to bactericidal on the 3 bacteria. Most notably, the P. multocida-challenged chickens were effectively protected by FF/TAP combination at lower dosages. Present study demonstrated the potentiality of a novel combination regimen of the same class of antibiotics against P. multocida infections.
DISCOVERY OF NEW ANTIMICROBIALS FROM ACTINOALLOMURUS

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Background

These past few years have been changing dramatically the landscape of infectious diseases, showing that investment on drug discovery should not be disregarded as a steady emergence and spread of antibiotic resistance keeps occurring and giving origin to more multi-drug resistant strains. One possible source to enhance the antibacterial arsenal is to look on what Nature has been using to solve the very same problem. With the myriad of species waiting to be scrutinized, it is wise to give priority to those less exploited but phylogenetically related to known producers.

Objectives

This project is part of a screening program designed to assess the metabolic potential of the recently described genus Actinoallomurus, and to deliver new molecular structures.

Methods

The program includes small-scale fermentations followed by in vitro and in vivo bioactivity tests on fermentation broths extracts. Positive samples are then subjected to a combination of bioassays and chemical analysis (HPLC, MS and NMR and a query within ABL (a proprietary antibacterial database) to determine the novelty of active compounds and discard those already known.

Conclusions

Our study shows Actinoallomurus carries the biosynthetic machinery to create a vast range of different active molecules. The dereplication process of 200 Actinoallomurus microbial extracts revealed the potential to produce a broad range of chemical classes, from lantipeptides to new hyperhalogenated angucyclines or aromatic polyethers. The diversity of the clusters of secondary metabolism found in the sequenced genome of an Actinoallomurus, seems to corroborate the theory that this genus may become an important producer of new active molecules.
Background

The human pathogen *Staphylococcus aureus* is responsible for many hospital-associated and community-acquired infections. The emergence of methicillin-resistant *S. aureus* (MRSA) and vancomycin-resistant *S. aureus* strains has prompted research into staphylococcal vaccines and preventive measures. Staphylococcal clumping factor A (ClfA), iron surface determinant B (IsdB) and gamma hemolysin B (HlgB) are key virulence factors and opsonic activity plays an important role in successful vaccine development against *S. aureus* infections.

Objectives

In this report, the BALB/c mice were immunized with a recombinant protein (CIH) composed of selected antigenic regions of ClfA, IsdB and HlgB of *S. aureus*. After the immunization, opsonophagocytic activity assay (OPK) was performed to all serum samples.

Methods

After exposing the opsonized bacteria with peritoneal macrophages, the phagocytic capacity of macrophages exposed with serum from mice immunized with recombinant protein CIH showed a significant increase as compared with macrophages exposed with serum from mice immunized with PBS (p = 0.002).

Conclusions

Our finding indicated that immunization of BALB/c mice with recombinant protein CIH promotes the phagocytosis of *S. aureus* and enhances the clearance of the *in vivo* pathogen.
Background

Due to the resistance of microorganisms to antibiotics, studying alternative methods such as plant extracts becomes very important.

Objectives

Evaluating the in vitro antifungal activity of Persea americana extract on C. albicans biofilm and its cytotoxicity in macrophage culture (RAW 264.7).

Methods

For the determination of the minimum inhibitory concentration (MIC), microdilution in broth (M27-A2 standard) was performed. Thereafter, the concentrations of 12.5, 25, 50, 100 and 200 mg/mL (n=10) with 5 min exposure were analyzed on mature biofilm in microplate wells for 48 h. Saline was used as control (n=10). After treatment, biofilm cells were scraped off and dilutions were plated on Sabouraud dextrose agar. After incubation (37°C /48 h), the values of colony forming units per milliliter (CFU/mL) were converted to log10 and analyzed (ANOVA and Tukey Test, 5%). The cytotoxicity of the P. americana extract was evaluated on macrophages by MTT assay.

Conclusions

Results: The MIC of the extract was 6.25 mg/mL and with 12.5 mg/ml there was elimination of 100% of planktonic cultures. Regarding the biofilms, a significant reduction (p

Conclusion: P. americana extract showed antifungal activity against C. albicans biofilm, whereas the concentration of 50 mg/mL presented the best results.
EVALUATION OF MYRTUS COMMUNIS AND ZIZIPHUS LEAVES ANTIBACTERIAL ACTIVITY ON IMP-TYPE METALLO-BETA-LACTAMASE PRODUCING PSEUDOMONAS AERUGINOSA STRAINS

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Background

Pseudomonas aeruginosa is one of the leading nosocomial pathogens. In recent years, inappropriate use of antibiotics is the cause of untreatable ‘superbugs’ and high death rates in clinics in worldwide. One of the mechanisms of resistance to carbapenem antibiotics in P. aeruginosa is metallo-β-lactamases (MBLs) production that hydrolyzes all carbapenems that are the last generation of drugs for treatment of infections caused by MDR P. aeruginosa.

Objectives

The aim of this study is evaluating the effects of Myrtus communis leaves and ziziphus leaves methanolic extract on IMP-Type metallo-beta-lactamase producing P. aeruginosa strains.

Methods

This experimental study was carried out on hospitalized burn patients during 2012-2013. Antibiotics and extracts susceptibility tests were performed by disc diffusion and broth microdilution methods. MBL detection was performed by Combination Disk Diffusion Test (CDDT). The bla(VIM) and bla(IMP) genes were detected by PCR and sequencing methods.

Conclusions

It was found that among 83 imipenem resistant P. aeruginosa strains, 48 (57.9\%) were MBL producers. PCR and sequencing methods proved that these isolates were positive for blaIMP-1 genes, whereas none were positive for bla (VIM) genes. All MBL-producing P. aeruginosa were resistant to Meropenem, Imipenem, Ceftazidime, Amikacin, Tobramycin, Ciprofloxacin, Aztreonam, Piperacillin/Tazobactam, Ceftriaxone, Cefepime and Carbenicillin. The result of MIC for imipenem was 128(μg/ml) for all strains. The MIC (mg/ml) and MBC (mg/ml) results of Myrtus...
*communis* was 6.25 and 12.5 (mg/ml) for all isolates, respectively. Although the ziziphus leaves methanolic extract represent any significant result on this isolates.
IDENTIFICATION AND CHARACTERIZATION OF A NOVEL CRYPTIC CATIONIC ANTIMICROBIAL PEPTIDE FROM THE THIRD DOMAIN OF LIFE

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Background

The huge increasing of bacterial strains resistant to classical antibiotics is threatening the public health security, thus the need of developing new antimicrobials has become critical. Cationic Antimicrobial Peptides (CAMPs) represent suitable alternatives, since they do not lead to selection of resistant strains. Indeed, CAMPs act by disrupting the membrane integrity and/or by altering its potential, so that, to acquire resistance bacteria need to remodelate the membrane composition. Such modifications could be detrimental to bacterial cells, thus dramatically reducing the onset of resistant strains.

Objectives

Although proteins involved in the immune response of multicellular eukaryotes represent a natural source of CAMPs, new methodologies for their identification are required to encounter the rising demand of more active and stable CAMPs.

Methods

A powerful bioinformatics tool has been recently set up and used to predict the antibacterial activity of cryptic CAMPs (i.e. embedded in the primary structure of proteins). Given their hydrophobic/cationic nature, sequences of transcription factors were chosen as source of choice for the discovery of new cryptic CAMPs. In particular, our analysis focused on extremely stable proteins from thermophilic Archaea.

Conclusions
Among the identified CAMPs, one from the DNA-binding protein Stf76 (here named PepC) has been structurally and functionally characterized. This peptide has the capability of inducing both membrane fusion and pore formation. Interestingly, the DNA-binding activity was retained opening the way to speculate about novel mechanisms of action. Moreover, its antimicrobial activity has been proved on clinical strains isolated from patients affected by cystic fibrosis.
ANTAGONISTIC ACTIVITY OF MARINE SPONGE-ASSOCIATED BACTERIA COLLECTED AT WIMEREUX (FRANCE) AGAINST CLINICAL STRAINS

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Background
Marine sponges are an important-and yet understudied-reservoir of new antimicrobials, a majority of which are produced by associated microbes.

Objectives
We tested the antibacterial activity of 110 bacterial strains isolated from two sponges (Halichondria panicea and Hymeniacidon perelvis) collected off rocky substrate in the low intertidal zone at Wimereux, France.

Methods
Strains were isolated on various media (R2A, Brain Heart Infusion (BHI)-agar and Marine agar) and tested against three target clinical strains: Pseudomonas aeruginosa ATCC 27853, Escherichia coli ATCC 25922 and Staphylococcus aureus ATCC 2913. Antagonistic activity was measured in double layer agar assays.

Conclusions
Fifty strains (45%) were able to inhibit the growth of at least one clinical strain (most often S. aureus). Antibacterial-producing strains belonged to Gamma-Proteobacteria (88%, mostly Pseudomonas spp.) and Firmicutes (12%, among which Lactococcus spp. and Vagococcus spp.). The production of antimicrobial substances turned out to be modulated by the medium formulation, its salinity (tested from 0.5 to 3.5%) and the temperature used in antagonism assays (tested from 8 to 25°C). The best results were obtained with BHI-agar, the lowest salinity and highest temperature tested. In all cases, antimicrobial substances were not soluble, as the antagonistic activity was lost when cultures were filtered on 0.2 mm-pore membranes. This study confirmed the potential of sponge-associated microbes for the production of antibacterial substances.
Background

With the emerging threat of infections caused by multidrug-resistant bacteria and scarce prospects of newly introduced antibiotics in the future, bacteriophages (phages) are currently being reconsidered as alternative therapeutics. However, the full breath of phage diversity suitable for treatment of bacterial infections is still largely unexplored.

Objectives

This study aims at providing an overview of the general occurrence of bacteriophages (with therapeutic potential) in natural environments and at devising strategies for their rapid characterization and classification at molecular level.

Methods

Novel phages were isolated from various aquatic systems (e.g. from general and hospital wastewater, activated sludge samples from sewage plants, streams, rivers, ponds and lakes). Environmental samples were pre-incubated with clinical isolates of multidrug-resistant bacteria. Enriched bacteriophages were subsequently obtained with the double agar layer plaque assay. “PhiSigns” (http://www.phantome.org/phisigns/) was used for phage identification based on the detection of signature genes. Genomic comparison of phages was performed based on digitized fluorescent restriction-fragment-length-polymorphism-analysis (fRFLP).

Conclusions

For roughly 50% of tested bacterial strains lytic bacteriophages were found against multidrug-resistant Pseudomonas aeruginosa, Klebsiella pneumoniae and Enterobacter sp. with waste water samples harbouring the highest phage diversity. By contrast no bacteriophages were isolated against multidrug-resistant Staphylococcus aureus and Acinetobacter baumannii. PhiSigns combined with fRFLP proofed to be a useful tool for rapid phage identification, which is the base for their proper selection for ultimate whole genome sequencing. S. aureus and A. baumannii phages require additional isolation steps such as FeCl₃-precipitation and/or use of...
subinhibitory antibiotic-concentrations for enhancing plaque visibility on agar plates.
ANTIFUNGAL ACTIVITY OF ESSENTIAL OIL FROM ABIES HOLOPHYLLA MAXIM AGAINST DERMATOPHYTES

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Background
Dermatophytes are well known for one of world-wide cause of diseases such as tinea pedis, tinea cruris, and tinea corporis. Dermatophytes infections are increasing of taking antifungal agents, accordingly increased side effects like toxicity and tolerance. That was why searches on natural antifungal medicine have motivated. Many kinds of woody essential oils have antifungal activity, especially essential oils of conifers. \textit{Abies holophylla} Maxim (AH) is widespread in East Asia and its essential oil was known to have effective antifungal actions against \textit{Aspergillus fumigatus}.

Objectives
Therefore, this study was to investigate the antifungal activity of AH essential oil against dermatophytes, such as \textit{Epidermophyton floccosum}, \textit{Trichophyton mentagrophytes} and \textit{Trichophyton rubrum}, and to determine the potential effective compound as dermatitis treatment.

Methods
To evaluate the potential antifungal activities of AH essential oil and its fractions, Minimum Inhibitory Concentration (MIC) measurement and agar dilution method were used with morphological observation. Also, their major constituents were analyzed by GC/MS. The morphological changes of the dermatophytes exposed to the AH essential oil were observed by electron microscopes.

Conclusions
As the results, the highest activities were identified in the fraction containing borneol, limonene, 3-carene, and camphene. Also, morphological observation using electron microscopes (SEM and FEM) showed a dramatic changes of cell wall of dermatophytes exposed to the AH essential oil. In conclusion, AH essential oil and its constituents were expected to be used as antifungal agent or raw material for dermatitis therapy.
New antimicrobials for resistant organisms

IN VITRO ANTI-STAPHYLOCOCCAL ACTIVITY OF SECONDARY METABOLITES FROM DROSERA BINATA COMBINED WITH SILVER NANOPARTICLES

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Background

Antibiotic resistant Staphylococcus aureus is a serious nosocomial pathogen and the major cause of life threatening infections, like burn wound infections. There is an urgent need to overcome antibiotic resistance by developing new antimicrobial treatment. The use of active plant secondary metabolites simultaneously with silver nanoparticles (AgNPs) against S. aureus is based on multi-target strategy.

Objectives

The aim was to find an alternative anti-staphylococcal treatment based on the combination of Drosera binata secondary metabolites and silver nanoparticles.

Methods

Plant extracts were obtained from in vitro cultured D. binata tissues. Antibacterial activity was determined by Broth Microdilutions Method. Checkerboard Titration technique was used to study interactions between AgNPs and extracts or secondary metabolites. Composition of the extract was established by the HPLC-MS analysis. The MTT Assay was used to evaluate cytotoxicity of antimicrobial agents towards in vitro cultured human keratinocytes.

Conclusions

Both silver nanoparticles and the D. binata extract possess antibacterial activity towards various S. aureus strains regardless of their antibiotic resistance. The synergistic bactericidal activity was observed when AgNPs and plant extracts were combined. This allowed to significantly reduce the bactericidal concentrations of AgNPs and the extract by 50% and 97%, respectively. Among all extract constituents only the chlorine derivative of plumbagin (ChPL) showed synergistic bactericidal activity with AgNPs. Moreover, reduced bactericidal concentrations of mixtures of
AgNPs with the extract or ChPL were observed to be non-cytotoxic towards human keratinocytes. Obtained results indicate that combination of AgNPs and secondary metabolites from *D. binata* tissues may provide an alternative antimicrobial treatment.
New antimicrobials for resistant organisms

ANTIBACTERIAL AND ANTIFUNGAL ACTIVITY OF NEWLY SYNTHESISED BENZOXABOROLES, BENZOSILOXABOROLES AND PYRIDOXABOROLES

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Background

Benzoxaboroles, and related compounds, are a particularly interesting group of synthetic agents. Their diverse biological activities and widespread applications in medicinal chemistry have undergone intense scientific investigation.

Objectives

We have examined whether new benzoxaborole-based derivatives and their analogues possess antimicrobial activity and are substrates for multi-drug-resistance efflux pumps.

Methods

The antibacterial and antifungal activities of newly synthesised compounds: 3 benzoxaborole-, 3 benzosiloxaborole- and 6 pyridoxaborole-based derivatives were evaluated. Antimicrobial activity was screened using the disc-diffusion test (0.4mg of the derivative was applied to a filter paper disc). The MIC and MBC values were estimated according to CLSI and EUCAST recommendations. During determination of MIC values ± PAbN (80mg/L), the concentrations of tested compounds in the broth medium ranged: 0.39-400mg/L. The 19 reference ATCC strains of Gram-positive, Gram-negative bacteria and yeast were used in this study.

Conclusion:

Generally, the most potent antibacterial and antifungal activities were found for fluorine substituted benzosiloxaboroles – analogues of benzoxaboroles possessing a silicon atom for carbon substitution at the 3-position (MICs 25-400mg/L and 0.78-100mg/L, respectively). Interestingly, the B(OH)₂-substituted 3-phenylbenzoxaboroles were found to be the most potent against Gram-positive cocci: Staphylococcus sp. (MICs 12.5-50mg/L), Enterococcus sp. (MIC 200mg/L) and yeasts (MIC 50mg/L). Furthermore, we demonstrated that benzoxaboroles were removed by efflux pumps present in Gram-negative bacteria. Of note, only two pyridoxaborole-based compounds were found to be weakly active against Staphylococcus sp.

These data demonstrate structure-activity relationships of the tested derivatives and
highlight the need for further investigations into the antimicrobial properties of new benzoxaboroles and related compounds.
APPLICATION OF A HIGH-THROUGHPUT ASPERGILLUS FUMIGATUS CELL WALL STRESS REPORTER SYSTEM: IDENTIFICATION OF SYNTHETIC PEPTIDES INCREASING THE SENSITIVITY FOR ANTIFUNGAL MEDICINES

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Background

Increased resistance to currently used antifungal compounds and the fact that these agents are often harmful to man and environment have resulted in a growing demand for new antifungals, which selectively act on cellular processes that are unique to fungi. To meet this demand, we have established in A. niger a luciferin/luciferase based reporter system for high-throughput screening of natural products for identification of potential new antifungal drugs.

Objectives

Our system allows us to identify compounds that specifically inhibited the fungal cell wall biosynthesis.

Methods

Recently the system has been developed for A. fumigatus and non-Aspergillus species. The system was validated by analyzing its performance with several antifungal drugs which are commonly used in the clinic like Caspofungin, Amphoterin B, Voriconazol (V fend) as well as Nikkomycin. A dose response behavior of the compounds could be demonstrated together with in time different types of stress responses. Importantly, whether the compound tested is fungistatic or fungicidal could be clearly indicated.

Subsequent analysis of different synthetic antimicrobial peptides, HTX1-4, revealed also a moderate effect on cell wall stress induction indicated by an increase in Lux activity.

Conclusions

Interestingly, incubation of these peptides at sublethal concentrations together with some of the antifungal medicines showed a considerable increase in lux activity.
and a significant increase in sensitivity of *A. fumigatus* for these medicines. Growth test with sublethal concentrations of HTX-2 on EBV-LCL, K562 cells and PHA-blasten, and also cytotoxicity- and genotoxicity tests with NCTC2544 cells did not show any negative/toxic effects on these human cell lines.
Background

Lantibiotics are potent peptide antimicrobials active against Gram-positive bacteria, including multidrug resistant pathogens. They display various posttranslational modifications, with lanthionine rings and dehydroamino acids as key signatures of this family of antimicrobials. In addition, some lantibiotics contain other enzymatic modifications that have a role in their activity.

Objectives

Our purpose is to combine different enzymes from diverse lantibiotic production systems in order to hypermodify the model lantibiotic nisin as a proof-of-principle. This can set a methodology for the design and modification of other (antimicrobial) peptides.

Methods

We have combined, in either Lactococcus lactis or Escherichia coli, the modification machinery of nisin (the dehydratase NisB, the cyclase NisC) with the hydroxylases MibO and CinX from the microbisporicin and the cinnamycin machineries, respectively; the C-terminal decarboxylase GdmD from the gallidermin cluster; and the reductase LtnJ from the lacticin 3147 system.

Conclusions

We have generated nisin derivatives containing hydroxyproline, hydroxyaspartate, a C-terminal aminovinyl-cysteine, or D-alanine in their structure. Moreover we have confirmed that these additional enzymes do not require leader peptide recognition to modify their substrate.

Our results provide an insight for the application of Synthetic Biology for the design and à la carte modification of (antimicrobial) peptides.
New antimicrobials for resistant organisms

THERAPEUTIC MONOCLONAL ANTIBODIES AGAINST THE GLOBALLY SPREAD MULTI-DRUG RESISTANT ESCHERICHIA COLI CLONE ST131-O25B:H4

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Background
The globally spread multi drug resistant E. coli clone ST131-O25b:H4 is responsible for a significant proportion of multi-drug resistant extraintestinal infections. By definition, isolates of this clone express the O25b antigen, whose unique structure has been resolved recently.

Objectives
We aimed to develop humanized monoclonal mAbs against the O25b antigen that may provide an alternative adjunct or stand-alone therapeutic option against this E. coli lineage.

Methods
Murine mAbs against the unique O25b antigen were generated by standard hybridoma technique. Following a primary screening, selected antibodies were subjected to humanization. Specificity and binding characteristics were assessed by immunoblots and ForteBio using purified antigens. Surface staining of ST131-O25b isolates irrespective of the capsular type expressed was shown by flow cytometry. Upon surface binding, the humanized antibodies induced complement mediated bactericidal activity in vitro. Passive immunization with the humanized mAbs elicited high level of protection at low mAb doses in a murine model of bacteremia. All whole cell assays were performed with bacteria grown in standard culture medium as well as in depleted pooled human serum in order to mimic in vivo-like conditions.

Conclusions
Humanized mAbs against E. coli O25b antigen were developed that elicited bactericidal activity in vitro and showed protection in vivo. Prophylactic passive immunization of colonized individuals or adjunct therapy of infected patients by mAbs may replace/substitute antibiotic therapy against this drug resistant clone. Relevant cases could be identified by the co-developed companion diagnostic tool.
FREE AND IMMobilIZED PHOTOSENSITIZERS FOR ERADICATION OF PATHOGENIC BACTERIA

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Background. The problem of controlling microbial infections remains one of the most serious challenges faced by mankind, despite the development of very effective antimicrobial agents. Bacterial resistance to antibiotics, which for decades were considered to be capable of combating any bacterial infection, actually destroyed hopes for combating pathogenic bacteria. One of alternatives to antibiotic treatment is photodynamic antimicrobial chemotherapy based on visible light-activation of dyes called photosensitizers. In the dark, these photosensitzers are non-toxic or have low toxicity. No bacterial resistance to photosensitizers has been reported to date.

Objectives. The present report is dedicated to the antibacterial activity of the photosensitizers Rose Bengal and Methylene Blue against Gram-positive Staphylococcus aureus. The antimicrobial activity of both free and immobilized photosensitizers was examined.

Methods. Rose Bengal and Methylene Blue were immobilized on polystyrene, polycarbonate and poly(methyl methacrylate) by dissolution in chloroform with further evaporation of the organic solvent and the formation of thin polymeric films. Immobilization of these photosensitizers on polyethylene and polypropylene was performed by dissolution in the melted polymers. The free and immobilized photosensitizers showed high efficiency in eradication of S. aureus as tested by the MIC, the colony-forming units count and measurement of the inhibition zone.

Conclusions. Photodynamic inactivation of S. aureus by Rose Bengal and Methylene Blue opens prospects for developing an alternative to antibiotic treatment of bacterial infections.
ANTIMICROBIAL PEPTIDES OF MARINE ORIGIN AND THEIR MOLECULAR CHARACTERIZATION

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Background

Marine organisms are continuously exposed to microbial population and are equipped with effective immune molecules to combat microbial invasion. Antimicrobial peptides are important components of innate defense mechanisms exhibited by animals. These organisms produce a range of AMPs varying greatly in size, structure and other physicochemical properties. Investigating novel antimicrobial peptides from marine organisms can provide new insight into the immune response of these organisms and a possibility of discovering new and effective drugs in medicine/aquaculture.

Objectives

Screening marine organisms for discovering novel antimicrobial peptides

Methods

Total RNA was extracted from haemocytes and cDNA was synthesized. PCR amplification was done using gene specific primers for AMPs. Amplicons were cloned and sequencing of the inserts were done at SciGenom, India. Homology search was performed by using BLASTn and BLASTp at NCBI and physio-chemical characterization by ProtParam Tool. Phylogenetic tree was constructed based on amino acid sequences by Neighbor joining method using MEGA 5.0

Conclusions

Four novel ALF isoforms were characterized, among which two were identified from crabs (Scylla serrata and Portunus pelagicus) and two from shrimp (Fenneropenaeus indicus). Four novel isoforms of crustins were also identified and characterized from Scylla serrata, Scylla tranquebarica, Portunus pelagicus and Fenneropenaeus indicus. Two penaeidin isoforms were characterized from Fenneropenaeus indicus and Penaeus monodon and a hepcidin from Chlorophtalmus bicorins. The marine environment, with its enormous biodiversity remains a largely untapped reservoir of bioactive peptides. Recombinant expression and bioactivity testing of these peptides
will definitely contribute to the design and formulation of potential peptide drugs for application in medical/ aquaculture industry.
Background

Today we witness the global health threat of antimicrobial resistance. This phenomenon relies largely on the erosion of the effective antimicrobials and has drawn the attention of researchers from the traditionally used antimicrobials to the so-called experimental compounds.

Experimental compounds are bioresearch compounds that have been subject to a certain degree of development, but have been put away either due to project deprioritization or lack of effectiveness against the application in question. They may, however, be very valuable in other applications; for instance, in diseases sharing similar biochemical pathways. These compounds are a rich source of new and, possibly, better antimicrobials. The question is: How can researchers gain access to them? We will discuss our efforts in developing a compound sharing facility and how we will collaborate with the community to facilitate open-innovation models for the antimicrobial compounds discovery.

Objectives

Our goal is to advance research in the field of antimicrobial resistance by repurposing rare and one-of-the-kind experimental compounds to the whole of academic community. For that, we developed ResearchAnt Hub (RAH). RAH is powered by the ResearchAnt Foundation, which is a not-for-profit organization. RAH is a repository and a platform that fosters researchers worldwide to use and share their experimental compounds.

Methods

Our repository will contain thousands of experimental compounds that were developed or manufactured by privately- and academically-led researchers during the course of their work.

Conclusions

RAH is a unique not-for-profit open-innovation initiative. It represents a major step towards the discovery of new drugs to counteract today’s ever-increasing antimicrobial resistance problem.
Background
Antibiotic resistant human pathogens are diminishing our ability to treat infections and novel antimicrobials are urgently needed to replace our current arsenal of antimicrobials. *Pseudomonas aeruginosa* strains have been shown to produce antimicrobial compounds including bacteriocins, pigments and secondary metabolites that show promise as novel antimicrobial therapeutic agents.

Objectives
To assess the antimicrobial activity of four environmental *P. aeruginosa* isolates from University College London, Hospital (UCLH).

Methods
Four *P. aeruginosa* isolates were obtained from various sites in UCLH. Cross streak assays using indicator organisms including *Escherichia coli, Staphylococcus aureus* and *Salmonella typhimurium* were conducted to determine the spectrum of inhibition of these *P. aeruginosa* strains. A spot-on lawn assay was used to compare the ability of each strain to inhibit the growth of *S. aureus*. To determine if the *P. aeruginosa* strains secreted compounds that were inhibiting *S. aureus*, this indicator strain was grown in cell free media that the *P. aeruginosa* strains were previously grown in.

Conclusions
Competitive hospital environments may be a rich source of bacteria producing novel antimicrobials. The results of the above experiments show that these four hospital environment acquired *P. aeruginosa* isolates are capable of inhibiting a wide range of bacteria. Some strains had a greater ability to inhibit *S. aureus* than others. All of the *P. aeruginosa* strains secreted inhibiting compounds into the media they were grown in. We are currently characterising the genetic and biochemical basis of these compounds.
Background: Staphylococcus aureus including methicillin-resistant S. aureus (MRSA) is recognized as a globally important pathogen in food-borne disease. Therefore bacteriocins have been focused on new weapon that are considered safe and effective substances as food additives to prevent food-borne pathogens.

Objectives: To control MRSA of agricultural product, identify and characterize a antimicrobial peptide (bacteriocin) produced by Staphylococcus epidermidis.

Methods: S. epidermidis were isolated from leafy green vegetables and identified by API Staph ID test and 16s rRNA sequence analysis. Antimicrobial activities of S. epidermidis against antibiotic resistant S. aureus and other Gram (+) bacteria were tested using agar diffusion assay. Purification of the active compound was achieved using a combination of ammonium sulfate precipitation, dialysis and reverse-phase high-performance liquid chromatography (RP-HPLC). The effects of heat, pH and enzymes on bacteriocin activity were determined using S. aureus CCARM3723 as indicator organism.

Conclusions: S. epidermidis S-199 strain showed broad and strong antimicrobial activity. Purified antimicrobial peptide was heat-labile but stable in a wide range of pH. And the antimicrobial peptide showed resistance to α-amylase, lipase, and RNase, and DNase. However, antimicrobial activity was lost after trypsin, pronase E, pepsin, α-chymotrypsin, proteinase K, and papin treatment.
New antimicrobials for resistant organisms

ROLE OF PRODIGIOSIN AS A MICROBicide

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Background

Number of microbial species are metabolically wired to produce some pigments constitutively. Prodigiosin is such a pigment having prodigious biological activities in accord to its name. Microbicides are compounds that can be applied inside the vagina or rectum to protect against sexually transmitted infections (STIs). Discovery of drugs, targeting multiple steps in multiple diseases is of great interest.

Objectives

To evaluate prodigiosin for its microbicide potential and study its mechanism of actions in restricting sexually transmitted infections like \textit{E. coli}, \textit{C. albicans}, HSV, HIV, \textit{Trichomonas vaginalis}. ii) To check cytotoxicity of prodigiosin against human cells.

Methods

96 well plate assay used for antibacterial assay followed by its biochemical investigations after treatment with prodigiosin at LC\textsubscript{30} concentration, cell associated and cell free assays were used to check anti HIV-1 (CXCR4 and CCR5) potential, while ONPG for HSV, cytotoxicity of prodigiosin was determined by MTT assay.

Conclusions

Prodigiosin induces leakage in bacterial cell membrane and simultaneously inhibits activity of few essential enzymes. Prodigiosin inhibits viral entry and cell fusion for HSV and HIV (IC\textsubscript{50} - 0.01 µg/ml). It also inhibited \textit{T. vaginalis} (1.2 µg/ml). The use of pigment found on safer side against human cells (PBMC (78±3.1 µg/ml) and HeLa cells (70.2±5.3 µg/ml)). One compound having all these activities is a remarkable feature. Prodigiosin can be formulated as a microbicide (drug) and applied on the affected area with a cream base so as to treat the STIs. In conclusion prodigiosin can be used in treatment modality for protection from STIs.
Background
Actinomycetes are prolific producers of natural products, including anticancer compounds, antifungals and antibiotics. The treasures that lie hidden in the actinomycete genomes may well be our final resource in the battle against the rapidly emerging infectious diseases associated with multi-drug resistance.

Objectives
We aim to understand the triggers and cues that elicit antibiotic production in the soil as well as in the laboratory. This knowledge is then translated into technologies for the activation and mining of silent pathways.

Methods
To uncover novel antimicrobials, we use a combination of ecological insights, systems biology, NMR-based metabolomics and genome mining to uncover the regulatory mechanisms that control antibiotic production. We then apply this for the discovery of novel antimicrobials.

Conclusions
One major control system revolves around the nutrient sensory protein DasR, which pleiotropically controls antibiotic production. Many other control systems are undoubtedly in place to allow actinomycetes to respond appropriately to challenges by competitors and fluctuations in nutrient composition in the habitat. Furthermore, once a bioactivity is elicited under specific growth conditions, novel approaches are needed to rapidly identify the bioactivity of interest and link it to a specific gene cluster. Molecular and ecological insights to elicit antibiotic production will be discussed, and examples of novel antibiotics identified by these approaches are presented.
FEMS-1522
New antimicrobials for resistant organisms

ACTIVITY ANALYSIS OF LYSOSOMES AS A FUNCTIONAL CELL ORGANELLE IN SACCHAROMYCES CEREVISIAE

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Background
Lysosomes are functional cell organelles which have antimicrobial and anticancerous activity. Lysosomal enzymes are easily integrated or released when exposed to some stimuli factors in all eukaryotic cells.

Objectives
In order to verify features special function of lysosomes, several conditions were used to stimulate \textit{in vitro} functions of lysosomes. In addition, we confirm lysosomal safety and activity \textit{in vivo} for using antimicrobial and anticancerous materials.

Methods
Proteomic analysis of lysosomal enzymes in response to the enhanced and repressed condition \textit{for in vitro} activity using 2-DE. Among up-regulated protein dots response to oxidative stress, the dot with highest increase was chosen to overexpress in \textit{Saccharomyces cerevisiae} and then confirm the lysosomal activity. In addition, we injected lysosomes into rat model infected with bacteremia, and then analysis bacterial growth in blood of rat according to the reaction time.

Conclusions
The antimicrobial and anticancerous activity of lysosomes was enhanced in cloned \textit{S. cerevisiae in vitro} assay. In addition, it is safe to inject lysosome into normal rat. Moreover, when being injected into rat model infected with bacteremia, lysosome showed its ability to inhibit bacterial growth in blood. Therefore, increases in endogenous levels of lysosomes may have various applications such as antimicrobial agents and apoptosis-inducing materials for cancer cells and it is very promising to use the organelles to improve \textit{in vitro} and \textit{in vivo} functions.

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EFFECT OF ACID TREHALASE (ATH) ON IMPAIRED YEAST VACUOLAR ACTIVITY

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Background
Acid trehalase (ATH) has been suspected to have two different localizations, vacuolar lumen and cell surface, among which the localization at cell surface is believed to be responsible for extracellular trehalose utilization.

Objectives
A lot of efforts were made to confirm the exact localization of ATH in yeast cells. Along with this, the trafficking pathway responsible for the delivery of ATH into yeast vacuoles was also studied with the purpose to provide better control on the transportation of this protein in yeast cells.

Methods
ATH was overexpressed in yeast cells grown on trehalose-containing medium to observe the impact of this enzymes on yeast vacuolar activity. After that, 4 groups of typical proteins representing the 4 main trafficking pathways in yeast were also selected to analyze their expression pattern via 2DE.

Conclusions
The overexpression of ATH was observed to decrease vacuolar activity. Moreover, ATH was confirmed again to locate in both cell surface and vacuoles. Finally, it is interesting to discover that on medium containing trehalose, the ATH’s localization at cell surface, but not vacuoles, is prioritized to utilize extracellular trehalose for cell growth. The mechanism behind that is suggested via the down regulation of MVB pathway that sorts ATH into vacuoles.

Acknowledgement: This work was carried out with the support of “Cooperative Research Program for agriculture Science & Technology Development (Project No:PJ01052701)’ Rural Development Administration, Republic of Korea.
Background

In the last few decades, the emergence of bacterial resistance to antibiotics has become a common phenomenon. As a consequence, the effectiveness of antibiotic treatment of bacterial infection has progressively decreased. Therefore, it's necessary to develop and test new antimicrobial compounds against multidrug-resistant bacteria. Recently, the employment of metallic nanoparticles has emerged as an alternative to the use of organic compounds as antimicrobial agents: different metals and metalloids nanoparticles have indeed shown a promising bactericidal capability.

Objectives

The aim of the present study was to evaluate the antimicrobial activity of selenium and tellurium nanoparticles of bacterial origin against both planktonic and biofilm culture of three pathogenic strains: Escherichia coli JM109, Pseudomonas aeruginosa PAO1 and Staphylococcus aureus ATCC 25923.

Methods

To measure the antimicrobial activity of these nanoparticles we determined both the minimum biocidal concentration (MBC) and the minimum biofilm eradication concentration (MBEC) using the MBEC assay™. We also observed the effect of increasing concentrations of nanoparticles on biofilm structure using Confocal Microscopy (CLSM).

Conclusions

Our results indicate that both Se⁰ and Te⁰ nanoparticles possess antimicrobial and biofilm eradication activity. In particular Se⁰ nanoparticles exhibited antimicrobial activity at lower concentration. The activity seemed to be dependent on the dimension of the nanoparticles: indeed, the highest activity was shown by the nanoparticles smaller in size. The key observation is that bacteria grown in biofilm mode didn’t exhibit a higher level of resistance against the nanoparticles antimicrobial action. Results described in this study suggest a possible application of these nanoparticles as an effective antimicrobial agent.
Background

With the increasing availability of next generation sequencing technology, whole genome sequencing (WGS) methods are increasingly being used for bacterial typing. Key challenge is the ability to rapidly extract the relevant information from large sequence data files.

Objectives

We present two pipelines for high resolution WGS-based molecular typing: whole genome multilocus sequence typing (wgMLST) and whole genome single-nucleotide polymorphism analysis (wgSNP). Both strategies are compared using data from a *Staphylococcus aureus* outbreak.

Methods

Using an in-house developed wgMLST schema that extends the core genome schema from Leopold *et al.* (2014), we apply two independent allele calling approaches, an assembly-free and a BLAST-based allele calling algorithm, to determine locus presence and detect allelic variants in a quality-controlled manner. The wgSNP pipeline, tuned to reduce false positives while maximizing resolution, detects SNP variants by mapping the WGS reads to a reference sequence internal or external to the data set.

For both methods, all calculation-intensive data processing steps are performed on the BioNumerics® Calculation Engine, deployed locally or in the cloud.

Conclusions

The BioNumerics® 7.6 software and its integrated Calculation Engine offer a powerful platform where both wgMLST and wgSNP can be performed to provide a robust,
portable and high resolution picture of molecular typing data. The polyphasic approach allows for validation both between WGS analysis techniques and traditional techniques such as MLST or PFGE. We illustrate this by comparing core genome MLST (cgMLST), wgMLST and wgSNP, thus establishing a clear picture of the differences in resolution between those analysis techniques.
Background

Tuberculosis is a treatable disease, yet remains a major worldwide health problem. In the last 10 years approximately 80 million people contracted the disease. The development of techniques for rapid diagnosis based in molecular technologies would greatly facilitate worldwide efforts to prevent further spread of the disease. One potential diagnostic sequence, the DR locus, is present in all isolates of M. tuberculosis complex bacteria. It is useful for molecular typing of M. tuberculosis because of its fortuitous absence in non-tuberculosis strains of mycobacteria.

Objectives

In this study, we attempted to combine the specificity of molecular inversion probe (MIP) technology with the sensitivity of pyrosequencing in order to detect a short conserved 18 bp sequence included in DR locus in 25 isolates of M. tuberculosis.

Methods

DNA from 25 M. tuberculosis isolates were extracted and submitted to the MIP reactions. A MIP was designed against the DR sequence of M. tuberculosis strain H37Rv and the MIP reactions were performed; the MIPs were then submitted to pyrosequencing reactions.

Conclusions

We design a molecular inversion probe for a specific region of M.tuberculosis genome. Probing this MIP with 25 M.tuberculosis genomic DNA and determining their sequences by pyrosequencing, we detect a common sequence among all strains of M.tuberculosis. Also, we determine the minimal amount of DNA needed (50 ng) in order to obtain a good readout. By introducing a modification on pyrosequencing
methodology we were successful in detecting M.tuberculosis DNA even in presence of a very small amount of DNA (500fg).
FEMS-2810
New approaches for typing

TYPING OF ENTEROBACTERIACEAE BY ERIC-PCR, REP-PCR AND MALDI-TOF MS METHODS

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Background: The fast and reliable fingerprinting methods for typing of Enterobacteriaceae are widely required in food microbiology for tracking sources and ways of food contamination. ERIC-PCR and REP-PCR genotyping use as markers specific DNA sequences (enterobacterial repetitive intergenic consensus sequences or repetitive extragenic palindromic sequences, respectively) and the changes of their dispersal, detectable by methods. In comparison MALDI-TOF MS phenotyping uses the conformity of protein profiles. As used typing markers mirror the differences among strains only partially, also compared to other features as the antibiotics resistance and biofilm formation, their combination could propose more precise approach.

Objectives: The aim was to compare the discriminatory power of ERIC-PCR, REP-PCR and MALDI-TOF MS fingerprinting methods for different Enterobacteriaceae isolates, also with respect to their antibiotics resistance and biofilm formation.

Methods: Fifty six isolates of Enterobacteriaceae, including genera Escherichia, Enterobacter, Klebsiella, Serratia and Raoultella, isolated from food and clinical sources in Czech Republic in 2007-2014, were genotyped by the same ERIC-PCR and REP-PCR methods (maximal length of analysed fragments 3000 bp). MALDI-TOF MS typing was done by the ethanol-formic acid extraction in Bruker Autoflex Speed MALDI-TOF mass spectrometer. The resistance for 12 antibiotics and the ability to form biofilm in different media at 25 °C were tested.

Conclusions: REP-PCR is more discriminative then ERIC-PCR in most species. The discriminatory power of MALDI-TOF MS phenotyping depends on the range of used protein peaks (e.g. application of specific intensity criteria) and the used clustering methods, but never gives the false positive results in comparison to the genotyping methods.
USE OF MATRIX-ASSISTED LASER DESORPTION/IONIZATION- TIME OF FLIGHT (MALDI-TOF) MASS SPECTROMETRY FOR ROUTINE ANALYSIS IN A DRINKING WATER TREATMENT PLANT

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Background
Drinking water treatment plants employ different treatments depending on source water quality to assure adequate chemical and microbiological quality according to the Directive 98/83/EC. The study of bacterial communities throughout the treatment plant could provide a basic understanding of the effects of water processing that can be used to improve the management in drinking water treatment plants.

Objectives
To assess the use of MALDI-TOF MS for routine analysis in a drinking water treatment plant to study bacterial diversity throughout the different treatments in comparison with other methods such as the PhenePlate® system, the API 20E strip and 16S rRNA gene sequencing.

Methods
A total of 366 colonies isolated from different points and seasons throughout the water treatment process including feed water (40), sand filtration (36), ultrafiltration (44) and reverse osmosis (191) were used in the study. The colonies were analyzed by MALDI-TOF MS by direct colony inoculation on the plate. The colonies were also biochemically fingerprinted using the PhenePlate® system, clustered according to their similarity and a representative strain was selected for 16S rRNA gene sequencing and API based identification.

Conclusions
The diversity decreased along the different treatments, with a total of 20 genera being detected. The use of MALDI-TOF MS was reliable compared to the PhenePlate® system and has the advantage of being faster and relatively cheap compared to other technologies such as the 16S rRNA sequencing. The API web was not sufficiently reliable for water monitoring since the identification of the majority of the strains failed compared to the other methods used.
MOLECULAR DIVERSITY OF CANDIDA ALBICANS ISOLATED FROM IMMUNOCOMPROMISED PATIENTS, BASED ON MLST METHOD
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Background

Multilocus sequence typing (MLST) is used as a highly discriminatory method for tracing sources and routes of transmission, and the genetic relatedness of isolates.

Objectives

The aim of the study was to assess the genetic diversity among C. albicans strains isolated from oropharynx and bronchoalveolar lavage (BAL) of immunocompromised patients.

Methods

A total of fourteen epidemiologically unrelated clinical isolates of C. albicans from three hospitals in northern Iran were tested. Seven loci of housekeeping genes were sequenced for all fourteen isolates.

Conclusions

The fourteen isolates were placed in 10 clonal clusters (CC) while two isolates were singletons, by eBURST analysis. Most of the isolates were belong to CC461 of eBURST analysis from the clade 11 and two isolates assigned to CC172 from the clade 15.

In conclusion, Genetic diversity is variable among unrelated strains obtained from different patients at different times and places, and epidemiologically unrelated strains usually are not genetically closely related and classify in different clusters.
MULTILOCUS SEQUENCE TYPING STRAINS OF YERSINIA PSEUDOTUBERCULOSIS ISOLATED IN RUSSIA

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Background: The epidemic spread of a special clinical and epidemiological form of pseudotuberculosis, Far East Scarlet-Like Fever (FESLF) was firstly described in Russia (Far East, Vladivostok) in the late 1950s [1].

Objective: To characterize Yersinia pseudotuberculosis strains isolated in Russia (Far East, Siberia, Europe) in 1973-2014 years with the multilocus sequence typing (MLST) method to ascertain a position of the strains in a phylogenetic structure of the species.

Methods. Totally, 80 Y. pseudotuberculosis strains isolated from the stool of patients with clinical signs of FESLF (n=54), vegetables (n=11) and rodents (n=15) were included in the study. The MLST method developed by Achtman et al. (http://www2.warwick.ac.uk/mlst) was used. The MLST scheme is based on 7 housekeeping genes.

Conclusion. The majority of Y. pseudotuberculosis strains isolated from patients belonged to sequence type ST2 (85.2%, serotype 01b). Five and three clinical strains belonged to ST26 (9.3 %, serotype 01b) and ST32 (5.5%, serotype 03), respectively. Y. pseudotuberculosis serotype 01b strains isolated from vegetables belonged to ST2 (81.8%) and ST14 (18.2%). Strains isolated from rodents (01b serotype) belonged to ST2 (60%), ST42 (20%), ST64 (13.3%), ST32 (6.7%). Serotype O3 rodent isolates belonged to ST14 (5.3%). MLST results showed the Y. pseudotuberculosis ST2 serotype 01b clone prevails in Russia among all studied sources including patients, vegetables, and rodents.

PHENOTYPIC AND GENOTYPIC IDENTIFICATION OF SALMONELLA USING ANTISENSE PEPTIDE NUCLEIC ACID PROBE - MULTITYPING.

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Background
Development of novel strategies for rapid and specific identification of bacteria pathogens is urgently needed to aid successful medical and epidemiological interventions in disease outbreaks.

Objectives
In this study we evaluated the feasibility of altering growth phenotypes by silencing specific essential genes with antisense probes as a rapid bacterial diagnostic tool.

Methods
Salmonella Typhimurium and Escherichia coli strains were treated with antisense Peptide Nucleic Acid (PNA) probe targeting Salmonella ftsZ at concentrations 0-3.0 µM. Cultures were grown in non-selective Muller Hinton broth (MHB), selective Rappaport-Vassiliadis Soya Peptone (RVS), and modified Rappaport-Vassiliadis Soya Peptone-low salt (mRVSs) and cell morphologies were examined by microscopy.

Results
If ftsZ silencing is successful, cells are expected to elongate, and we here observed that treatment with the Salmonella anti-ftsZ PNA treatment elongated Salmonella grown in MHB and modified RVS low salt (mRVSs), but not in RVS. In contrast, E. coli did not elongate when treated under the same conditions. Likewise, untreated Salmonella cells did not elongate. Also, modified RVS low salt media retained Salmonella selectivity and enrichment qualities.

Conclusions
Anti-ftsZ PNA treatment can provide both phenotypic and genotypic identification of Salmonella, and the method can be combined with selective growth conditions. We are further testing this method using artificially contaminated water and milk.
DEVELOPMENT OF A QPCR TO DETECT PSEUDOMONAS AERUGINOSA IN CULTURE-NEGATIVE SAMPLES FROM CYSTIC FIBROSIS PATIENTS AND MONITOR THE EFFECTIVENESS OF ANTIBIOTIC TREATMENT

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Background

Early detection of Pseudomonas aeruginosa in cystic fibrosis (CF) samples is pivotal for disease eradication by antibiotic treatment. However, standard cultural methods are hampered by slow growth and alginate overproduction of mucoid variants, and possibly by viable but non-culturable state induction.

Objectives

This work aimed at developing a qPCR for improved P. aeruginosa infection diagnosis in sputum from CF patients and assessment of the effectiveness of antibiotic treatment.

Methods

Thirty-three sputum samples that were negative by culture assays were collected from 28 CF patients. DNA was extracted by the QIAamp DNA kit after centrifugation to remove free DNA. Real-time PCR assays targeting the species-specific gene ecfX were performed.

Conclusions

Six/33 samples were positive for P. aeruginosa by qualitative Real-time PCR (LOD: 60 cells/ml) and were analyzed by qPCR (LOQ: 3x10^2 cells/ml). P. aeruginosa abundance exceeded the LOQ in three samples, which carried 1x10^3, 1.6x10^4 and 2.8x10^6 cells/ml, respectively. Three successive samples from a culture-negative, symptomatic patient, showed bacterial counts <LOQ (May 7); 1x10^3 (still culture-negative but now receiving inhaled antibiotic therapy; May 11), and <LOQ (at the end
of antibiotic therapy; May 28). Four months later the patient was positive by cultural assays.

The matching qPCR and clinical results highlight the value of molecular assays in *P. aeruginosa* detection and quantification in CF. Dead cells cannot account for the whole difference found between PCR and culture data. Dormant forms in the pulmonary biofilm, coupled with the greater sensitivity of PCR, are likely to account for a large part of this difference.
Background

With the ever-expanding role of extended-spectrum β-lactam (ESBL) -harboring Enterobacteriaceae in causing serious infections, rapid diagnostic assays of such resistant organisms are highly desired to improve patient screening and hospital infection control practices as well as to reduce inappropriate antibiotic use.

Objectives

To reach this goal, we developed an amperometric detection of the β-lactamase activity in blood culture samples using disposable carbon screen-printed sensors in the presence of a well-chosen cephalosporin substrate.

Methods

Using an ESBL enzyme isolated from a clinical strain, we showed that the intensity of a specific anodic peak current resulting from the catalytic hydrolysis of the β-lactam ring was proportional to the amount of ESBL. A novel susceptibility assay for the rapid and specific identification of ESBL-producing bacteria in blood culture samples was then proposed. This assay was based on a two-step protocol performed within 2 hours : (1) subculturing of blood culture samples in the presence or absence of cefotaxime and/or the potassium clavulanate (ESBL inhibitor) for a few hours followed by, (2) incubation of the subculture filtrates with the redox substrate which hydrolysis was monitored by amperometry. This assay allowed the reliable and specific identification of various types of β-lactamases-producing bacterial species (including ESBLs, penicillinases, cephalosporinases) in blood cultures.

Conclusions

Owing to its low cost, portability, simplicity and its ability to perform measurements in turbid media, the electrochemical approach, which did not require prior strain isolation, holds great promise for the rapid screening of β-lactamases in clinical and other complex samples.
FEMS-2135
New diagnostic approaches

RAPID AMPEROMETRIC DETECTION OF ESCHERICHIA COLI IN WASTEWATER BY MEASURING β-D GLUCURONIDASE ACTIVITY WITH DISPOSABLE CARBON SENSORS
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Background
Methods based on the direct measurement of β-D-glucuronidase (GLUase) activity in the presence of the 4-methylumbelliferyl-β-D-glucuronide substrate are well-established to specifically monitor Escherichia coli (E.coli) in water samples without any cultivation step. However, because of the interferences from auto-fluorescence or light-quenching particle in complex samples, the fluorescence detection is often not suitable for the analysis of environmental samples.

Objectives
To overcome this drawback, the amperometric quantification of GLUase activity was investigated and applied to the rapid and specific detection of E. coli in wastewater samples.

Methods
For this purpose, the p-aminophenyl β-D-glucopyranoside (PAPG) was selected as electrochemical substrate for GLUase measurement and the p-aminophenol (PAP) released during the enzymatic hydrolysis was monitored by cyclic voltammetry with disposable carbon screen-printed sensors. The intensity of the measured anodic peak current was proportional to the amount of GLUase, thus providing a measurement of the number of E.coli in the sample (using a calibration curve). Following an optimized protocol, we were able to detect E. coli cells in the range of 5 x 10⁴ to 10⁸ per filter within 2 h. The amperometric assay was applied to the determination of fecal contamination in raw and treated wastewater samples and it successfully compared with conventional bacterial plating methods and uidA gene detection by quantitative PCR.

Conclusions
Owing to its ability to perform cell density measurements in turbid media, the GLUase amperometric method is a reliable tool for the rapid and decentralized quantification of viable but also non-culturable E.coli in complex environmental samples.
New diagnostic approaches

RAPID DETECTION OF SALMONELLA SPP. USING AN ISOTHERMAL AMPLIFICATION BASED SYSTEM, DNABLE

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Background

The rapid, cost-effective, and robust detection of pathogens continues to be a major focus for the livestock industry. While PCR has filled the niche for a highly sensitive detection, PCR inherently burdens the end user with cumbersome sample preparation, complex and closed instrumentation with long time–to-result (TTR) (1.5-3h post-enrichment, excluding sample preparation). Other DNA amplification methods have shortened the TTR but have not been able to provide a robust and reliable alternative to PCR. The need for improved testing solutions continues to grow with growing intricacies of production logistics.

Objectives

Here we present an isothermal DNA amplification tool called DNAble for Salmonella spp. detection. This system provides a distinctive advantage over other technologies as it combines high specificity and sensitivity with minimal sample preparation leading to a significantly reduced TTR (30-40 min post-enrichment, including sample preparation).

Methods

Salmonella detection was demonstrated with an analytical sensitivity of <50cp of genomic DNA per reaction or <10 cells post-enrichment in multiple test matrices. Inclusivity and exclusivity data indicate 100% accuracy across 34 Salmonella serotypes and 29 non-Salmonella tested strains.

Conclusions

DNAble requires minimal sample preparation due to its robust chemistry and is compatible with a range of detection equipments. Salmonella spp. detection system also includes an internal amplification control, Salmonella selective media supplement that improves current enrichment, and a user-friendly sample preparation kit. Results are provided by a portable and simple DNAble reader or user provided thermocycler. The inherent robustness, quick TTR, and broad range instrument compatibility support use of DNAble for Salmonella detection.
Background

Fecal pollution of water can lead to health problems due to presence of infectious microorganisms especially *Enterococcus faecalis* (*E. faecalis*) which may be derived from human sewage or animal sources. Hence, detection of these microorganisms is of much interest using different methods. Electrochemical DNA biosensors, which employ an immobilized DNA as the biological recognition element, are currently under intense investigation in detection of several microorganisms relying on the conversion of the base-pair recognition event into a useful electrical signal.

Objectives

The aim of this study was to demonstrate a rapid and inexpensive method for accurately detection of *E. faecalis* in contaminated water via fabrication of a specific and selective DNA electrochemical biosensor.

Methods

A modified DNA oligonucleotide with 5’ amino modifier C12 was designed based on a house keeping gene glucose 6-phosphate dehydrogenase of *E. faecalis*. The multi walled carbon nanotubes (MWCNT) were allowed to form amide bond between their carboxylic acid groups and amino groups of oligonucleotides. Hybridization was performed by incubating the DNA/self-assembled MWNTs with complementary DNA oligonucleotides, non-complementary DNA oligonucleotides, genomic DNA and artificial contaminated water under specific conditions for 30 min at room temperature. The hybrid-electrodes were successively transferred into hybridization indicator of methylene blue for 30 min without any potential followed by several washings and subsequently electrochemical measurements.

Conclusions

Such single-use electrochemical biosensors have great promise for decentralized water testing for presence of *E. faecalis* in an accurate and rapid approach.
Background

Acute exacerbations of chronic obstructive pulmonary disease (AE COPD) are predominantly caused by microbial pathogens. Molecular techniques may improve the speed and sensitivity of respiratory infections diagnostics. Currently there is limited knowledge on the best method for DNA extraction from sputum samples.

Objectives

This study is a part of the European JRP HLT08 INFECT-MET. The aim was to evaluate the DNA yield of two different extraction methods, manual and automated, in treated and untreated sputum samples from COPD patients.

Methods

A total of 47 good-quality sputum samples from COPD patients between March 2014 and October 2014 were included in the study. Twenty samples were left untreated and 27 were treated with NALC-NaOH. All samples were split in two aliquotes and DNA was extracted with manual (QIAamp DNA Mini Kit, Qiagen GmbH, Germany) and automated (DNA Extraction Kit, DiaSorin, Italy/NorDiag Arrow) method. Concentration of DNA was quantified (Qubit Fluorometer, Invitrogen, USA).

Conclusions

Out of 47 sputum samples 41 (87,2%) had higher DNA concentration with automated extraction method. Among 20 untreated sputum samples automated method had a higher DNA yield in 14 samples by 68,1%, lower in 5 by 43,1 % and equal in 1 sample. In 27 treated sputum samples automated extraction method had a higher DNA yield in all 27 samples by 60,9%.
Automated extraction method proved to be more efficient in extracting DNA from untreated and treated sputum samples. Automated extraction method is much easier to perform and less time consuming.
DEVELOPMENT OF DUAL-FUNCTION ELISA FOR EFFECTIVE ANTIGEN AND ANTIBODY DETECTION AGAINST H7 AVIAN INFLUENZA VIRUS

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¹Animal Health Biotechnology, Temasek Life Sciences Laboratory, Singapore, Singapore

Background

Outbreaks in poultry involving influenza virus from H7 subtype have resulted in human infections, thus causing a major concern for public health, as well as for the poultry industry. Currently, no efficient rapid test is available for large-scale detection of either antigen or antibody of H7 avian influenza viruses.

Objectives

In the present study, a dual function ELISA was developed for the effective detection of antigen and antibody against H7 AIVs.

Methods

The test was established based on antigen-capture-ELISA and epitope blocking ELISA. The two Mabs 62 and 98 which were exploited in the assay were identified to recognize two conformational neutralizing epitopes on H7 HA1. Both of the epitopes exist in all of the human H7 strains, including the recent H7N9 strain from China and > 96.6% of avian H7 strains. The dual ELISA was able to detect all of the five H7 antigens tested without any cross reaction to other influenza subtypes. The antigen detection limit was less than 1 HA unit of H7. For antibody detection, the sensitivity and specificity of the dual ELISA was evaluated and compared to HI and microneutralization using immunized animal sera to different H7 strains and different subtypes of AIVs. Results indicated that antibodies to H7 were readily detected in immunized animal sera by the dual ELISA whereas specimens with antibodies to other AIVs yielded negative results.

Conclusions
This is the first dual-function ELISA reported for either antigen or antibody detection against H7 AIVs. The assay was highly sensitive and 100% specific in both functions rendering it effective for H7 diagnosis.
NEOPTERIN AND SOLUBLE CD14 LEVELS IN CASES WITH UNDETERMINATE AND CONFIRMED HIV-1 INFECTIONS RELATED WITH HIV-1/2: DOES UNDETERMINATE HIV-1 PATTERN SHOW A REAL HIV-1 INFECTION?

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Background

HIV-1/2 testing algorithms are currently controversial. The most important problem and difficulty about this debate is undeterminate HIV-1 infections (UHIV-1). Studies for an optimised algorithm for UHIV-1 are under research.

Objectives

In this study, we aimed to determine neopterin and sCD-14 levels in cases with UHIV-1 and confirmed HIV-1 (CHIV-1) infections and evaluate the relationship between UHIV-1 and real HIV-1 infection and also aimed to evaluate the diagnostic values of two markers, especially in situations when PCR were not available and western blot were undeterminate after ELISA.

Methods

Eighty-eight cases with UHIV-1, 100 patients with CHIV-1 and 100 healthy control group (HCG) were included. Neopterin and sCD-14 levels were determined by competitive and sandwich ELISA (nmol/mL, µg/mL, quantitatively) methods, respectively. Statistical tests were performed using SPSS 21.0 and p < 0.05 was considered significant.

Conclusions

Results:
Table 1. Neopterin and sCD-14 levels of UHIV-1 and CHIV-1 cases and healthy control group

<table>
<thead>
<tr>
<th>Markers</th>
<th>Undeterminate HIV-1¹ (n:88)</th>
<th>Confirmed-HIV-1² (n:100)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HCG³(n:100)</td>
<td></td>
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</tr>
</tbody>
</table>
Neopterin, mean (min-max)  5.73 (1.58-38.71)  15.77 (1.19-127.19)  7.95 (3.16-41.08)
sCD14, mean (min-max)  3.65 (1.86-6.17)  4.20 (1.53-7.59)  3.51 (0.97-3.97)

P value: Neopterin: 1x2; p:0.004, 1x3; p:0.754
sCD14: 1x2; p:0.034, 1x3; p:0.787

Table 2. Diagnostic test performances of neopterin and sCD14 in CHIV-1 infections

<table>
<thead>
<tr>
<th>Markers</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
<th>Kappa coefficient</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neopterin (%)</td>
<td>38</td>
<td>82</td>
<td>76</td>
<td>57</td>
<td>20</td>
</tr>
<tr>
<td>sCD14 (%)</td>
<td>50</td>
<td>100</td>
<td>100</td>
<td>67</td>
<td>50</td>
</tr>
</tbody>
</table>

As conclusion,, we couldn’t suggest UHIV-1 pattern as real HIV-1 infection and we concluded that two markers were useless as surrogate markers for real HIV-1 infections.
Background

Isolated Anti-HBc pattern (IAH) positivity causes diagnostic difficulty in laboratory diagnosis of HBV infections. In addition, this pattern is a big problem for clinical diagnosis and blood transfusion.

Objectives

In this study, we aimed to determine neopterin and sCD-14 levels in cases with IAH and chronic active hepatitis (CAH) and to evaluate the relationship between IAH and real HBV infection and also aimed to evaluate the diagnostic values of two less sophisticated and cheaper surrogate markers in situations when the molecular methods were not available.

Methods

IAH (n:102), KAH (n:70) and healthy control group (HCG), (n:100) cases were included in the study. Neopterin and sCD-14 levels were determined by competitive and sandwich ELISA (nmol/mL, µg/mL, quantitatively) methods, respectively. Statistical tests were performed using SPSS 21.0 and p < 0.05 was considered significant.

Conclusions

Table 1. Neopterin and sCD-14 levels in IAH, KAH and HCG cases.

<table>
<thead>
<tr>
<th>Markers</th>
<th>IAH</th>
<th>KAH</th>
<th>HCG</th>
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</table>
Neopterin, mean (min-max) (n:102) 8.67 (3.56-67.15) (3.16-41.08) 15.43 (4.75-43.27) 7.95
sCD14, mean (min-max) (n:70) 3.61 (1.82-6.17) (0.97-3.97) 4.20 (1.96-6.78) 3.51

p value: Neopterin: 1x2; p:0.002, 1x3; p:0.915
sCD14 : 1x2; p:0.006, 1x3; p:0.821

Table 2. Diagnostic test performances of neopterin and sCD14 in real HBV infections

<table>
<thead>
<tr>
<th>Markers coefficient</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>PPV</th>
<th>NPV</th>
<th>Kappa</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neopterin (%)</td>
<td>59</td>
<td>82</td>
<td>69</td>
<td>75</td>
<td>42</td>
</tr>
<tr>
<td>sCD14 (%)</td>
<td>51</td>
<td>90</td>
<td>78</td>
<td>73</td>
<td>44</td>
</tr>
</tbody>
</table>

As conclusion, we couldn’t suggest IAH pattern as a real HBV infection and we concluded that two markers were useless as surrogate markers for the diagnosis of HBV infections.
DETECTION OF THE OOMYCETE PYTHIUM INSIDIOSUM BY REAL-TIME PCR
OF THE EXO-1,3-B-GLUCANASE GENE

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Background
Pythiosis is a life-threatening infectious disease caused by the pathogenic oomycete
Pythium insidiosum. The disease has high rates of morbidity and mortality. Timely
diagnosis permits prompt treatment and a better prognosis. PinsEXO1, encoding an
exo-1,3-β-glucanase, is a novel and efficient target for identification of P. insidiosum
by conventional PCR.

Objectives
In this study, we aimed to develop a real-time PCR approach targeting PinsEXO1
and compare its performance to conventional PCR for the detection of P. insidiosum.

Methods
Genomic DNA samples were prepared from 35 culture-proven P. insidiosum isolates
and 58 culture-proven fungi (served as the control), for PCR analyses. Both
conventional and real-time PCR assays were positive for all P. insidiosum strains
tested, while all control fungi were negative. Turnaround time for conventional PCR
was 4 hr, while that of real-time PCR was 1.5 hr. The minimum DNA template
required for successful PCR amplification by conventional and real-time PCR were 1
ng and 1 x 10⁻⁴ ng, respectively.

Conclusions
The real-time assay retained high detection sensitivity and specificity. It showed a
substantially improved analytical sensitivity and turnaround time that could improve
diagnoses of pythiosis.
DEVELOPMENT OF A NOVEL MALARIA ANTIBODY ASSAY UTILIZING ANTIGENS FROM ALL 5 HUMAN PATHOGENIC PLASMODIUM SPECIES

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Background

The proper diagnosis of Malaria disease is essential to provide early treatment and improve the prognosis of patients. Transfusion-transmitted Malaria is rare, but it may produce severe problems in the safety of blood Transfusion and blood related products due to the lack of reliable procedure to evaluate donors potentially exposed to malaria.

Objectives

Microscopy, still considered the gold standard for diagnosing malaria. It is time consuming and requires trained expertise. ELISAs are known to be ideal for high throughput screening with high sensitivity and specificity, but it also requires trained personal and an equipped laboratory. Line Blots are often used as confirmatory tests since they provide high sensitivity and specificity. There is nearly no lab equipment needed to perform this kind of assay. In addition, blots can also be used in automated processes for high throughput screening.

Methods

Here we show an improved diagnostic performance of the new antibody detection Systems (ELISA and Lineblot) utilizing early and late antigens of all 5 human pathogenic Plasmodium species (P. falciparum, P. vivax, P. ovale, P. malariae, P. knowlesi) compared to test systems only relying on antigens derived from one or two Plasmodium species. Assays with a limited number of antigens often fail to detect antibodies from certain regions of the world. For evaluation purpose, we collected samples from all over the world, including samples from newborns.

Conclusions

We evaluated the performance of ELISA and Lineblot directly in endemic countries with samples of patients who presented symptoms akin to malaria infection in local hospitals.
INVESTIGATION OF ORTHOPAEDIC IMPLANT ASSOCIATED INFECTIONS: ALTERNATIVE APPROACHES TO LABORATORY PROCESSING AND CULTURE

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Background

Since its introduction in the 1960s, joint replacement (arthroplasty) has become a common procedure. Around 15% of revisions are due to ‘septic’ loosening caused by biofilm formation on the implant. Infection rates are now much lower; however, there is still a risk associated with each procedure. UK SMI B44: is a diagnostic tool aimed at practising professionals in the field of microbiology. It describes and recommends methods for the microbiological investigation of orthopaedic implant associated infection.

Objectives

The aim of this poster is raise awareness of the various methods of sample preparation (including the use of glass beads and sonication) and culture. Timely diagnosis and treatment can help reduce the likelihood of systemic dissemination of infection. Therefore, in most cases, this UK SMI recommends up to 5 days culture using either cooked meat broth or continuous monitoring blood culture system.
Methods

Produced by the Standards Unit, UK SMIs are developed, reviewed and updated through a wide consultation process with users and other stakeholders. The process follows the AGREE tool and resulting documents reflect best evidence based practice. Where evidence is not available, the documents are based on national working group consensus decisions.

Conclusions

B44 provides guidance on the best minimum practice when investigating orthopaedic implant associated infection. It is a useful resource for laboratories which aims to help drive pathology modernization by recommending alternative technologies for timely laboratory diagnosis of infection.
MASS SPECTROMETRY PROTEOTYPING FOR THE DETECTION,
CHARACTERISATION AND IDENTIFICATION OF INFECTIOUS BACTERIA
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E. Kristiansson⁶, E. Moore⁷
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Background

Diagnosing infectious diseases requires a range of methodologies for the comprehensive detection, characterisation and identification of infectious bacteria. Proteomics-based approaches represent an alternative to traditional methods of characterising microorganisms, elucidating expressed biomarkers of genome information, applied to ‘proteotyping’ microorganisms at all taxonomic levels.

Objectives

Apply state-of-the-art and novel techniques of tandem mass spectrometry (MS) analyses of expressed proteins for species- and strain-level identification and detection of antibiotic resistance and virulence, with the focus on optimizing MS-based proteomics analyses for ‘proteotyping’ infectious bacteria.

Methods

Intact bacteria or cell fractions were immobilised, via membrane-gold interactions within a flow-cell. Within the flow-cell, proteins were subjected to enzymatic digestion to generate peptides, which were identified, using LC-MS/MS. Following database matching, peptides were used for bacterial species-level identification, and for detection of antibiotic resistance and sub-species typing. To demonstrate proteotyping capability and differentiate closely related species, mixtures of different Streptococcus spp. were analysed. Results of Streptococcus spp. identifications from clinical samples were confirmed by standard microbiology, including cultivation of bacteria in selective media, PCR, DNA sequencing and MALDI-TOF MS analyses.

Conclusions

Proteotyping of infectious bacteria, using LC-MS/MS enabled the differentiation and identification of Streptococcus spp. by ranking identified expressed proteins, according to the number of peptide matches to genome sequence information available in databases, and was applied to detection in clinical samples. The methods
have been shown to enable reproducible, sensitive, rapid and cost-effective typing of bacteria for diagnostics of infectious diseases.
APPLICATION OF MOLECULAR TECHNIQUES TO THE RAPID DETECTION AND CONFIRMATION OF HIV

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Background
Western Blot (WB) is the gold standard as HIV-1 confirmatory test, which has high cost and complexity in processing, our setting send the reactive samples to a national reference center for confirmation and reporting of results takes at least 15 days.

Objectives
The aim of this work was to standardize a technique multiplex PCR as a complementary test for the detection and confirmation of HIV-1 infection in Oaxaca, Mexico.

Methods
Standardization of the multiplex PCR was perform using primers designed to amplify the gag, pol and env genes of M, N, O groups and other specific to Subtype B. For the standardization, we use DNA of three patients that was previously characterized as HIV positive and samples sequenced for HIV-1 (pol, gag, env). We used this multiplex PCR in DNA samples from PMBC of 600 patients. All patients were screening by HIV rapid test. To confirm HIV infection, both WB and multiplex PCR were perform.

Conclusions
We obtain concordant results with multiplex PCR and those of WB. The implementation of a multiplex PCR as a supplemental test for confirmation of HIV-1 infection is reliable and safe. Its implementation has the possibility of decreasing the time of detection and reporting of confirmatory results to a day, which is essential, for example, in cases of pregnant women the confirmation or discard of HIV must be timely, with the aim to start prophylaxis on time.

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INFLUENCE OF STORAGE CONDITIONS ON THE SPECTRAL STABILITY OF BACTERIAL SUSPENSIONS IDENTIFIED BY MALDI BIOTYPER

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Background

Although direct smear technique is an excellent approach for the preparation of samples for MALDI-TOF MS identification, the usefulness of this approach is limited to BSL-1/BSL-2 bacteria, it is not recommended for MSP creation and it requires fresh material for an optimal performance. Formic acid extraction is proposed as a more precise method for MALDI Biotyper identification that includes a step in which bacteria are suspended in 70% ethanol which could potentially be used for the storage of bacterial suspensions in their optimal growth phase. However, there is a lack of information regarding the impact of storage of bacterial suspensions in ethanol on the reliability of the MALDI-Biotype identification.

Objectives

A study was carried out aiming to assess the influence of time (90 days) and temperature (room temperature and 4°C) on the identification by MALDI Biotyper.

Methods

Two bacterial suspensions (E. coli and S. aureus) stored in ethanol 70% were included. Half of the aliquots (n=27) were conserved at room temperature and the other half at refrigeration until the day of analysis (0, 5, 8, 12, 19, 26, 42, 71 and 90 days).

Conclusions

Our data suggest that E. coli and S. aureus ethanol suspensions can be stored up to 90 days for MALDI Biotyper identification in laboratory conditions. Although it has to be validated, it may apply to other bacteria and could greatly facilitate laboratory work-flow and shipment of bacterial suspensions.
Background

To elucidate the molecular mechanisms involved in persistency/latency of the H. pylori infection or in its progression towards serious diseases, it is necessary to analyze the host pathogen interaction in vivo. The circulating antibody repertoire represents an important source of diagnostic information, serving as biomarker to provide a ‘disease signature’.

Objectives

The aim of this work is the identification of H. pylori epitopes responsible for host immunoresponse modulation though a discoverydriven approach that couples ‘phage display’ and deep sequencing.

Methods

We used an approach for identifying novel antigens by screening gDNA libraries created from the pathogen genome, directly with sera from infected patients. Three phage display libraries from three H. pylori strains (HP26695, HPB128, HPSS1) have been constructed by using βlactamase ORF selection vectors (Di Niro et al., 2010). Genomic DNA was sonicated, fragments cloned into the filtering vector, after transformation libraries of 1x10^6 clones were obtained and sequenced by 454 technology.

Conclusions
More than 93% of HP CDSs were represented in the phage genomic library therefore being representative of the whole *H. pylori* antigenic ORFeome. Putative antigens were selected from libraries using sera from patients affected by *H. pylori* presenting increasing degrees of infection: i) autoimmune gastritis and pernicious anemia; ii) gastric adenocarcinoma; iii) MALT lymphoma. The results show that the diversity of the libraries obtained after selection is significantly reduced. Furthermore, individual ranks, for each infection condition, have been compared highlighting the pattern of putative antigens, shared by all the conditions, and some that can distinguish the different stages of infection.
DETECTION OF CRYPTOSPORIDIUM OOCYST RNA BY SURFACTANT EXTRACTION TREATMENT AND RT-PCR: INHIBITION OF REVERSE TRANSCRIPTION BY SDS AND ITS SUPPRESSION USING NONIONIC SURFACTANTS

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Background
Cryptosporidium causes the symptoms of cryptosporidiosis including watery diarrhea, vomiting and fever. Cryptosporidium oocysts possess a robust wall, which is extremely resistant to chlorine used for potable water disinfection. Successful DNA detection from the robust oocyst usually requires complicated procedures including DNA extraction such as freeze-thaw cycling and enzyme treatment. Therefore, we previously developed a surfactant extraction treatment (SET) method for extracting DNA from the oocysts with only sodium dodecyl sulfate (SDS).

Objectives
We examined the inhibition of reverse transcription-PCR (RT-PCR) by SDS and its suppression using four kinds of nonionic surfactants in order to develop a new method for detecting 18S rRNA from the oocyst by SET and RT-PCR.

Methods
The template DNA/RNA was amplified in the presence of SDS and/or the nonionic surfactants; Tween 20, Tween 80, Triton X-100, Triton X-114 by RT-PCR to evaluate the inhibition of reverse transcriptase and DNA polymerase. The oocysts were incubated in a 0.1% SDS solution to extract DNA/RNA.

Conclusions
DNA/RNA amplification was inhibited at 0.01% SDS but its inhibition was suppressed by adding any of the nonionic surfactants at 5%. There was a difference in the suppression efficacy of the nonionic surfactants in tests that used 0.1% SDS. 18S rRNA molecules are present in high copy numbers in viable cells, which increases detection sensitivity. Therefore, we succeeded to detect 18S rRNA equivalent to $10^{-4}$ oocysts by SET and RT-PCR. This work was supported in part by JSPS KAKENHI Grant Number 25420559 and the Kurita Water and Environment Foundation.
NEW DIAGNOSTIC APPROACHES IN INFECTION DETECTION BASED ON THE IMMUNE SYSTEM DERIVED ENZYMES IN WOUND FLUID

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Background

Quick and proper wound diagnostics could avoid wound infections of currently 5-10% of post-surgical - and 25% of chronic wound patients. Apart from standard methods, this study focusses on the wound fluid, harboring enzymes of the human immune system and reflecting a critical wound status upon increasing activity.

Objectives

The determination of the enzyme levels of the neutrophil derived enzymes human neutrophil elastase (HNE), myeloperoxidase (MPO) and lysozyme (LYS) was assessed and correlated to the bacterial burden of the wounds [1]. New diagnostic approaches were based on these enzymes and include electrochemical and optical sensor systems for fast diagnosis of infections in chronic wounds [2].

Methods

The enzyme activities of HNE, LYS and MPO were directly monitored in wound fluid of affected patients via biochemical investigations. Both electrochemical sensors such as measuring the hydrogen peroxide consumption by MPO as current decrease and lateral flow devices based on release of dyes were developed.

Conclusions
Determination of the enzyme activities showed more than 6 to 10 fold higher substrate conversion in infected wound fluids when compared to non-infected wound fluids and clearly correlated with the elevated bacterial burden (Fig1). Consequently, both the electrochemical sensor as well as lateral flow devices were able to differentiate between infected and non-infected wounds.


Molecular surveillance on nasopharyngeal carriage of Streptococcus pneumoniae in children vaccinated with conjugated polysaccharide pneumococcal vaccines


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Background
As a commensal of the upper respiratory tract, Streptococcus pneumoniae is a potential pathogen causing respiratory and invasive diseases. Following implementation of pneumococcal conjugate vaccination (PCV) for infants, surveillance studies have proven essential for monitoring direct (carriage of serotypes targeted by vaccine, VTs) and indirect effects (changes in carriage of non-vaccine serotypes, NVTs).

Objectives
To compare the detection of pneumococcal carriage and serotypes in a unique study setting using both conventional culture and molecular methods, in nasopharyngeal samples from healthy PCV-vaccinated infants in two large, cross-sectional surveillance studies on PCV-effects in the Netherlands.

Methods
Nasopharyngeal samples were collected from 1182 11- and 24-month old children (n=591 each) during autumn/winter 2010/11 (n=584) and 2012/2013 (n=598). Following conventional culture on plates selective for S. pneumoniae, DNA extracted from all bacterial growth was tested by quantitative-PCR (qPCR) for the presence of pneumococci and a panel of serotypes, including serotypes targeted by the thirteen-valent PCV (PCV13).

Conclusions
There was a correlation (Spearman's rho=0.980; p<0.001) between the frequency of serotypes detected using qPCR and prevalence according to conventional culture. Moreover, we observed a trend (p=0.056) for an underestimation of carriage detection for serotypes not targeted by PCV7 or PCV10 (in use in the Netherlands) compared to PCV10-serotypes when detected using the culture-method alone. We found no evidence of a hidden circulation of serotypes rarely detected by culture or those targeted by vaccination. This suggests that surveillances based on the culture method alone do not underestimate carriage of VTs in immunised children.
Background

In the field of natural product research, finding new sources of bioactive compounds is of primary importance. In this respect, microorganisms have provided a large number of biologically active molecules.

Objectives

Recently, the use of fungal co-cultures for the induction of new natural products has emerged as a promising field in drug discovery. For the success of such studies, a key element is the development of a co-culture methodology that provides high reproducibility of metabolite induction patterns and that is compatible with high throughput analytical procedures.

Methods

To tackle this issue, a method based on 12-well-plate miniaturized Petri dishes compatible with high throughput UHPLC-TOF-MS metabolomics [3] has been developed. This strategy was used to screen for metabolite induction in co-cultures of various fungal species. This approach provided a satisfactory reproducibility and was used for the identification of induced biomarkers.

Conclusions
This study demonstrates the consistent induction of new metabolites through co-culture. Moreover, the developed strategy is generic and can be applied to other types of microorganisms that can grow on solid media and that are part of the myc- or microbiome. This 12 well-plate approach and the adapted data mining strategy were validated by the untargeted metabolomic study of a model co-culture (Eutypa lata versus Botryosphaeria obtusa responsible for confrontation zone lines in Vitis wood). This procedure is currently used for screening novel metabolite induction in various fungal co-cultures.
Background
Cyclodipeptide synthases (CDPSs) constitute a novel family of enzymes that use charged tRNAs to synthesize cyclodipeptides in biosynthetic pathways dedicated to the synthesis of diketopiperazines, a large class of secondary metabolites with noteworthy biological activities. To date, 11 CDPSs have been biochemically characterized, showing the incorporation of hydrophobic amino acids in cyclodipeptides (essentially Phe, Leu, Tyr and Trp). Biochemical and structural characterizations of three CDPSs identified catalytic residues and allowed to propose a ping-pong mechanism for cyclodipeptide formation. However, bioinformatic searches in databases identified in microbial genomes numerous putative CDPS whose activities are not predictable. Furthermore, some of these CDPSs showed differences in predicted catalytic residues.

Objectives
Our objective was to provide a better characterization of the CDPS family of enzymes notably by determining the cyclodipeptide-synthesizing activity of a large set of unrelated putative CDPSs identified in databases.

Methods
We used bioinformatic tools to select the CDPS candidates for biochemical characterization and to analyze their amino acid sequences. As previous work showed the recovery of cyclodipeptides in culture supernatants upon CDPS expression in *Escherichia coli*, CDPSs were produced in this host and cyclodipeptides were searched in culture supernatants by HPLC coupled to mass spectrometry.

Conclusions
The determination of the activities of 41 novel CDPSs enlarged the chemical diversity synthesized by these enzymes, as they were shown to incorporate 17 of the 20 proteinogenic amino acids. CDPSs can be classified into several phylogenetically distinct subfamilies characterized by specific functional subsequence signatures, suggesting differences in catalysis by members of the two subfamilies.
IDENTIFICATION OF ANTIMICROBIAL LIPOPEPTIDES OF BACILLUS STRAINS OBTAINED BY DIFFERENT WAYS OF EXTRACTION USING MALDI-TOF MASS SPECTROMETRY

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Background

Bacillus cyclic lipopeptides of surfactin, iturin, fengycin, and recently discovered kurstakin families have well-recognized potential in biotechnology and biopharmacies. Analytical method such as matrix-assisted laser desorption ionization-time of flight mass (MALDI-TOF) spectrometry proved very effective for detection and identification of various antimicrobial substances.

Objectives

The present study was aimed at investigating how different mode of extraction of crude lipopeptides produced by different Bacillus strains influence the amount of crude lipopeptide and highlights the best extraction mode for further use.

Methods

Five proven lipopeptide producers were grown under previously determined conditions and consequently extracts were obtained by ethyl acetate extraction and by combination of acid precipitation and methanol extraction. Also, cell-free supernatants in the form of aqueous extracts were examined. Extracts were subjected to MALDI-TOF-MS analysis in the acquisition mass range 800-1700 Da.

Conclusions

Mass spectra obtained from all extracts tested, showed clearly three peak clusters corresponding to different lipopeptide families. Cluster of peaks, corresponding to iturin and surfactin families, with similar intensities and points to the dominant presence of these compounds in all strains and extracts, was observed. However, for peaks within the mass range of 1450-1540, corresponding to fengycin family, the best detection was achieved for ethyl acetate extracts. The peaks in the mass range of kurstakin family, for most cases, were more pronounced in the aqueous extract, while the lowest detection was obtained for methanolic extraction. In conclusion, the most efficient was ethyl acetate extraction, which gives the most prominent peaks together with the aqueous extract as a positive control.
Background

Antibiotic-associated hemorrhagic colitis (AAHC) develops during antibiotic-driven intestinal dysbiosis and is caused by production of the bacterial pyrrolobenzodiazepine tilivalline.

Objectives

To understand the function of this secondary metabolite during health and disease we study the regulation of tilivalline biosynthesis and degradation.

Methods

Genes involved in tilivalline biosynthesis were identified via transposon mutagenesis of the clinical isolate AHC-6. Genes encoding a 3-deoxy-D-arabino-heptulosonate-7-phosphate (DAHP) synthase (aroX), a dehydroquinate-synthase (aroB) and two non-ribosomal peptide synthases (npsA and npsB) were identified as essential genes for tilivalline production. Adjacent genes npsA, npsB and aroX are located on a pathogenicity island present in all toxin positive Klebsiella isolates.

Under laboratory conditions tilivalline is detected in conditioned medium of bacterial cultures. Cytotoxicity towards human epithelial HeLa cells appears at the end of the exponential phase (8 h), reaches a maximum during stationary phase and decreases after 40 h to undetectable levels at 48 h. The loss of cytotoxicity can indicate a growth-dependent regulation of biosynthesis and/or inactivation of tilivalline. Emergence of degradation products in tilivalline-depleted supernatant was analyzed by HPLC-MS and NMR. Regulation of tilivalline biosynthesis genes in vitro was monitored via qRT-PCR.

Conclusions

We find that transcription of npsA and npsB is regulated in a growth dependent
manner. Current studies focus on effects of host and microbial factors on tilivalline biosynthesis and cytotoxicity.
BURKHOLDERIA GENOME MINING FOR NONRIBOSOMAL PEPTIDE SYNTHETASES REVEALS A GREAT POTENTIAL FOR NOVEL LIPOPEPTIDES SYNTHESIS

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Background

Many of natural compounds produced by microorganisms, are synthesized by non-ribosomal peptides synthetases (NRPS), working as assembly lines. Among them, lipopeptides as those produced by Bacillus and Pseudomonas, may play an important role in sustainable agriculture, used as biopesticides to kill plant pathogens.

Objectives

As lipopeptides are synthesized by NRPS, we have decided to screen and identify in silico all NRPS gene clusters of burkholderia, especially those producing siderophores and lipopeptides, potentially implicated in biocontrol of phytopathogens.

Methods

We performed in silico detection and analyses of NRPS genes present in 48 gapless complete genomes of Burkholderia available in the NCBI, by following Florine, a workflow we have developed especially with this aim.

Conclusions

Genome analyses revealed 161 clusters containing NRPSs, with the potential to synthesize at least 11 novel products. Although most strains produce the main siderophore ornibactin or malleobactin, a cluster corresponding to a new siderophore, called phymabactin, was identified in Burkholderia phymatum STM815 and the cluster for cepaciachelin biosynthesis was, for the first time, identified in Burkholderia ambifaria AMMD. Elsewhere, the cluster for the antifungal burkholdin was detected in both the genomes of B. ambifaria AMMD and Burkholderia sp. KJ006. We also
identified a novel lipopeptide called burkhomycin specifically produced by *Burkholderia pseudomallei* strains. This study revealed the significant of the genus *Burkholderia* as a promising source of bioactive compounds. It also gave new insights on the non-ribosomal synthesis exemplified by the identification of dual C/E domains in lipopeptide NRPSs, as frequently found in *Pseudomonas* strains.
FEMS-1299
Secondary metabolites, metabolomics

BACTERIAL TRIGNONELLINE METABOLISM, FROM THE GENES TO THE METABOLITES: A STEP FORWARD INTO THE METABOLIC LANDSCAPE OF BACTERIA
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Background
Trigonelline (N-methylnicotinate), a pyridine alkaloid, is produced by plants to resist drought stress. Its catabolism by soil bacteria is presumed to proceed through methylamine and succinate but no pathway is described until now.

Objectives
Our model organism is Acinetobacter baylyi ADP1, a strictly aerobic, nutritionally versatile soil bacterium which is easily amenable to genetic manipulations. Biological tools and resources have been developed in the laboratory to enable the integrative analysis of its metabolism. Since it grows well on trigonelline, we wanted to elucidate this pathway at the finest level.

Methods
By comparative genomics, we could identify a conserved cluster of 10 genes involved in trigonelline breakdown. We dissected the pathway by combining bacterial physiology with mutant strains, classical enzymology with recombinant enzymes and untargeted LC/MS based metabolomic approaches.

Conclusions
In reaction mixtures with trigonelline, we could detect succinate semialdehyde and succinate along with three new metabolites, which were produced on a preparative scale, purified and structurally characterized by 1D- and 2-D NMR. Unexpectedly, the first intermediate exhibited a rare 5-hydroxy-butyrolactone structure in equilibrium with an open form, which resulted from a successive reduction-oxidation and cleavage of the pyridine ring followed by a recyclization. This reaction scheme remains unprecedented for a natural pyridine compound and may have valuable spin-off effects in the field of bioremediation since pesticide derivatives or pyridine itself may also be catabolized by this way.
In addition, we showed that trigonelline metabolism is connected with the metabolism of compatible solutes in the cell, extending its physiological function in bacteria beyond nutrition.
Background

Soil microorganisms produce a range of secondary metabolites like antibiotics, toxins, biosurfactants, siderophores. Beside these metabolites soil microorganisms are capable of emitting another class of secondary metabolites so called volatile organic compounds (VOCs). Volatiles are low molecular mass compounds (100-300 Da) with high vapor pressures, low boiling points and lipophilic character. These properties facilitate evaporation and diffusion through both water- and gas-filled pores in soil and rhizosphere environments. As compare to other microbial secondary metabolites volatiles are relatively less studied.

Objectives

The major aim of our study was to obtain more insight in the role of volatiles in microbial interactions.

Methods

In order to obtain insight in the importance of volatiles in the inter-specific interactions between soil microorganisms we performed several experiments in glass Petri dishes plates, which were designed as such that growth of different microorganisms occurred in physically separated areas within a common atmosphere. In this way we studied the role of volatiles in fungal-bacterial and bacterial-bacterial interactions. The volatiles were analysed by using thermal desorption GSMS.

Conclusions

The obtained results revealed that volatiles play important role in the interaction between soil microorganisms. Here we will report on bacterial volatiles with (1) antimicrobial activity and (2) on volatiles acting as infochemicals affecting the behavior, growth and gene expression in responding bacteria. Furthermore we will report on the effect of fungal volatiles on bacteria and on the importance of interspecific interactions for the production of volatile.
ANALYSIS OF THE DRAFT WHOLE GENOME SEQUENCE OF THE HUMIDIMYCIN PRODUCER STREPTOMYCES HUMIDUS F-100.629

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Background

Characterization of new secondary metabolite biosynthetic clusters is gaining increased attention because of their potential to produce new molecules with medical and biotechnological application. Most biosynthetic pathways in actinobacteria are cryptic or silenced in standard laboratory conditions, requiring alternative approaches to identify and these new clusters.

Humidimycin (MDN-0010) is a peptide structurally related to the antiviral siamycins that potentiates the antifungal activity of caspofungin against Aspergillus fumigatus and Candida albicans (1). Humidimycin is produced by the strain Streptomyces humidus F-100.629.

Objectives

This study has been focused on a gene mining of S. humidus F-100.629 to identify the humidimycin biosynthetic pathway and other potential cryptic biosynthetic gene clusters.

Methods

Whole genome sequence was obtained with Illumina. Assembly and annotation were performed using standard bioinformatics tools.

Conclusions

The draft genome size of S. humidus F-100.629 was estimated in 8.4 Mb. The biosynthetic cluster of humidimycin was identified and 50 putative secondary metabolite clusters were predicted using the antiSMASH application (2), including PKS-I, PKS-II, NRPS, terpenes, siderophores and bacteriocin, suggesting the genomic potential of the strain as producer of diverse natural products.

MICROEVOLUTION OF THE BIOACTIVE PEPTIDE ANABAENOPEPTIN IN CYANOBACTERIA

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Background

Bloom-forming cyanobacteria such as the genus Planktothrix sp. form a large number of bioactive peptides, e.g. the family of the Anabaenopeptins (AP) through nonribosomal peptide synthesis (NRPS).

Objectives

In total 140 Planktothrix strains, isolated from various climatic zones, were phylogenetically assigned into three major lineages that differed in ecosystem type colonization. We analyzed the distribution and recombinations of the AP synthesis gene cluster (apn).

Methods

The presence or absence of the apn gene cluster was determined by polymerase chain reaction (PCR) and sequencing of PCR products.

Conclusions

We found that the majority of strains of lineage 1, occurring in shallow lakes, carried specific apn remnants in parallel to the full gene cluster. The recombinations occurred within adenylation domains resulting in rare structural variants as well as a fusion of part of the ABC transporter domain from another NRPS gene cluster. In contrast, strains of lineage 2, isolated from deep lakes, did not contain apn remnants but always contained the full gene cluster. Strains of lineage 3, isolated from tropical regions, did not carry any sequence related to apn. Overall evolutionary diversification of apn genes was congruent with ecological diversification. Because of the occurrence of apn remnants and recombination events within the apn in lineage 1 we hypothesize that its ancestor lost the apn genes but some of the genotypes regained it through horizontal transfer.
A NEW GENE CLUSTER FOR MYCOSPORINE-LIKE AMINO ACID BIOSYNTHESIS IN CYANOBACTERIUM: HETEROLOGOUS EXPRESSION, PURIFICATION, AND CHARACTERIZATION

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Background

Mycosporine-like Amino Acids (MAAs) are an important class of secondary metabolites known for their protection against ultra violet (UV) radiation and other cellular functions, such as scavenging of free oxygen radicals, regulation of osmotic balance, and resistance to thermal and salt stress and desiccation. Cyanobacteria are microorganisms that colonize diverse ecological niches exposed to the UV light and produce MAAs. Because of its efficient sunscreen role, the MAA shinorine from cyanobacteria is utilized commercially in sunscreen creams.

Objectives

The biosynthesis of MAA was recently solved and showed that the structural diversity is directly encoded by the gene cluster. Bioinformatic analysis of the genome of the soil strain Cylindrospermum stagnale revealed a new gene cluster homolog to the MAAs biosynthetic gene cluster. Analysis of this gene cluster presents five genes instead of three/four common biosynthetic genes in other cyanobacteria.

Methods

The gene cluster was successfully cloned and heterologously expressed in E. coli. The MAA was extracted from recombinant E.coli cultures and identified through HPLC and a UV absorption maxima of 310 nm. Structural characterization is underway, combining the data from mass spectrometry and NMR.

Conclusions

The characterization of MAA gene cluster from Cylindrospermum stagnale shows a new gene organization suggesting that it codes for a novel biosynthesis pathway. The new MAA, potentially produced by this terrestrial strain, might play a role for combating the terrestrial environmental stress(s).
Background

Phycocyanin (PC) is the major pigment of the phycobiliproteins in cyanobacteria. The commercial applications of PC could be divided into two sections; the first is an ingredient for nutrient elements, natural dyes for food and cosmetics, and the second is a potential remedy in oxidative stress-induced diseases.

Objectives

The aim of this study was to increase the PC content in the isolated cyanobacterium (Nostoc sp.) with nitrogen-free medium and chromatic acclimation.

Methods

Nostoc sp. was cultured for 10 days in 1-L cylinder flasks containing 800 ml nitrogen-free BG11 medium. For chromatic acclimation, two types of light sources were compared, i.e. plant-fluorescent lamp (only blue and red wavelengths are provided) and white fluorescent lamp.

Conclusions

Under fluorescent lamp condition, the maximum biomass production was 2.24 g DCW/L (dry cell weight g/L). High biomass production could be achieved by nitrogen fixation of Nostoc sp., even without nitrogen supply. The maximum biomass concentration was higher at 2.65 g DCW/L under plant lamp condition. It means that blue and red lights in plant lamp are more critical in promoting cell growth. Cell color also changed from brown to green through chromatic acclimation by plant lamp. Higher PC content (15%) in cells was obtained and maintained throughout the cultivation. By changing light source from white to plant lamp, biomass production and PC content have been increased. Cost-effective PC production was possible,
due to its nitrogen fixation ability. More simplified PC extraction process was also devised for more efficient PC production.
ENDOPHYTIC MICROBIOTA OF THE AMAZON: ANTIBIOTIC ACTION OF METABOLITES IN PATHOGENS PRESENT IN WOUND

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Background
The endophytic microorganisms (EM) of the Amazon are potential producers of secondary metabolites with biological activity that can be used on infected wounds.

Objectives
This research investigated the antibiotic action of ME from the Amazon front of pathogenic microorganisms present in the in wound healing process.
Methods
The antibiotic activity of 171 secondary metabolites of endophytic microorganisms (MSME) were analyzed against Staphylococcus aureus, Pseudomonas aeruginosa, Escherichia coli and Candida albicans by diffusion in agar. In the test were used 50µL of each MSME extract; ampicillin, tetracycline, ciprofloxacin and fluconazole, all in 2% as positive control according to the test microorganism and DMSO as a negative control. The readings of the zones inhibition were carried out in 24 and 48 hours. The mean, standard deviation, ANOVA and Tukey's test (p <0.05) were performed for analysis.

Conclusions
The largest zone of inhibition with significant differences for C. albicans were MSME DfGa2 1.2 G4 (2.88 ± 0,71cm) and DfGa2 1.2 G5 (2.72 ± 0.69cm), for S. aureus were MSME DfGa2 1.2 G4 (2,52±0,83cm) and DfGa2 1.2 G5(2,32±0,45cm), for P. aeruginosa were MSME DfGa2 1.2 G5 (2,24±0,30cm). For E.coli there was no significant difference between the means of the zones of inhibition. The authors concluded that the samples DfGa 2 1.2 G4 and DfGa 2 1.2 G5 had better antibiotic action for the tested pathogens, suggesting a formulation of a gel and, subsequently, its evaluation in vivo of the activity of healing wounds in rats.
ANTIFUNGAL ACTIVITY OF METABOLITES FROM STREPTOMYCES DIASTATOCHROMOGENES (NO.1628) AND ITS ACTIVE PRINCIPLE

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Background
The fermentation broth by Streptomyces diastatochromogenes (No.1628), isolated from a soil sample collected in Mountain Tianmu, Hangzhou, China, had strong antifungal activity against Alternaria solani, Botrytis cinerea, Fusarium oxysporum and Rhizoctonia solani.

Objectives
The aim of this research is to isolation and identification of active compounds from S. diastatochromogenes 1628.

Methods
A bioactivity-guided approach was employed to isolate and determine the chemical identity of bioactive constituents with antifungal activity from S. diastatochromogenes 1628.

Conclusions
By the mycelium growth rate assay, it was showed that the n-butanol extract dosedependently inhibited growth of above tested phytopathogenic fungi with a potency at the concentration of 175 mg/L that was equal to that of commonly used antifungal agents at the concentrations tested. Furthermore, n-butanol extract of S. diastatochromogenes (No.1628) could effectively suppress and control cucumber rhizoctonia rot caused by R. solani. Through bioassay-guided fractionation, the n-butanol extract from S. diastatochromogenes (No.1628) afforded a new tetraene macrolide, \((7E, 12Z,13E,15E,17E,19E)\) -21 -((4-amino-3,5-dihydroxy -6-methyltetrahydro -2H -pyran-2-yl)oxy) -12-ethylidene- 1,5,6,25 –tetrahydroxy -11 -methyl-9-oxo-10,27-dioxabi-cyclo[21.3.1]heptacosa-7,13,15,17,19–pentaene-24-carboxylic acid (1), together with tetrin B (2), tetramycin A (3), toyocamycin(4) and anisomycin (5). The structures of compounds were established on the basis of spectroscopic analyses. This new compound strongly inhibited hyphal growth of R. solani and B. cinerea with IC\(_{50}\) of 0.20 and 1.53 μg/mL, respectively. Our study demonstrated that S. diastatochromogenes (No.1628) is a promising source of natural bioactive and novel metabolites and has wide application prospect in biocontrol field.
FEMS-2681
Secondary metabolites, metabolomics

PANCREATIC LIPASE INHIBITORY ACTIVITY OF ENDOPHYTIC FUNGUS ASSOCIATED WITH AEGLE MARMELOS: A BIORESOURCE FOR NOVEL ANTIOBESITY DRUG

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Background

Obesity is a disease resulting from improper balance between energy intake and expenditure and is increasingly becoming a major cause of preventable mortality. Pancreatic lipase (PL) is considered as one of the safest target for diet-induced anti-obesity drug development.

Objectives

In the process of exploration of new PL inhibitors, we have screened culture filtrates of 200 endophytic fungi from different medicinal plants using qualitative and quantitative assays. The organism was identified and partial purification of the bioactive moiety was carried out using analytical techniques.

Methods

Rhodamine olive oil plate assay and PNPL (p-nitrophenyl laurate) as substrate was used for quantitative assessment of PL activity inhibition.

Conclusions

The qualitative assays indicated potential PL inhibition in 27 isolates. Further, only organic extract of these culture filtrates exhibited PL inhibition in range between 30-100%. #6 AMLWLS exhibited complete inhibition of PL with an IC₅₀ of 2.12 μg/ml which was comparable to the Orlistat exhibiting an IC₅₀ value of 2.73 μg/ml. Purification of the extract by column chromatography led to isolation of the pure compound which was identified as a tetra peptide molecule on the basis of C¹³ NMR, HR/MS and biochemical analysis. Further molecular phylogenetic tools and morphological studies were used to identify the isolate #6 AMLWLS as Fusarium incarnatum species.

Hence endophytic fungal isolate #6 AMLWLS stands out as a potential candidate for anti-obesity therapy
Background
Bioprospecting using endophytic fungi is a promising source for the detection of novel compounds with biological activities of interest in health care, especially for diseases such as leishmaniasis (neglected tropical diseases) and cancer.

Objectives
The objective was to evaluate the cytotoxicity and the leishmanicide activity of endophytic fungi extracts isolated from *H. suaveolens* (L) Poit and *C. lanceolatum* Pohl against tumor cell lines MCF-7 and MDA-MB-231 and promastigotes forms of *L. amazonensis*.

Methods
We have evaluated 25 fungal culture extracts obtained by maceration of the mycelium in ethyl acetate (EtOAc). The fungi have isolated from *H. suaveolens* and *C. lanceolatum*. The extracts were tested against tumor cell lines MCF-7 and MDA-MB-231 and promastigotes forms of *L. amazonensis*. The test used was a colorimetric method with tetrazolium salt 3- [4,5-dimethylthiazol-2-yl] -2,5- diphenyltetrazolium bromide (MTT). The concentration of the extracts ranged from 100 to 12.50 µg/mL.

Conclusions
Extracts of 5 fungal isolates (*Taifanglania curticatenata*- F27; *Marasmius* sp.-21C, *Fusarium oxysporum*-33C; *Macropomina phaseolina*-46C and *Trichoderma spirale*-66C) inhibited the proliferation of promastigotes at a low IC(50) of between 25.61 and 52.60 µg/ml. Six extracts (*M. phaseolina*- F1/46C; *T. curticatenata*-F27; *Cladosporium flabelliforme*- F29; *Neosartorya pseudofischeri*-F36, and *Corynespora cassiicola*-72C) were cytotoxic to both cell lines at a low IC(50) of between 23 and 48.72 µg/ml. Our results indicate that the medicinal plantas living in wetlands Brazilian shelter an interesting bioactive fungal community that is able to produce leishmanicidal and antitumoral molecules. These molecules may be used to develop new leishmanicidal and anticancer drugs.

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VERIFICATION OF PEPTAIBOL PRODUCTION ABILITY OF A TRICHODERMA STRAIN SELECTED BY A BACTERIUM-BASED SCREENING METHOD

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Background

Filamentous fungi are producers of a large number of valuable secondary metabolites due to their promising biological effects. Identification of these compounds and characterization of their biological activities are the important goals of research working at the interface of chemical and biological sciences.

Objectives

We aimed to verify the peptaibol production of a Trichoderma strain selected from the Szeged Microbiological Collection by a Micrococcus luteus-based agar plate screening method.

Methods

Mass spectrometric analysis of putative peptaibols produced by a T. pleuroticola strain have been carried out via on-line reversed-phase high performance liquid chromatography (HPLC) coupled to electrospray ionization ion trap mass spectrometry (ESI-IT-MS) after the solid phase clean-up of culture extracts. Initially, the separation and the mass spectrometric parameters were optimized using alamethicin standard peptaibol mixture and the unknown components produced by the selected isolate were investigated and determined.

Conclusions

Our mass spectrometric measurements confirmed the results of the biotest selection method used for the detection of peptaibols. Some of the produced peptaibols showed the characteristic mass spectral components of trichorzianins, including TA IIIb, TB IVb, TAP-14a and TA VII. However, besides the known molecules, seven new trichorzianin-related compounds were also detected; these contained some
amino acid changes in the core trichorzianin sequence. Comparing the amount of the
different components, compounds with a molecular mass of 1909 amu were present
in the highest amount. This research was supported by OTKA grant K-105972 from
the Hungarian Scientific Research Fund and a bilateral grant from “Stiftung Aktion
Österreich-Ungarn”.

PURIFICATION AND STRUCTURAL ELUCIDATION STRATEGY OF PEPTAIBOLS PRODUCED BY TRICHODERMA STRAINS

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Background

An outstanding group of secondary metabolites are peptaibols, produced mainly by fungal species from the genus Trichoderma. These compounds show a wide spectrum of biological activities including antibacterial, antifungal and antiviral effects.

Objectives

We aimed to develop a simple and easy-to-follow peptaibol characterization strategy including purification and structural elucidation steps, which can help to analyze the peptaibol-producing abilities of various isolates within the genus.

Methods

Our characterization strategy includes the mass spectrometric analysis of the culture extracts of possible peptaibol-producing Trichoderma strains, the semi-preparative purification of the selected compounds, identification of amino acids in their sequences, and chiral analysis of the identified building blocks. These tasks require the use of microbiological methods for strain identification and for different cultivations and the application of separation techniques such as solid phase extraction, normal and reversed-phase HPLC and mass spectrometry.

Conclusions

During the study, the different methodological steps were optimized and their application served the detailed characterization of some Trichoderma strains from the point-of-view of their peptaibol profiles. The examined isolates are belonging to the species T. aggressivum, T. atroviride, T. longibrachiatum, T. orientale, T. pleuroticola, T. pleurotum and T. virens. The peptaibol compositions showed high level of diversity and their amounts presented in a wide range. A number of already described compounds have been identified from species other than the original producer and novel peptaibols were also defined. This research was supported by OTKA grant K-105972 from the Hungarian Scientific Research Fund and a bilateral grant from “Stiftung Aktion Österreich-Ungarn”.
Background

Ophiobolins are a remarkable family of secondary metabolites with various biological activities such as calmodulin antagonism, cytotoxic, antimicrobial and nematocid effects.

Objectives

Our aim was to develop a rapid and efficient method for purification of different bioactive sesterterpene-type fungal secondary metabolites belonging to the ophiobolin family.

Methods

Based on our preliminary results, a Bipolaris oryzae strain was selected and cultivated in PDB. The ferment broth was extracted with ethyl-acetate and concentrated. The purification of the crude extract was carried out using subsequential semi-preparative liquid chromatographic techniques. Initially, normal phase column chromatography was carried out using the mixture of ethyl-acetate and n-hexane, and the fractions containing large amount of potential ophiobolin compounds were further purified on a semi-preparative reverse phase HPLC system with the mobile phases of acetonitrile and water. After each purification step, the collected fractions were analyzed on an analytical HPLC system to determine their purity. The purified compounds were identified using mass spectrometric and NMR experiments.

Conclusions

The selected B. oryzae strain produced several ophiobolin analogues. The purified compounds synthesised in the highest amount proved to be ophiobolin A; its identity was confirmed with HPLC-MS and NMR experiments. Besides ophiobolin A, four other ophiobolin analogues were also purified including 3-anhydro-ophiobolin A,
ophiobolin I, 6-epi-ophiobolin A and 3-anhydro-6-epi-ophiobolin A with the purities over the 95%. This research was supported by grant TÁMOP-4.1.1.C-12/1/KONV-2012-0012. Csaba Vágvölgyi thanks the visiting professor program, Deanship of Scientific Research, King Saud University, Riyadh.
Background

Rhamnolipids are potent biosurfactants with high potential for industrial applications. They are currently produced with the opportunistic pathogen *Pseudomonas aeruginosa* during growth on hydrophobic substrates such as plant oils. The heterologous production of rhamnolipids enables the use of a non-pathogenic host and cheaper substrates such as glucose. Glucose also has the advantage of simpler purification, e.g., via coupled foam fractionation and adsorption [1].

Objectives
To engineer a non-pathogenic rhamnolipid producer converting glucose or other renewable resources efficiently into the target product.

**Methods**

Introduction of the biosynthesis pathway for rhamnolipids from *P. aeruginosa* in the non-pathogenic *P. putida*. Metabolic engineering of secondary metabolite producers implicitly relies on high flux through central carbon metabolism. Our strategy relies on the ability of *P. putida* to reroute metabolic resources into a peripheral pathway when a high demand exists, e.g., initiated by high transcriptional activity.

**Conclusions**

We previously presented a recombinant *P. putida* able to produce 0.2 g/L rhamnolipids [2]. For optimization, several techniques were used. A synthetic promoter library was developed using an approach based on degenerated primers [3]. This technique yields an array of transcriptional activity, potentially allowing the identification of optimal enzyme activity for high flux towards rhamnolipid synthesis. The second step was eliminating a pathway competing for a common precursor. The best producing strain was able to reach a titer of 3 g/L with a yield of 40% and a very high specific rhamnolipid synthesis rate, which is possible because the flux through the precursor providing pathways is significantly increased in the engineered strain.
FUNCTIONAL CHARACTERIZATION OF A GENE INVOLVED IN THE BIOSYNTHESIS OF MYCOPHENOLIC ACID IN THE FUNGUS PENICILLIUM ROQUEFORTI

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Background

Penicillium roqueforti is a filamentous fungus used in the ripening of blue-veined cheeses. Previous studies have shown that this organism produces several secondary metabolites, including mycophenolic acid, a meroterpenoid with immunosuppressive activity of great pharmaceutical interest. However, the genes involved in the synthesis of this metabolite in P. roqueforti have not been described or characterized so far. In other organisms such as P. brevicompactum, it has been described that the mycophenolic acid biosynthetic cluster consists of seven genes, among them mpaDE, which encodes for a natural fusion enzyme consisting of a cytochrome P450 and a hydrolase.

Objectives

To evaluate the role of the mpaDE gene in the biosynthesis of mycophenolic acid in P. roqueforti.

Methods

The genome of P. roqueforti was analyzed using bioinformatic tools and an ORF that corresponds to the hypothetical orthologue of mpaDE was identified. The expression of this gene was down regulated using RNAi-silencing technology. For this purpose, P. roqueforti was transformed with an appropriate genetic construct, and several transformants strains showing dramatic reductions in the levels of mpaDE transcript were selected. Finally, extracts from these transformants were analyzed by HPLC.

Conclusions

P. roqueforti wild-type showed high levels of mycophenolic acid. On the contrary, the transformants showed barely detectable levels of this metabolite. These results indicate that mpaDE is involved in the biosynthesis of mycophenolic acid in P. roqueforti.

This work was supported by FONDECYT 1120833 y DICYT-USACH.
A UNIQUE GLUCOSE SENSOR GCR1 OF THE METHYLOTRPHIC YEAST HANSENULA POLYMORPHA CAN TRANSPORT MONOSACCHARIDES

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Background
Several proteins involved in glucose sensing, signalling and transport have been described for methylotrophic yeast Hansenula polymorpha. Among them Gcr1, a peculiar C-tale-less glucose sensor, required for glucose repression and regulation of hexose transport.

Objectives
The aim of this work was to further elucidate the H. polymorpha Gcr1 protein function in glucose transport.

Methods
Wild type NCYC495 (WT) and mutant strains of H. polymorpha, classical methods of molecular genetics and microbiology were used.

Conclusions
The closest homologue for glucose sensor Gcr1 of H. polymorpha is the functional high-affinity monosaccharide transporter involved in H⁺-dependent glucose symport – MstA from Aspergillus niger. Regulated HpGcr1 production in hexose transporter-less Saccharomyces cerevisiae mutant did not restore growth of this strain on different glucose concentrations. Similarly to Gcr1, functional high-affinity glucose transporters of Trichoderma harzianum Gtt1 and Aspergillus nidulans HxtA did not complement growth deficiency of S. cerevisiae hxt null on different carbohydrate substrates. Grown on methanol-containing medium WT-derivative strains with constitutive expression of GCR1 and HXT1 encoding low-affinity hexose transporter were more sensitive to exogenous glucose antimetabolite 2-deoxy-D-glucose (2-DOG) (concentration 0.3 mM) compared to the WT strain. The toxic effect of 2-DOG was more acute on medium with lower pH. Therefore, Gcr1 can possibly transport glucose toxic analogue into cells, thus increasing its harmful effects.

H. polymorpha protein Gcr1 that most probably results from a horizontal transfer from Aspergillus fungi, is essential for glucose catabolite repression of genes involved in metabolism of alternative carbon substrates and "very-high affinity" glucose transport.
FEMS-1893
Signalling

DESIGNING FUNCTIONAL HETERODIMERS TO DECIPHER SIGNAL TRANSDUCTION IN BVGS

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Background

The virulence of Bordetella pertussis, the whooping cough agent, is regulated by the two-component system BvgAS. BvgA is a classical response regulator while BvgS is an unorthodox sensor-kinase harboring receiver and Histidine phosphotranfer domains. BvgS contains three putative perception domains, two periplasmic Venus Flytrap domains linked by a transmembrane segment to a cytoplasmic PAS domain. The kinase of BvgS is active by default and is turned off by negative modulators such as nicotinate or sulfate ions.

Objectives

BvgS functions as a homodimer, and its phosphorylation cascade works in trans. We took advantage of this property to construct a merodiploid strain in which only heterodimers, but not homodimers, are functional to decipher the signal transduction mechanism involved in BvgS.

Methods

We introduced point mutations in the various sensor domains to test their effect in heterodimers. The same substitutions in the context of a wild type phosphorylation cascade affect either the basal kinase activity of BvgS or its ability to respond to modulation, as seen using a lacZ reporter gene under the control of a virulence gene promoter.

Conclusions

When substitutions affecting the response to modulation are introduced in the periplasmic VFT domains of BvgS, the WT monomer is able to partly compensate for the defect of the other monomer, yielding intermediate phenotypes. In contrast, the substitutions introduced in the PAS domain appear to be dominant over a wild type PAS domain. This suggests that signal transduction is achieved through different mechanisms in the periplasmic and cytoplasmic moieties of BvgS.
THE STRUCTURE AND FUNCTION OF ISOLATED N-TERM NAL AMPHIPATHIC HELIXES OF MEMBRANE STRESS CONTROLLERS PSPA AND VIPP1

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Background

All cell types maintain the integrity of their membrane. One widely distributed plasma membrane stress response system in bacteria is the phage shock protein (Psp) system. Its central component PspA has a counterpart Vipp1. Both proteins are directly responsible for membrane maintenance under stress conditions while PspA also negatively regulates its own expression via interaction with the AAA+ ATPase PspF.

Objectives

The PspA and Vipp1 N-terminal amphipathic helix A (ahA) is implicated in their membrane binding, and ahB in PspA is required for negative regulation. A direct interaction between ahA and the lipid bilayer for effector function and ahB and PspF for regulatory function has yet to be determined.

Methods

Purified short peptides derived from the amphipathic helixes were used to directly probe membrane binding and regulatory function. Defined membrane vesicle in vitro assays and CD spectroscopy characterised the functionalities of the ahs and the downstream effects on membrane stability and transcription regulation.

Conclusions

We observed direct membrane-binding of ahA derived peptides and in the case of Vipp1 an accompanying change in secondary structure from random coil to alpha
helical upon membrane association. Binding specificity by varying the membrane
anionic lipid content and stored curvature elastic stress linked the functionalities of
the peptides with those of full length proteins. ahB of PspA inhibited the ATPase of
PspF proving its direct regulatory role. These findings establish synthetic peptides
can probe PspA structure-function and target regulation of the Psp response, which
could be of interest for controlling pathogens where Psp response is important and
conserved.
NEW INSIGHTS IN THE ACTIVATION OF THE PSEUDOMONAS AERUGINOSA VIRULENCE REGULATOR SIGMA VREI AND THE PUMA3 CELL-SURFACE SIGNALING SYSTEM

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Background

Cell-surface signaling (CSS) is a regulatory mechanism used by gram-negative bacteria to modify gene expression in response to environmental signals. CSS regulation is extensively present in the human opportunistic pathogen Pseudomonas aeruginosa. Most CSS systems of this bacterium regulate iron uptake. However, the PUMA3 CSS system regulates virulence. P. aeruginosa PUMA3 is constituted by the VreA receptor, the VreR anti-sigma factor and the sVreI extracytoplasmic function (ECF) sigma factor. VreR binds to and keeps inactive sVreI in absence of the PUMA3 inducing signal. The role of VreA is to sense the presence of the signal and to transduce it to the VreR anti-sigma factor allowing the release and activation of sVreI and the expression of the PUMA3 regulated genes.

Objectives

The PUMA3 inducing signal is still unknown. However it has been recently shown that expression of the vreAIR operon is induced in low phosphate (Pi). In this condition, some degree of sVreI activity is also observed despite the fact that the VreR anti-sigma factor is also produced. In this work, we aimed at analyse the role of VreA and VreR in both the expression and the activation of sVreI in low Pi.

Methods

P. aeruginosa vreR and vreA deletion mutants has been constructed by allelic exchange. Gene expression was monitored using lacZ transcriptional fusions. sVreI stability has been assayed by Western-blot.

Conclusions

Full activation and stability of sVreI in low Pi requires the removal of VreR and the presence of an additional inducing signal that targets this process.
IMPACT OF POTENTIAL IN VIVO DIVK DIMERIZATION ON CAULOBACTER CRESCENTUS CELL CYCLE

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Background

Response regulators are crucial actors of two-component systems and have often been shown to dimerize. We focus on DivK, an essential single domain response regulator of the aquatic α-proteobacterium C. crescentus. DivK is part of a phosphorelay controlling developmental events and belongs to the same protein subfamily as CheY and Spo0F, two response regulators known to dimerize. A previous study has highlighted DivK dimerization in vitro (Guillet et al., The Journal of Biological Chemistry, 2002) but the relevance of this dimerization in vivo is still a mystery.

Objectives

The goal of our research is (i) to assess whether DivK is able to dimerize in vivo and (ii) to determine whether this dimerization mediates DivK subcellular localization and/or the interaction with potential partners. We are also interested in investigating the importance of DivK phosphorylation in its dimerization.

Methods

Under native conditions, DivK appears as a doublet which is not visible under reducing conditions. Interestingly, the relative intensity of the DivK bands varies when the PleC phosphatase or the DivJ kinase, both acting on DivK, are knocked-out. In addition, the doublet is upshifted in the DivK_{D90G} mutant, whose subcellular localization is affected. The nature of this doublet is currently under investigation. We are also trying to identify DivK partners by MS-MS analysis of DivK co-immunoprecipitates obtained from various genetic backgrounds.

Conclusions

Preliminary results suggest that DivK is indeed able to dimerize in vivo and to bind different partners. This dimerization could be triggered by phosphorylation.
FEMS-2171
Small regulatory RNAs

USE OF ANTISENSE RNA TO MODULATE HSP GENE EXPRESSION IN OENOCCUS OENI
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Background

Oenococcus oeni is a wine-associated lactic acid bacterium responsible for wine malolactic fermentation. Wine exhibits harsh and challenging conditions: low pH, low temperature, nutrient-poor and presence of ethanol leading to O. oeni stress response. Understanding of the mechanisms involved in stress tolerance is essential to improve O. oeni development in wine.

Stress adaptation of O. oeni has been studied and many hsp genes, induced in oenological conditions, have been identified. The hsp\textit{18} gene, encoded the small Hsp Lo18, have been particularly characterized. In addition to its chaperon activity, Lo18 could have a lipochaperon-function and maintains membrane integrity in stress conditions.

Objectives

Up to now, characterization of O. oeni genes was restricted due to lack of genetic tools for gene replacement. Accordingly, we focused our work on gene inactivation by using antisense RNA approach to modulate hsp gene expression in O. oeni. With the goal of understanding the function of O. oeni hsp genes in vivo, we have developed an efficient expression vector to produce antisense RNA targeting hsp\textit{18} mRNA and reduce sense RNA transcript.

Methods

After transformation into O. oeni, transformants were exposed to multiple stresses inducing hsp\textit{18} gene expression: heat shock, acid shock and presence of ethanol.

Conclusions

We highlighted that in vivo antisense inhibition of hsp\textit{18} expression strongly affects the survival of O. oeni in stress conditions. This study presents a new efficient genetic tool for O. oeni and demonstrates for the first time the use of antisense technology for modulating gene expression in O. oeni.
DIFFERENTIAL EXPRESSION OF SMALL RNAS IN RESPONSE TO CHEMICAL STRESS
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Background
Bacterial sRNAs are often expressed in response to changing environmental conditions and function to modulate gene expression. There are numerous documented connections between sRNAs and stress, where sRNAs regulate important processes in response to metabolite, envelope, oxidative, iron deficiency, anaerobic and pH stress.

Objectives
Although chemical stress is routinely encountered in microbial processing applications due to the toxicity of substrate, intermediate or product compounds, the cellular response and the involvement of sRNAs in this process is poorly understood. We have used an RNA sequencing approach to map the E. coli sRNome during chemical stress and high cell density fermentations with the aim of gaining insights into the chemical stress response and identifying sRNAs with roles in stress tolerance that have potential applications in the design and optimization of future production strains.

Methods
RNA sequencing libraries were prepared from RNA isolated from E. coli MG1655 cells subjected to chemical stress with twelve compounds including organic acids, amino acids, and organic solvent-like compounds. The E. coli MG1655 strain was also grown under high cell density fermentation conditions, where cells were harvested in exponential batch, glucose-limited exponential fed-batch, transition and stationary growth phases.

Conclusions
Over 250 novel intergenic sRNAs have been identified with this approach, adding to the roughly 200 previously reported sRNAs in E. coli. A significant fraction of the sRNAs exhibit specific expression patterns during high cell density fermentation and
a group of them are differentially expressed in the presence of multiple chemicals, suggesting they may play regulatory roles during these stress conditions.
Background
Bacterial sRNAs are 50-400 nucleotide-long RNA molecules, most of which exert their regulation by base-pairing with their target mRNAs. In many bacteria, including *Escherichia coli*, the interaction between an sRNA and its targets is mediated by the Hfq protein, resembling the situation in eukaryotic cells where the interaction of a microRNA and its targets involves Argonaute. While it is widely accepted that miRNAs guide Argonaute proteins to their specific targets through sequence complementarity, the dynamics of sRNA-Hfq-target interaction, to the best of our knowledge, has not been yet fully determined.

Objectives
As a first step towards this question, we study the relationship between the levels of Hfq-bound sRNA and its target mRNAs. More specifically, we aim to measure the change in Hfq-bound target mRNA levels following a change in the expression level of their sRNA regulator.

Methods
To this end we applied RNA-seq to measure the levels of total RNA and RNA co-IPed with Hfq from three *E. coli* strains (MG1655 Hfq-FLAG, MG1655 hfq::kan and MG1655 Hfq-FLAG rnc14::Tn10) with and without overexpression of the sRNA MicA.

Conclusions
Our results indicate that most MicA target mRNAs, as well as MicA itself, were enriched in Hfq when MicA was overexpressed. In contrast, most target mRNAs of other sRNAs were depleted. In conclusion, our initial results indicate that there is a relationship between sRNA and target mRNA levels on Hfq, offering a strategy to identify putative novel targets.
Background

RNA sequencing has recently revolutionized transcriptome analyses. As a result, novel non-coding RNAs have been discovered in several organisms and subsequent studies have revealed an increasing number of new regulatory mechanisms at the level of RNA. The lactic acid bacterium *Lactococcus lactis* is of high importance for biotechnological applications, specifically in the dairy industry. The stress response systems are very well studied for this bacterium. In addition, global regulators of carbon and nitrogen metabolism have also been studied in detail. However, the presence and roles of non-coding RNAs with regulatory functions has not been assessed until now.

Objectives

We aimed to identify novel RNA elements, such as small regulatory RNAs, antisense RNAs and sORFs. In addition, expression of these novel RNAs under various stress conditions was assessed, and for one novel sRNA, the function was studied in depth.

Methods

By using differential RNA sequencing (dRNA-seq) data, a transcriptome landscape of *L. lactis* was constructed that was used for manual and automated mining using TSSer. Northern hybridization was used to confirm 12 putative regulatory RNAs.

Conclusions

This study used both manual and automated approaches to identify hundreds of novel RNA species. Experimental validation was performed for a selection of these potential RNA regulators. The role of one non-coding RNA located in the 3'UTR of *argR*, the regulator of e.g. the arc-operon in *L. lactis*, is currently being examined. This small RNA, called ArgX, appears to stabilize the arc-operon. Results of this sRNA ArgX and its role in regulating arc, will be presented.
FEMS-1755
Small regulatory RNAs

A GROUP B STREPTOCOCCUS SMALL RNA PRESERVES CELL INTEGRITY
AND CONFFERS RESISTANCE TO A CATIONIC ANTIBACTERIAL PEPTIDE
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Background

Gram-positive bacteria reduces the effectiveness of cationic antimicrobial peptides
(CAMPs), such as colistin, and preserves cell integrity by the addition of D-alanine to
lipoteichoic acids (LTAs). Thus far, no small RNA (sRNA) has been described to be
involved in the regulation of the genes involved in the D-alanylation of LTAs in the
human neonatal pathogen Group B Streptococcus (GBS).

Objectives

Here, we present the first study describing the functional characterization of a GBS
sRNA. We highlight its pleiotropic role in maintaining cell integrity and colistin
resistance.

Methods

The GBS sRNA was identified by RNAseq and its expression was confirmed by
northern blot analyses. The biological role of the sRNA was investigated by analyzing
a deletion mutant (Δsrna). The interaction region between the sRNA and its mRNA
target was predicted in silico and confirmed by in vivo mutagenesis.
Conclusions

The northern blot analyses showed that the sRNA is mainly expressed at the onset of the stationary phase until late stationary phase. Microscopic observations showed that the Δsrna strain forms significantly shorter chains, and is more sensitive to colistin (MIC = 16 µg/mL) when compared to the wild-type (MIC > 256 µg/mL). Interestingly, the in silico predicted mRNA target is downregulated in the mutant strain. Mutations in the sRNA sequence likely involved in RNA-RNA pairing are sufficient to provoke similar chains alterations than the Δsrna strain. All together, these results suggest that the sRNA is required for colistin resistance and cell integrity due to a direct regulation of the D-alanylation pathway.
DEVELOPMENT OF OAPV3 VIRAL-LIKE PARTICLES FOR INVESTIGATION OF SHEEP HUMORAL IMMUNE RESPONSE TO PAPILLOMAVIRUS INFECTION
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Background

Papillomaviruses are a diverse group of small, non-enveloped, double stranded DNA viruses that cause proliferations of the stratified squamous epithelium of the skin and mucosa in a wide variety of vertebrate hosts. Papillomaviruses play an important role in human cancer development, and have been isolated from a number of animal malignancies. Recently we identified a novel ovine papillomavirus, OaPV3, associated with squamous cell carcinoma. Like other papillomaviruses, OaPV3 is not able to productively infect cells in culture. The lack of in vitro factories for viral production hampers the investigation of pathogenetic mechanisms and interaction with host immune response.

Objectives

Main goal of this study is the production of non-infectious OaPV3 viral-like particles (VLPs) for evaluating humoral immune response in naturally infected sheep.

Methods

Non-infectious VLPs were obtained by expressing the L1 major capsid protein in insect cells using recombinant baculoviruses. Two sheep were immunized with OaPV3 VLPs. Sera were collected after each of the 3 immunisations. VLPs were used in western blotting and ELISA to test reactivity of 90 sera collected from sheep sampled in flocks naturally infected by OaPV3.

Conclusions

Data indicate a specific immune response of immunised sheep. In naturally infected flocks, only sheep with clinical lesions showed detectable antibodies. OaPV3, like human cutaneous papillomavirus seems to escape humoral response by playing “hide and seek” with host immune system.
Background
Papillomaviruses play an important role in human cancer development, and have been isolated from benign and malignant proliferative skin lesions. However, the association of papillomaviruses with cutaneous lesions has been poorly investigated in sheep. To date, three ovine papillomavirus types have been classified. OaPV1, OaPV2 belong to Deltapapillomaviruses genus and were found in cutaneous fibropapillomas of australian merinos. OaPV3 belongs to a new genus, Dyokappa, and has been recovered from cutaneous squamous cell carcinomas of sarda breed sheep.

Objectives
In this study we investigated unknown ovine papillomaviruses in cutaneous fibropapillomas of sarda breed sheep.

Methods
Rolling circle amplification was performed on total DNA extracted from skin lesions biopsies, and resulted in the presence of a novel ovine Papillomavirus (OaPV4). The entire OaPV4 genome was cloned and sequenced. IHC with L1 antibodies and ISH with E6 dig-probe were carried out in order to establish viral cellular tropism.

Conclusions
A novel papillomavirus, OaPV4, was identified. Viral L1 shares 82.5% nucleotide homology with the closest relative (OaPV1). Maximum likelihood phylogenetic tree placed OaPV4 in a monophyletic clade including all Deltapapillomaviruses so far described.

OaPV4 infects both epithelial cells and fibroblasts and has a role only in benign lesion development. This is the first report of a Deltapapillomavirus 3 species in European sarda breed, and this support the existence of an OaPV1/OaPV4 common ancestor infecting sheep before radiation and evolution of modern sheep breeds.
EQUUS ASINUS PAPILLOMAVIRUS (EAPV1) PROVIDES NEW INSIGHTS INTO EQUINE PAPILLOMAVIRUS DIVERSITY

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Background

The Papillomaviridae family consists of a large and diverse group of viruses characterised by a double strand, covalently linked circular genome typically ranging from 7 to 8 kb in size, and causing proliferative lesions in animals and human.

Objectives

Aim of this study was to investigate papillomavirus diversity in Equidae and to establish the presence of PV in species related to horse. We detected a novel papillomavirus (EaPV1) from healthy skin and from sun associated cutaneous lesions of an Asinara (Sardinia, Italy) white donkey reared in captivity in a wildlife recovery centre. EaPV1 genome is 7467 bp long, and shows some characteristic elements of horse papillomaviruses, including a small untranslated region between the early and late regions. A typical E6 ORF is missing. EaPV1 DNA was detected in low copies in normal skin of white and grey donkeys of the Asinara Island, and does not transform rodent fibroblasts in standard transformation assays. Pairwise nucleotide alignments and phylogenetic analyses based on concatenated E1-E2-L1 amino acid sequences revealed the highest similarity with the Equine papillomavirus type 1.

Methods

The entire genome of EaPV1 was cloned, sequenced, and characterised. Methods included: DNA extraction, traditional and real-time PCR, RCA, cloning, sequencing and phylogenetic analyses.

Conclusions

The discovery of EaPV1, the prototype of a novel genus and the first papillomavirus isolated in donkeys, confirms a broad diversity in Equidae papillomaviruses. Taken together, data suggest that EaPV1 is a non-malignant papillomavirus adapted to healthy skin of donkeys.
BACTERIOSTATIC SYNERGISTIC EFFECT OF RECOMBINANT HUMAN LACTOFERRIN, SILVER AND ZINC NANOPARTICLES ON PATHOGENIC ENTERIC BACTERIA ISOLATED FROM DIARRHEAL CALVES

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Background
The spread of multidrug-resistant Enterobacteriaceae is complicating the treatment of diarrhea in neonatal calves. Growing resistance of microorganisms to antibiotics requires to find novel effective antibacterial drugs.

Objectives
To evaluate the bacteriostatic effect of human lactoferrin (hLf), silver nanoparticles (AgNPs), zinc nanoparticles (ZnNPs) and their combinations on pathogenic enteric bacteria isolated from diarrheal calves.

Methods
Recombinant hLf was isolated from milk of transgenic goats generated by microinjecting sequence encoding hLf cDNA to the pronuclear. AgNPs and ZnNPs were prepared by chemical methods. The radii of particles were between 20 and 25 nm. E.coli, Pr. mirabilis and Kl. pneumoniae used in these experiments were isolated from diarrheal calves. Isolated bacteria were resistant to ciprofloxacin and erythromycin. To evaluate the bacteriostatic effect of hLf, nanoparticles and their combinations the broth microdilution minimal inhibitory concentration (MIC) testing was applied.

Conclusions
The results suggested that the MIC value of AgNPs against E.coli, Pr. mirabilis and Kl. pneumoniae was about 2-4 μg/ml; ZnNPs activity was lower (about 8-16 μg/ml). The synergistic antimicrobial effect between hLf and NPs was also evident. The MICs of hLf decreased when used together with NPs. So the MIC of hLf against all of the tested bacterial strains was reduced from 4-8 mg/ml when used alone to 0.5-2 mg/ml when used in conjunction with 1 μg/ml of AgNPs. The synergistic activity of hLf and ZnNPs combination was less pronounced. The best synergistic combination was AgNPs and hLf that allows to recommend it for the treatment of diarrhea in neonatal calves.
FEMS-2137
Veterinary microbiology

EFFECTS OF COLD STRESS ON BACTERIAL ISOLATION AND IMMUNE CELLS IN CAMPYLOBACTER JEJUNI INFECTED SPF CHICKENS.
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Background

Stress activate the hypothalamic-pituitary-adrenal axis and the sympathetic nervous system resulting in a series of neural and endocrine adaptations. In industrial production the cold stress are one of the most important problem in the first week of poultry production. Many microorganisms are present in the livestock and poultry products are considered main source of Campylobacter jejuni to humans.

Objectives

Our aims were challenge chickens with cold stress and Campylobacter jejuni (ATCC33291) infection to assess the role of stress in the interaction host-pathogen

Methods

Thirty-two Leghorn SPF birds were housed in isolators divided into three groups: C (negative control not shown), CI (Campylobacter jejuni infected - 10^4 ufc) kept for 21 days in thermal comfort and the group CI+CS (Campylobacter jejuni infected - 10^4 ufc + cold stress) was subjected to cold stress for 6 h per day during 7 days at a 19°C. At 7 and 21 days old, bacterial isolation from liver and cecum, blood flow cytometry to CD45, CD3, CD4 and CD8 antibodies, plasma corticosterone (CS) (Fig.1) and blood glucose (BG) levels were performed in the birds.

Conclusions

We observed high BG and CS (p<0.005) during entire experiment in CI+CS group. Also, the cold stress increased bacterial isolation at 21days in the liver and cecum (Fig. 2). At 21days of age, we observed a increase of CD3+ (p<0.05) CD4+ and CD8+ peripheral cells in CI+CS group (Fig.3). We may conclude that high CS and BG in birds subjected to cold stress affect negatively the bacterial clearance and immune cells distribution.
Plasma Corticosterone in SPF Chickens from 7 and 21 days old.

Campylobacter jejuni isolation from SPF Chicken with 21 days old.
CD4+ and CD8+ cells from SPF Chicken blood at 21 days old.
MYCOPLASMA AGALACTIAE ESCAPES SHEEP NEUTROPHIL EXTRACELLULAR TRAPS (NETS)

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Background

Mycoplasmas are the smallest self-replicating bacteria that evolved a parasitic lifestyle through regressive evolution. The ability of mycoplasmas to modulate immune responses allows the establishment of persistent infection resulting in chronic inflammatory diseases. Mycoplasmas evolved different strategies to escape host immune system. Despite mechanisms of humoral response evasion are known, resistance to innate response is still insufficiently investigated. To date, studies of interaction between mycoplasmas and granulocytic neutrophils are still lacking. NETs are a mixture of antimicrobial peptides and proteins "caught" in a backbone of DNA filaments that trap bacteria, which are cleared by macrophage through phagocytosis. It has been demonstrated that Staphylococcus aureus (S. aureus) breaks down NETs thorough its SNase and induces macrophages apoptosis. Recently, we demonstrated the in vivo expression of MAG_5040, the SNase homologue of M. agalactiae. We speculate that MAG_5040 might also be an important virulence factor of mycoplasmas.

Objectives

Main goal of this study is to investigate the interaction of M. agalactiae with NETs and to verify if MAG_5040, similarly to its S. aureus homologue, allows M. agalactiae to avoid NETs-mediated neutrophils extracellular killing.

Methods

Sheep neutrophils, either untreated or activated with PMA and/or MAG_5040, were incubated with M. agalactiae. After 3 hours, extracellular DNA was stained with SYTOX Orange and images were detected with a confocal microscope. Differences in fluorescence intensity were also calculated.

Conclusions

Data suggest that M. agalactiae is able to induce NETs formation and their digestion ex vivo, suggesting a novel mechanism for escaping sheep innate immune response.
PERSISTENT CONTAMINATION OF SMALL AND MEDIUM Sized POULTRY FARM ENVIRONMENTS BY CLOSTRIDIUM PERFRINGENS IN THE UNITED KINGDOM

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Background

Clostridium perfringens is a cause for increasing concern due to its involvement in severe infections in both humans and animals, especially poultry. Although C. perfringens is part of the natural microbiota of the intestinal tract in poultry, it is an opportunistic bacterium which can be pathogenic resulting in the possibly severe disease - necrotic enteritis.

Objectives

The aim of the present study was to determine the occurrence of C. perfringens on one single sampling occasion from poultry enterprises widely distributed across the United Kingdom. The samples were toxin genotyped by PCR and analysed by PFGE and antimicrobial testing.

Methods

172 samples were collected from 50 poultry establishments described as 'free range' operations. None of the enterprises either prior or during the study reported necrotic enteritis in their poultry. Samples for bacterial isolation were grown on tryptose sulphite cycloserine (TSC) agar and quantified; typical colonies of C. perfringens putatively identified and confirmed biochemically. The samples were subjected to toxin production identification, ribosome typing, PFGE and antibiotic susceptibility testing.

Conclusions

The presence of cpa, cpb, etx, iA, cpb2, cpe, and NetB toxin genes were determined by PCR. 89.7% C. perfringens isolates were classified as type A and the remaining 10.3% type C; no other toxin genes were identified. Quantitative estimates of numbers of isolates was higher than expected (from $10^4$ to $10^9$ CFU/g) from poultry...
farm sources, regardless of the cleaning protocol used to sanitise the housing environment. Attempts to correlate specific clusters of *C. perfringens* with specific locations will be presented.
MSSA AND MRSA CO-COLONIZATION DYNAMICS AND CLONAL DIVERSITY IN PIGS

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Background
Methicillin-susceptible Staphylococcus aureus (MSSA) and methicillin-resistant Staphylococcus aureus (MRSA) are colonizers of skin and mucosa. Studies about MSSA and MRSA in human patients showed a competition of these two microorganisms for colonization space in the anterior nares. Moreover, in humans one clone can be found rather than differing types of MSSA and MRSA. At present, it is unknown whether this is also true for animals.

Objectives
The aim of this study was to investigate the colonization dynamics of both MSSA and MRSA in pigs over a longer time period and investigate their clonal diversity. The results might be of interest for the development of control strategies. This is important as in farm animals, and particularly in pigs common MRSA are found that are transmissible to humans and the environment.

Methods
Eighteen pigs were sampled three times every ten weeks with a nasal swab. Additionally, environmental samples were taken. All samples were investigated for MSSA and MRSA, respectively. Spa-typing was done with up to five MRSA and MSSA isolates, respectively, found per sample and time point. Of almost 400 MSSA and MRSA isolated, 62 isolates were further investigated by microarray.

Conclusions
The results do not support the hypothesis of a competitive colonization of MSSA and MRSA in the anterior nares of pigs. Rather we found a changing status. Hence, highly identical clones of MSSA and MRSA are present in the anterior nares of pigs and their environment. CC398 and CC9 associated spa-types were the predominating clonal lineages found among MRSA and MSSA isolates, respectively.
VAGINAL BACTERIOLOGY RESULTS OF THE EWES INDUCED WITH DIFFERENT PARTURITION INDUCTION METHODS

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Background

Induction of parturition can result in some infective postpartum vaginal complications in ewes due to their side effects on parturition. Therefore, it is important to reveal whether induction methods responsible to the vaginal bacterial changing in ewes.

Objectives

This study aims to reveal vaginal bacteriology and antibiogram results of the ewes induced with different parturition induction methods.

Methods

Twenty-four curly-fleeced ewes (n=24) at 3rd pregnancy period on 138th day were divided as group I (n=6): control; group II (n=6): dexamethasone (16 mg, im); group III (n=6): aglepristone (5mg/kg, im) and group IV (n=6): dexamethasone (8mg, im) + aglepristone (2.5 mg/kg, im). For bacteriological examination, the vaginal swaps were taken from ewes at 138th pregnancy day and then they were induced as below. Parturitions had not any clinical complications. Vaginal swaps were repeated on days of postpartum 15 and 30. Bacterial identifications revealed that *Escherichia coli* was the dominant pathogen bacteria both pre-induction and postpartum days in all groups. In postpartum 15th day, *Acinetobacter* spp. was the other pathogens for only in group III. And in postpartum 30th day, *Acinetobacter baumannii* was the other pathogen in group III. Antibiotic susceptibility of these pathogens was performed by Kirby Bauer disc diffusion methods. *Escherichia coli* and *Acinetobacter baumannii* were 100.0% sensitive to enrofloxacin, *Acinetobacter* spp. was 100.0% sensitive to trimethoprim / sulfamethoxazole.

Conclusions

It has been considered that induction of parturition can affect the vaginal flora and can be responsible for the postpartum vaginal complications to be encountered.
CHARACTERISTICS AND ANTIMICROBIAL SUSCEPTIBILITY OF TRUEPERELLA PYOGENES ISOLATED FROM BOVINE MASTITIS CASES IN CHINA

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Background
Trueperella (T) pyogenes is an opportunistic pathogen causes suppurative diseases in dairy cows. However, the pathogenesis and antibiotics resistance of T. pyogenes are still unclear.

Objectives
T. pyogenes isolates morphological characteristics, the presence of genes (plo, cbpA, fimA, fimC, fimE, fimG, nanH, nanP, tet/Wi, erm/Xi, erm/Bi), biofilm formation, the cytopathological effects in intracellular assay and antimicrobial resistance were investigated.

Methods
T. pyogenes was isolated from 50 out of 275 clinical and subclinical bovine mastitis cases in China. Both pyolysin (plo) and collagen-binding protein (cbpA) virulence factor genes were detected by conventional PCR in all T. pyogenes isolates. The tissue culture plate method was used to assay the capacity of T. pyogenes for biofilm formation and showed that 90% of T. pyogenes isolates were able to form biofilms with different production amount. Minimum inhibitory concentrations (MICs) of 14 antimicrobial agents were determined and observed high susceptibility to rifampin (96%), while high resistance to trimethoprim–sulfamethoxazole (90%) and bacitracin (98%). Intracellular assay revealed that 4 different T. pyogenes isolates had different cytopathological effects on infected bovine mammary gland epithelial cells.

Conclusions
18% T. pyogenes isolates indicates that T. pyogenes is important contributors to bovine mastitis. Moreover, the within-host quantitative, spatial, high occurrence of multidrug-resistant, biofilm producing and temporal dynamics of T. pyogenes interactions are key factors to better understand how immunity acts on infections with bacteria and how they evade immune surveillance; thus, highlighting the need for prudent use of antimicrobial agents in veterinary medicine (This research was supported by projects No. 2012BAD12B03, No. 313054, No. 20120008110042, No. 2014M561102 and No. GDT20141100043).
BRUCELLOSIS IN GREY SEAL IN BALTIC SEA

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Background

Finnish Food Safety Authority Evira participates in the HELCOM monitoring program to assess the health status of seals in the Baltic Sea area. Organ samples are collected from hunted seals or, to a lesser extent, from seals found dead. In the year 2013 we detected for the first time brucellosis in one of these samples. There are two marine Brucella species, B. pinnipedialis affecting pinnipeds (sea lions, walrus and true seals) and B. ceti affecting cetaceans (dolphins, porpoises and whales). Brucellosis is a zoonosis.

Objectives

The aim of this study was to examine the distribution and possible infection routes of brucellosis in grey seals in the northern Baltic Sea.

Methods

Livers of grey seals (Halichoerus grypus) (n=95) were inspected for parasites and pathological lesions. Samples for bacteriological examination (n=18) were taken from livers with macroscopic inflammatory lesions (n=10) and from normal livers (n=8) for control and cultivated on fastidious anaerobe agar plates. The plates were incubated at 37 °C in microaerophilic atmosphere, and observed for bacterial growth daily for 10 d. The identification was based on Stamp stain, biochemical tests and PCR.

Conclusions

Brucellosis does not seem to be rare in grey seals in the northern Baltic Sea. Of 10 samples with lesions 3 were positive. All normal livers were negative. The bile ducts of positive livers contained abundant numbers of flukes (Pseudamphistomum truncatum), which were positive for Brucella sp. in PCR. Flukes might be one of the possible vectors of brucellosis in marine environments.
Background

*Mycobacterium avium* subspecies *paratuberculosis* (MAP) is an obligate, Gram positive, acid-fast bacterium which has adapted to the gastrointestinal tract of ruminants and other animals and can cause fatal inflammatory disorder called Johne’s disease. The potency of microorganism to infect human and its close association with Crohn’s disease is the main public health issue of the infection, worldwide.

Objectives

The aim of this study was to find out some information about the frequency of MAP in slaughtered animals and further characterization of its genome, in Southern Iran.

Methods

The study was conducted between October 2013 to August 2014 in which total of 450 specimens from ileocecal lymph node, ileocecal valve and surfaces of 150 slaughtered sheep were collected. The animals were categorized according to the sex and age. IS900 PCR assay was employed to confirm the positive cases.

Conclusions

Out of 450 specimens, 12 were found positive using IS900 PCR assay. All the positive samples were from the intestinal tissues and lymph nodes. Statistical analysis of the results, using Fisher's Exact test, confirmed no significant difference between the frequency of infections in the ram and ewe. Furthermore, infection in the lambs was significantly lower than sheep over 1.5 year age. In general, employing the IS900 PCR assay is recommended to monitor the MAP infected carcasses. The sequence analysis of the positive cases showed a 93% identity when it was compared with gene bank database.
Background
Ear infections are a common problem in dogs. The infections can be very persistent and difficult to treat. The common treatment is with antibiotics and in chronic infections the dogs go through multiple treatments.

Objectives
The aim of this study was to map the most common causes of ear infections in dogs and test their resistance to antibiotics.

Methods
In the period of 2010 to 2013 the Department of Microbiology received 246 samples from dogs for microbial culture. Of these 93 were ear samples and we managed to isolate pathogenic bacteria and/or fungi from 83 of dogs. In the study period the main cause of ear infections in dogs' ears was the bacterium Staphylococcus intermedius. In addition we commonly isolated Proteus mirabilis, Pseudomonas aeruginosa and Candida spp. The antibiotic sensitivity of the isolated pathogens varied considerably and we often saw multiresistant Proteus mirabilis and Candida spp.

Conclusions
Dogs with ear infections are usually treated with antibiotics as soon as the infection is detected. Our study shows that multi-resistant bacteria are common in dogs with recurrent ear infections. It is very important to detect the pathogen and obtain information about antibiotic sensitivity before starting treatment. This will ensure more effective treatment and possible prevent chronic ear infections caused by multiresistant bacteria.
GENOTYPIC RELATEDNESS OF SALMONELLA ENTERICA SEROVAR ENTERITIDIS ISOLATES FROM CHICKENS AND HUMANS IN KOREA

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Background
Salmonella enterica serovar Enteritidis is one of the most common serotypes implicated in Salmonella infections in both humans and poultry worldwide. Most Salmonella infections in humans are considered to be associated with foods of animal origins contaminated by Salmonella. Indeed, the most common sources of Salmonella Enteritidis infection are believed to be chickens and chicken products.

Objectives
This study was to estimate the genetic relationship between Salmonella Enteritidis isolates from chickens and humans in South Korea.

Methods
A total of 127 chicken isolates were collected from clinical cases, on-farm feces, and slaughtered chicken meat between 1998 and 2012 and 20 human clinical isolates were obtained from patients who showed clinical signs between 2000 and 2006 in Korea. We compared pulsed-field gel electrophoresis (PFGE) patterns and multilocus variable-number tandem-repeat analysis (MLVA) profiles of the isolates.

Conclusions
PFGE analysis with all the isolates revealed 28 patterns and MLVA identified 16 allelic profiles. PFGE and MLVA showed common types shared by most human and chicken isolates although there was relatively poor congruence between the two analyses. Overall findings demonstrate the transmission of the serovar Enteritidis between humans and chickens.
Background

Canine leishmaniasis is a zoonotic disease by the protozoan parasite *Leishmania* transmitted by the bite of an infected phlebotomine sandfly. *Leishmania infantum* is the most common and important cause of canine leishmaniasis worldwide. Other *Leishmania* spp. reported from dogs include *L. mexicana*, *L. donovani*, and *L. braziliensis*. Leishmaniasis can be categorized by two types of diseases in dogs: a cutaneous reaction and a visceral reaction also known as black fever, the most severe form of *leishmaniasis*.

Infection does not invariably lead to illness. In fact, most infected dogs remain asymptomatic and may never develop clinical manifestations. In endemic regions, the prevalence of disease is often less than 10% and only about 1 in 5 infected dogs are considered likely to develop clinical disease.

Diagnosis of canine leishmaniasis is based on the presence of clinical signs together with positive specific antibody assay.

Objectives

The aim of this work was to develop a serological ELISA assay to detect IgG and IgM antibodies against Leishmania in serum or plasma samples derived from all mammals.

Methods

Microtiterplates were coated with antigen preparations of *Leishmania infantum*. The presents of antibodies against Leishmania is detected by protein A/G-HRP. A sample collection of about 200 positive samples and 400 negative samples was used for development and evaluation of the assay.
Conclusions

Here we show the performance characteristic of the newly developed assay. Due to the improved antigen design, purification method and test setup a superior assay performance was achieved compared to other test methods.
Background
Bovine mastitis was one of the common and frequent disease in cows, caused huge economic losses to dairy industry.

Objectives
The objective of this study was to investigate the distribution of bovine mastitis pathogens and their antibiotic resistance in China.

Methods
A total of 3941 milk samples from clinical mastitis cows (n=963), subclinical mastitis cows (n=2772) and healthy cows (n=206), which were collected from 40 cities in 150 dairy farms in China. Bacteria were isolated and identified. And disc diffusion test (K-B method) was used to determine the antibiotic resistance of the main pathogens, including Streptococcus agalactiae, Streptococcus dysgalactiae, Staphylococcus aureus and Escherichia coli.

Conclusions
The results showed that 24 species of bacteria and fungi include 4337 microorganisms were isolated from 3445 milk samples (87.41%), the other 496 milk samples were detected as negative ones. A total of 2386 bacteria were identified as bovine mastitis related pathogens: Streptococcus agalactiae (35.54%), Streptococcus dysgalactiae (23.55%), Staphylococcus aureus (16.89%), Escherichia coli (12.99%), breast Streptococcus (5.28%), klebsiella (1.63%), Proteus (1.59%), Pseudomonas aeruginosa (1.22%), stellite Nocardia (0.59%), suppurative Corynebacterium (0.38%), Candida albicans (0.17%) and Streptococcus pyogenes (0.17%). the total pathogens detection rate was 62.12%. Drug susceptibility test showed that the four major pathogens have different resistance to all antibiotics, the resistance rate was 10%~100%. Especially to penicillin G, streptomycin, and cotrimoxazole respectively, the resistance rate was 30%~100%.
O SEROGROUPS OF ENTEROPATHOGENIC (EPEC) AND SHIGATOXIGENIC (STEC) ESCHERICHIA COLI FROM

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Background

*Escherichia coli* producing the attachment-effacement (AE) lesion (EPEC) and/or Shiga toxins (STEC) cause enteritis and (bloody) diarrhoea in young calves and in humans, and are also present in the intestines of healthy cattle. Besides the O157:H7 serotype, EPEC and STEC can belong to more than sixty O serogroups. Of them, 8 have been most frequently identified worldwide: O5, O26, O103, O111, O118, O121, O145 and O165, with some also causing diarrhoea in young calves.

Objectives

This study aimed at identifying the pathotypes and the O serogroups of STEC and EPEC isolated from <1-month-old diarrhoeic calves in Wallonia, Belgium.

Methods

(i) 233 enterohaemolysin-producing *E. coli* were isolated at ARSIA between November 2008 and February 2014 from diarrhoeic calves after growth on EHLY Medium®. They were tested with a triplex PCR targeting the *stx1*, *stx2* (Shiga toxins) and *eae* (AE lesion) genes.

(ii) triplex PCR-positive *E. coli* were assayed with two pentaplex PCR targeting the specific genes coding for the nine O serogroups listed above and for the O104 serogroup.

Conclusions

(i) 206 isolates tested positive with the triplex PCR. The most frequent pathotypes were *eae+stx1+* (102 isolates), *eae+* (78 isolates) and *eae+stx1+stx2+* (13 isolates).
(ii) the most frequent serogroups of EPEC and STEC were O26 (57 isolates) and O111 (36 isolates). A few additional isolates tested positive for the O103, O5, O145, O121 and O157 serogroups.

(iii) the future is to compare these EPEC and STEC with those isolated from healthy cattle and from humans, to identify host- and age-specific properties.
COMPARISON OF BACTERIOLOGICAL AND GENETIC IDENTIFICATION OF MAMMARY PATHOGENS IN MILK SAMPLES FROM COWS WITH CHRONIC MASTITIS IN WALLONIA, BELGIUM

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Background

Mastitis is the inflammation of the mammary gland following infection by most often bacteria. Clinical mastitis is characterized by macroscopic modification of the milk while only high cell counts observed during subclinical mastitis. According to the bacterial species, mastitis can also be subdivided as either contagious (Staphylococcus aureus, Streptococcus agalactiae, Mycoplasma bovis, coagulase-negative staphylococci, Corynebacterium sp., etc) or environmental (Escherichia coli and coliforms, Streptococcus uberis, Streptococcus dysgalactiae, Enterococcus sp., etc).

Objectives

The overall objective of this study is to compare the results of detection of mastitis-associated bacteria by classical bacteriology and by quantitative (q) PCR.

Methods

A total of 120 milk samples were collected from mammary gland quarters with high cell counts (> 300,000 Somatic Cell) between January and March 2014 in seven farms in Wallonia. Samples were inoculated onto Columbia 5% sheep blood, McConkey's, Chapman's and modified Edwards' agar plates. Isolated colonies were picked up and stored at -80°C till further use. After new growth, colonies were Gram stained and identified using appropriate API sets.

Conclusions

A total of 231 isolates grew on Columbia blood agar. Of them, 133 isolates also grew on Chapman's agar including 63 S. aureus and 63 CNS; 70 isolates on Edwards' agar including 14 S. agalactiae, 12 S. uberis, 6 S. dysgalactiae and 10 Enterococcus sp.; and 15 isolates on McConkey's agar including 3 E. coli, 3 Klebsiella sp. and 8 Pantoea sp. These results are being compared with the results of the qPCR detection assay.
Background
Mastitis is one of the most frequent and costly disease in dairy cattle caused by different bacteria.

Objectives
The purpose of this study was to determine and compare the prevalence of different staphylococci, their antimicrobial susceptibility and toxin production, obtained from mammary infected cows in Kosovo.

Methods
Total 152 cow’s samples with clinical mastitis were collected. Cows were of different breeds and different lactation. Eight different commonly used antibiotics were analyzed for antimicrobial activity against staphylococci while SET-RPLA kit for enterotoxin detection.

Conclusions
Our results showed that, staphylococci were present in 89 out of 154 cows from which 58 were coagulase negative and 31 coagulase positive. S. aureus was most dominant (27/89) followed by S. epidermidis (25/89) and S. chromogenes (15/89). Interestingly, staphylococci where found more frequent in some dairy cow breeds than others. Most of the strains (76/89) were resistant to two or more different antibiotics. The highest resistance was observed against penicillin and ampicillin (>65%). While the lowest resistance was against erythromycin (<3%). Regarding the enterotoxin production, 40 out of the 89 strains produced at least one of the toxin types with SEA and SEC being the most common. In Kosovo is very common antibiotic misuse because of lack of local laboratories, hence the study suggests that, the uncontrolled use of antibiotics in treatment of different infective diseases might cause resistance against many antimicrobials. Therefore, attention should be paid on selection of antibiotics; timing of treatment, dosage and the uncontrolled and careless use of antibiotics should be more strictly regulated.
MALDI-TOF AS A TOOL IN IDENTIFICATION OF AVIAN MYCOPLASMA SPECIES
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Background

MALDI-TOF is a key method in identification of microorganisms. Large databases of reference and field strains have been made commercially available. Identifying Mycoplasma species has always been a time-consuming process using serological tests and molecular methods which require a large number of primers.

Objectives

Presently, only 3 avian Mycoplasma species are available in commercial databases. To evaluate if MALDI-TOF is a useful tool, first a larger database needed to be constructed. Secondly, strains isolated from infected birds were compared and matched against the database.

Methods

Fourteen reference and 8 field strains, obtained from collections held at our facility and from the University of Liverpool were grown in broth. An ethanol/formic acid extraction method was performed on each strain and then spotted on a MALDI target plate. For each strain, minimum 20 mass spectra of high quality were acquired to create the MSP using MALDI Biotyper 3.1 software. Forty-nine clinical isolates from poultry and wild birds were matched. Twenty-seven were correctly identified with a matching score of ≥ 2.0, 6 with a score between ≥ 1.7-1.99 and 16 remained unidentified.

Conclusions

MALDI-TOF is a rapid and useful tool in identifying Mycoplasma species. Fourteen isolates did not match probably due to missing strains in the database or possible unknown Mycoplasma species. More clinical isolates will be tested in the future and all non-identified isolates will be verified with 16S sequencing.

Acknowledgments

This study was funded by the Federal Public Service of Health, Food Chain Safety and Environment (RT12/4 Mycoplasma)
Background

Bovine mastitis caused by *Prototheca* can assume high significance because of economic losses and the potential risk to public health. Many studies highlighted the high level of antibiotic resistance of *Prototheca*, as well as the unsuccessful therapies.

Objectives

To determine the *in vitro* susceptibility of *Prototheca* isolates to different antibiotics and alternative treatments based on essential oils.

Methods

Sixteen isolates of *Prototheca* were identified by multiplex PCR reaction, typed using RAPD-PCR, and tested by the disk diffusion method for their susceptibility to 28 antibiotics. The efficacy tests to essential oils (thymol, carvacrol and cinnamaldehyde) were performed using emitters releasing compounds in a controlled way in jars, containing the *Prototheca* strains streaked on Sabouraud agar.

Conclusions

All the strains were resistant to Amikacin, Ampicilin, Aztreonam, Cefepime, Ceftazidime, Chloramphenicol, Ciprofloxacin, Erythromycin, Fosfomycin, Imipenem, Levofloxacin, Meropenem, Mupirocin, Nitrofurantoin, Oxacillin, Penicilin G, Piperacillin, Quinupristin/Dalfopristin, Rinfamicin, Streptomycin, Tetracycline, Ticarcillin/clavulanic acid, Tobramycin and Vancomycin. In contrast, Colistin sulphate, Gentamicin, Kanamycin and Netilmicin were effective against all the strains tested.

The growth of *P. zopfii* genotype 1-2, and *P. blasckaeae* was completely inhibited by thymol, carvacrol and cinnamaldehyde.

The results of the study revealed 4 out of 28 antibiotics tested efficient against *Prototheca* spp. while all the strains were inhibited by the tested essential oils.
Thus, *in vivo* studies are needed to evaluate if these compounds could rise as alternative treatments for bovine mastitis caused by *Prototheca* spp.
CHARACTERIZATION OF VIRULENCE FACTORS AND IMMUNE-STIMULATING ANTIGENS IN VIBRIO SALMONICIDA

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Background

Vibrio salmonicida is the cause of cold-water vibriosis in farmed Atlantic salmon. The disease manifests as a hemorrhagic septicemia, with a mortality ranging from 3-90 %. In the 1980s, the disease was a major challenge to the Norwegian aquaculture industry, but it is now being effectively controlled by the use of oil-based vaccines. Nevertheless, in recent years there have been some reports of disease outbreaks. In 2013, 13 Norwegian fish farms were diagnosed with cold-water vibriosis, all localized in the three northernmost counties of Norway.

Objectives

The knowledge on virulence factors of Vibrio salmonicida and the subsequent immune response of the fish is limited. As bacteria adapt to their current environment, a bacterium isolated from a diseased fish will not be identical to the same species cultivated in vitro. The aim of this study was to explore the phenotype of bacteria present in vivo.

Methods

We have grown Vibrio salmonicida in semi-permeable implants in live fish and analyzed bacterial protein expression by two-dimensional gel electrophoresis and tandem mass spectrometry. The experiment was approved by the Norwegian Animal Research Authority (approval no. ID6228).

Conclusions

Our findings may contribute to the knowledge on disease progression in cold-water vibriosis, as well as to provide novel targets for more effective vaccines. Also, the
study may provide insights in the course of disease for other bacterial infections in fish.
IDENTIFICATION AND TYING OF LACTOCOCCUS GARVIEAE BASED ON MALDI TOF MS

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Background

MALDI-TOF-mass spectrometry (MS) has emerged as a powerful technique for the routine identification of microorganisms in clinical microbiology laboratories allowing an easier and faster diagnosis than conventional phenotypic and molecular methods. We have evaluated the suitability of MALDI-TOF for the rapid identification of \textit{Lactococcus garvieae} the etiological agent of lactococcosis, a septicemic infection affecting different wild and farmed fish species, and recognized also as an opportunistic emerging human pathogen.

Objectives

Assessment of the reliability of MALDI-TOF MS for the identification of \textit{Lactococcus garvieae}

Methods

A total of 35 isolates from different host and environments were obtained from the culture collection of VISAVET were used in the study. The identification results obtained by MALDI TOF were compared with those obtained by biochemical identification (API 32 Strep) and by a species- specific PCR assay (Zlotkin et al. 1998. J Clin Microbiol 36, 983-985). Mass spectra acquisition and analysis was performed on a Bruker UltraFlextrem platform (Bruker Daltonics) using MALDI Biotyper™ 3.0 software in the automatic mode using a matrix of saturated solution of α-HCCA. In addition, a subset of isolates was subjected to PFGE.

Conclusions

The majority of peaks were obtained in the range from m/z 2000 to 10000. The proteomic results matched with those of genotypic approach. Our results also showed some differences in MS spectra in \textit{L. garvieae} isolates recovered from different origins/hosts suggesting the possible identification of molecular biomarkers for \textit{L. garvieae}. Our study demonstrates that proteomics identification using MALDI TOF MS could be a reliable approach for identifying and discriminating this microorganism.
IDENTIFICATION OF ARCANOBACTERIUM PLURANIMALIUM BY MALDI-TOF MS, BY SEQUENCING 16S RDNA AND GENE PLA AND BY DEVELOPMENT OF A LOOP-MEDIATED ISOTHERMAL AMPLIFICATION (LAMP) ASSAY

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Background

Genus Arcanobacterium consists of four species namely A. pluranimalium, A. haemolyticum, A. phocae and A. hippocoleae. More recently, A. canis and A. phocisimile were described as novel species. A. pluranimalium was initially recovered from a harbour porpoise and a fallow deer, from a dog with pyoderma, from abortion material, semen, abscesses, viscera, navel illness and peritonitis of sheep and from milk samples of cows with mastitis. More recently, an A. pluranimalium strain was recovered following necropsy of a juvenile giraffe. However, the clinical importance of A. pluranimalium in animal infection remains unclear.

Objectives

Because of this widespread presence of A. pluranimalium in clinical samples of various animals there is an urgent need of phenotypic and genotypic tests for a rapid and reliable identification of this bacterial species.

Methods

The A. pluranimalium strains investigated in the present study were analysed phenotypically, by MALDI-TOF MS analysis and by 16S rDNA sequencing. In addition, the previously sequenced gene pla encoding pluranimaliulysin of A. pluranimalium was used for molecular identification of this bacterial species and for the development of oligonucleotide primers for a loop-mediated isothermal amplification (LAMP) assay.

Conclusions

Phenotypic methods, MALDI-TOF MS analysis, sequencing 16S rDNA and target gene pla and the newly designed pla LAMP assay of the present study allowed the identification of all nine investigated A. pluranimalium strains of ovine and bovine
origin and from a juvenile giraffe. This phenotypic and genotypic approach might help to elucidate the role this species plays in animal infections.
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SEQUENCING AND PHYLOGENETIC ANALYSIS OF GP51 GENE OF BOVINE LEUKEMIA VIRUS ISOLATES IN KOREA IN 2014
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Background

Enzootic bovine leukosis was first reported by serological method in Korea as early as in 1980s and recent prevalence in dairy cattle was over 50%. So far there has been no report on the bovine leukemia virus (BLV) genotype circulating in Korea since partial env gene sequence which belonged to genotype 1 was deposited in 2009.

Objectives

This study was performed to identify the genetic diversity of BLV in Korea.

Methods

Total 185 whole blood samples were collected from dairy cattle in two farms in southeast region in Korea. BLV gp51 genes were amplification for diagnosis as described previously by Fechner et al. Gp51 full gene (903bp) for sequencing was amplified. The gp51 PCR products were purified, sequenced and analyzed using DNAstar program and Mega 6 Program.

Conclusions

Total 185 samples were analyzed with PCR, and total 78 samples were positive for BLV env DNA nested PCR (37 samples out of 119 in farm A and 41 out of 66 in farm B). Further PCR reaction to amplify full gp51 gene was performed. 19 out of 37 positive samples from farm A and 31 out of 41 positive samples from farm B were amplified successfully. The genotype of sequences from farm A was divided into two types, genotype 1 and genotype 3. All 31 isolates from farm B were found to belong to genotype 1. This is the first report of BLV genotype 3 circulation in Korea.
DEVELOPMENT OF COMPETITIVE ELISA FOR THE DETECTION OF ANTIBODIES AGAINST SEVERE FEVER WITH THROMBOCYTOPENIA SYNDROME VIRUS (SFTSV) IN SUSCEPTIBLE ANIMALS

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Background

The Severe Fever with Thrombocytopenia Syndrome (SFTS) caused by SFTS virus (SFTSV) which belongs to the phlebovirus in bunyaviridae family is an emerging tick-borne mediated infectious disease for human, and the fatality rate is 12% in China. SFTSV seems to have wide host range in that SFTSV positive ticks were isolated from farm animals and wild rodents. Therefore, it is important to monitor SFTSV positive animals to prevent circulation or transmission of SFTSV.

Objectives

As there is no gold standard method to detect SFTSV specific antibodies from field animals, we are developing competitive enzyme-linked immunosorbent assay (c-ELISA) technique using nucleocapsid protein (NP) of SFTSV.

Methods

The recombinant NP of SFTSV was obtained by the expression of cloned genes in E. coli and the SFTSV was amplified in Vero cells for 7 days. The purified recombinant protein or formalin-inactivated SFTSV was used to immunize laboratory animals. Mono- and polyclonal antibody were tested for their antigen specificity in immunofluorescence assay (IFA) and western blot.

Conclusions

We developed direct c-ELISA using SFTSV specific monoclonal antibody, and generated the SFTSV positive serum from immunized cattle. The c-ELISA result shows 98.08% agreement in IFA result using 418 cattle field serum samples. We also observed the cELISA is applicable to detect SFTSV positive sera from immunized cattle, chicken and goat. This technique could be usefully used as a serological diagnostic method for various species of SFTSV susceptible field animals.
PHYLOGENETIC ANALYSIS AND MOLECULAR CHARACTERIZATION OF MYCOPLASMA BOVIS OUTBREAK STRAINS IN DENMARK

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Background

*Mycoplasma bovis* is a cattle pathogen causing broad spectrum of diseases including mastitis, pneumonia and arthritis. The infection is very difficult to control due to long-time persistence of the organism in asymptomatic carriers, lack of knowledge on the specific role of environmental factors in the development of clinical disease, poor response to antibiotic treatment and the lack of effective immunotherapy.

Objectives

Here we report whole-genome sequence analysis of 41 *Mycoplasma bovis* isolates collected during recent outbreak of disease in Danish cattle herds. Our objectives were to determine the extent of genomic diversity in the test population, identify dominant clones, reconstruct their phylogenetic ancestry and infer transmission history.

Methods

Genome sequencing was performed on an Ion Torrent PGM sequencer. Genomic libraries were prepared by using Ion Xpress Plus fragment library kit and amplified using Ion PGM Template 200 OT2 kit according to manufacturer’s instruction. Raw sequence reads were pre-processed using FastQC, mapped to reference genome (PG45) using Burrows-Wheeler aligner and single nucleotide polymorphisms (SNPs) were identified using REALPHY server.

Conclusions

Unique molecular markers were identified for individual strains. Phylogenetic clustering based on SNPs revealed two distinct clades indicating a recent common ancestry. Substantial variations in both the number and sequence identities of individual variable surface proteins were observed with respect to type strain PG45 that may have contributed to increased virulence. The genetic distances inferred from SNPs and temporal distances (sampling dates) showed a strong
negative correlation suggesting that introduction of strains into the herds took place much before sampling time.
Background

Bovine digital dermatitis (BDD) and Contagious ovine digital dermatitis (CODD), are infectious foot diseases causing severe lameness in cattle and sheep, respectively. The causative agents of digital dermatitis (DD) in dairy cattle are considered to be Treponema bacteria, however little work has been done regarding DD in beef cattle. Similarly, although it is suggested CODD emerged from BDD little has been done to investigate the aetiology of the disease in sheep or the transmission of DD.

Objectives

1. Isolate and characterise beef cattle and sheep DD treponemes from lesions and compare with dairy cattle DD treponemes.
2. Identify reservoirs of infection of DD.

Methods

Beef cattle and sheep DD lesions were subjected to culture and PCR analysis for the commonly detected DD treponemes from dairy cattle lesions.

The gastrointestinal (GI) tract of beef cows and sheep was investigated as an infection reservoir of DD treponemes by analysing GI tract tissues for the presence of treponemes. Farm equipment which comes into contact with lesions was analysed by PCR and culture methods for the presence of treponemes.

Conclusions

From 16S rRNA gene analysis, treponemes isolated from beef cattle and sheep lesions belonged to the groups of treponemes associated with dairy cattle lesions. The close similarity poses concerns for species-species transmission.

Several GI tract tissues were positive for DD treponemes, providing evidence towards the GI tract as an infection reservoir for DD treponemes. Additionally, farm equipment
was also found to harbour DD treponemes after coming into contact with an infected foot indicating a possible transmission route.
PROTECTING CHICKENS AGAINST CAMPYLOBACTER JEJUNI BY CAMELIDAE-DERIVED NANOBODIES

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Background

Campylobacter jejuni is a Gram-negative microaerophilic bacterium, and is the major cause of human gastro-enteritis worldwide. It is a zoonotic pathogen and chickens are generally accepted as the most important source of infections. Since the use of antibiotics in animal feeds is no longer an acceptable option and the prevention is limited to hygiene measures, novel control methods are needed.

Objectives

The purpose of this research is to investigate whether colonization of chickens by C. jejuni can be controlled by passive vaccination in a cost effective manner by using nanobodies, the antigen-binding domains of heavy-chain antibodies.

Methods

Anti-Campylobacter nanobodies were identified after immunization of an alpaca with heat-killed C. jejuni cells. The binding of these nanobodies on different C. jejuni strains, isolated from chickens and their environment as well as human clinical isolates, was verified. Nanobodies that are able to bind all of the tested strains were selected for further research. We presumed that these nanobodies recognize conserved epitopes expressed on C. jejuni. The ability of one of these nanobodies to agglutinate Campylobacter cells was tested. This nanobody was multimerised by linking it to magnetic beads. The results confirm that the nanobody recognizes antigens on living C. jejuni cells and that multimers of this nanobody are able to agglutinate the C. jejuni cells.

Conclusions
The nanobodies with a broad *Campylobacter* specificity can be interesting for diagnostic and therapeutic applications, such as for the reduction or inhibition of the colonization of chickens by *C. jejuni*. 
GENETICS AND BIOFILM ANALYSIS ON STRAINS OF E.COLI ISOLATED FROM ANIMALS WITH DIARRHOEIC PROBLEMS.

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Background

- *Escherichia coli* (*E. coli*) is a bacterium that commonly lives in the intestines of people and animals. There are four major categories of diarrheogenic *E. coli*, namely: enterotoxigenic *E. coli* (*ETEC*), enteroinvasive *E. coli* (*EIEC*), enteropathogenic *E. coli* (*EPEC*) and enterohemorrhagic *E. coli* (*EHEC*). These categories of *E. coli* differ in their epidemiology and pathogenesis and their O: H serotypes and for the presence or absence of several genes responsible for their virulence.

Objectives

- The objective of the study was the analysis of virulent genes and biofilm production in *E.coli* strains isolated from animals with diarrhoeic problems.

Methods

The strains were isolated from stool samples of animals affected by diarrhoea. Genetic analysis were performed by multiplex PCRs. Biofilm assays were performed by safranin method with O.D. readings at 492 nm on 96 wells plate.

Conclusions

The isolates were all coming from symptomatic animals from farms in which repeated episodes of diarrhoea are present. All isolates were negative for the *aggR* gene (Aggregative adherence fimbria), the *pet* gene (Plasmid-encoded toxin), *aap* (dispersin) and other genes. 10 out of 35 were positive for irp2 (yersinia bactin biosynthesis gene) and 18 out of 35 strains carried the *astA* gene which encodes for EAST1 (Entero Aggregative heat-StableToxin1). This protein is thought to play a role in EAEC pathogenicity even if not all EAEC strains harbour the *astA* gene. The analysis for biofilm formations on the first 22 isolates, showed low capability in biofilm formation. Only 8/22 showed values above 120 O.D., but further analysis will be performed with different stains.
DETERMINATION OF ANTIMICROBIAL RESISTANCE AND VIRULENCE GENES PATTERNS AMONG CAMPYLOBACTER FROM BOILERS IN POLAND

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Background

Campylobacter is the most common zoonotic agent deriving from broiler flocks and broiler carcasses in Poland. The increased resistance of this bacteria to antibiotics is a matter of special concern, representing a significant public health problem. The genetic features responsible for pathogenesis of campylobacteriosis are still under investigation.

Objectives

The aims of the study were to test Campylobacter strains for antimicrobial resistance patterns and to screen for the presence of virulence-associated genes involved in different steps of infection.

Methods

During the project, 29 broiler farms located in 15 voivodeships all over Poland were selected for sampling at the slaughterhouse level. In total, 252 isolates from caeca (150 C. jejuni and 102 C. coli) and 197 strains from the corresponding carcasses (142 C. jejuni and 55 C. coli) were tested. A microbroth dilution method was used to establish the antimicrobial resistance by the minimum inhibitory concentrations (MICs) to 7 antimicrobials. Additionally, the following genes: cadF, flaA, virB11, iam, flhA, cdtA, cdtB, cdtC, ciaB, docA and wlaN were detected by PCR.

Conclusions

The most common resistance profile of C. jejuni was ciprofloxacin-nalidixic acid and tetracyclines. C. coli were mostly resistant to ciprofloxacin-nalidixic acid-tetracycline and streptomycin. The results regarding virulence patterns demonstrated that the majority of C. jejuni strains had virulence and cytotoxin gene pattern: flaA, flhA, cadF, ciaB, cdtA, cdtB, cdtC and docA. Among C. coli, the simultaneous presence of the flaA, flhA, cadF, ciaB, cdtA, cdtB, iam and docA markers was most often found.
PRESENCE OF CAMPYLOBACTER JEJUNI IN DIFFERENT SPECIES OF ANIMALS, POSSIBILITY OF ITS INFLUENCE AS A RESERVOIR AND/OR VECTOR INFECTION OF BROILERS

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Background
Campylobacteriosis is the most common foodborne gastroenteritis in industrialized countries and the number of cases is growing continuously. This bacterial infection is mainly related to the consumption of poultry meat, when it is contaminated by *Campylobacter jejuni*. Although it is a known fact that *C. jejuni* has a highly prevalent in broiler flocks, the source of infection in herds remains still unclear. Nowadays, it would be interesting to knowledge the role of the different animals in the transmission of the infection

Objectives
The objective of this research was to determine the prevalence of bacteria in different types of animal species

Methods
Both birds and mammals remaining in direct or indirect contact with infected broilers for *C. jejuni* were analyzed. Individual animals were sampled by rectal swab and all samples were processed similarly, being the positive samples confirmed by PCR. This organism was isolated from all species that had been in direct contact with infected broilers, but it cannot be isolated from the animals not having direct contact with them.

Conclusions
According to the results obtained, it is concluded that *C. jejuni* can be transmitted between different animal species, acting as vector and / or reservoirs of this organism in broilers.
DETECTION OF HUMAN PAPILLOMAVIRUS DNA AND P53 CODON 72 POLYMORPHISM IN PROSTATE CANCER IN TURKISH POPULATION

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Background: The possible relationship between human papillomavirus infection, tumor suppressor gene p53 and prostate cancer (PCa) has been previously investigated, however, the etiological role of HPV infection and p53 codon 72 polymorphism in the PCa pathogenesis remains highly controversial.

Objectives: The aim of the study was to assess the possible role of human papillomavirus in the development of prostate cancer and the distribution of the p53 codon 72 polymorphism in Turkish population.

Methods: We investigated the presence of HPV in 60 formalin fixed and paraffin embedded prostate cancer tissues, as well as in 36 benign prostatic hyperplasia tissue (BPH) samples by using real-time PCR and pyrosequencing. Furthermore, the same group of patients were also screened for the presence of the Arg72Pro polymorphism of the p53 gene, a p53 polymorphism related to HPV.

Conclusions: No HPV DNA was detected while using real-time PCR in any of the 36 BPH samples. One of the sixty (1.67%) PCa samples was found to be positive for HPV DNA by using a real-time PCR. After using pyrosequencing analysis, the only HPV type detected in the PCa sample was HPV-57. Compared to the control group, Arg/Pro genotype was observed more frequently in prostate cancer patients (p=0.044), and also Pro allele frequency is higher in prostate cancer patients (p=0.021). Our findings do not support the involvement of HPV in the etiology of prostate carcinogenesis, and the proline allele could be a risk factor for the development of prostate cancer in the Turkish population.
Background

Viruses are the numerically dominant life forms on Earth. They represent a huge reservoir of genetic diversity, infecting organisms from small to large. In the oceans, every second $10^{23}$ viral infections take place.

Objectives

To fully comprehend the impact marine viruses can have on their host population dynamics and subsequently biodiversity, ecosystem functioning and global energy & matter fluxes, it is essential to study virus-host interactions under different environmental conditions.

Methods

Here I will present experimental results on physiochemical variables affecting algal virus growth characteristics, as well as field data showing how a changing ocean due to global warming can control the magnitude of viral lysis as compared to the traditional loss factor, grazing.

Conclusions

These examples illustrate the importance of environmental factors underlying the ecological importance of viral infection in marine ecosystems. Further research is warranted to identify more regulatory factors and understand the mechanisms by which they influence virus-host interactions.
Viral infections and host

RECOMBINANT LIPIDATED DENGUE ENVELOPE PROTEIN DOMAIN III INDUCES DURABLE NEUTRALIZING ANTIBODY RESPONSES AND PROTECTIVE IMMUNITY

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Background
The immunogen and immunopotentiator are two important components of vaccines. Combination of these two components into a single construct will have a great potential to increase the reactivity of immunogen. The lipid moiety of the recombinant lipoprotein provides a danger signal to activate antigen-presenting cells via toll-like receptor 2, which may further enhance immune responses.

Objectives
We selected dengue-4 envelope protein domain III (LD4ED III) as the immunogen and converted this candidate in lipidated form in Escherichia coli-based system. In the present study, the immunogenicity and protective immunity of LD4ED III in mice were evaluated.

Methods
Recombinant proteins were purified by immobilized metal affinity chromatograph columns. Groups of mice were immunized subcutaneously twice with 10 μg of D4ED III or LD4ED III in PBS at a 4-week interval. Mice immunized with PBS alone served as controls. Immune responses in mice were examined. All animal studies were approved and performed in compliance with the guidelines of the Animal Committee of the National Health Research Institutes.

Conclusions
The LD4ED III expressed in E. coli retains the proper conformation, which has the capability to compete with dengue virus for cellular binding sites. In addition, LD4ED III is more immunogenic than D4ED III to induce immune responses. Most importantly, LD4ED III elicits a durable neutralizing antibody response and inhibits viremia in LD4ED III vaccinated mice in the absence of exogenous adjuvant. These results suggest that lipidated dengue envelope protein domain IIIIs are potential dengue vaccine candidates.
THE ROLE OF ADENOVIRUS 36 AS A RISK FACTOR IN OBESITY: THE FIRST CLINICAL STUDY IN THE ADIPOSE TISSUE SAMPLES OF ADULTS IN TURKEY

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Background

Obesity which develops due to multifactorial reasons, was associated recently with human Adenovirus-36 (Ad-36).

Objectives

The aim of this study was to investigate the prevalence of Ad-36 antibodies in obese adults and also investigate the DNA of Ad-36 in their adipose tissue.

Methods

In this cross-sectional and case-control based study, 49 obese adults (BMI ≥ 30 kg/m²) and 49 non-obese adults (BMI ≤ 25 kg/m²) were included in this study as patient and control groups, respectively. Adipose tissue samples obtained by the lipoaspiration method, were studied by PCR methods. Simultaneously, the presence of Ad-36 antibodies and serum leptin and adiponectin levels were assessed by serum neutralization assay (SNA) and ELISA, respectively.
Conclusions

Ad-36 antibody was detected in 6 (12.2%) of 49 patients by SNA and was statistically significant (p<0.05). We couldn’t detect Ad-36 DNA in adipose tissue; however, we detected significantly higher Ad-36 antibody levels in the obese group compared to the non-obese group. Mean BMI and leptin levels were higher in the Ad-36 positive group, while adiponectin levels were found to be lower in the Ad-36 positive group. Although no statistically significant difference was found in cholesterol and triglyceride levels between the two groups (p > 0.05), lower mean serum cholesterol and triglyceride levels were found in the Ad-36 positive patients. There is a need for extended serial, particularly cohort and human-based, studies in order to have a clear understanding of the Ad-36-obesity relationship.
Viral infections and host

PREVALENCE OF ASTROVIRUS IN FECAL SPECIMENS OF CHILDREN WITH GASTROENTERITIS
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Background
Norovirus genogroup I (NV GI) and NV GII sapovirus (SaV) genogroups I, II, IV, and V, human rotavirus A(HRV), adenovirus serotypes 40 and 41 (AdV), and human astrovirus (AstV) are the most common causes of viral gastroenteritis and result in large outbreaks of viral diarrhea. An estimated 1.8 million children die each year from largely preventable enteric illnesses, with the large majority of these mortalities occurring in developing countries. Gastroenteritis is a major cause of morbidity and mortality in children worldwide.

Objectives
Astroviruses are one of the major viral pathogens responsible for gastroenteritis. In this study performs to determine the prevalence of astrovirus in fecal specimens of children with gastroenteritis.

Methods
In this study 2490 stool samples from children less than five years with gastroenteritis from five cities of Iran were collected. The prevalence of human astroviruses was tested in patients with acute gastroenteritis by using conventional duplex reverse transcription (RT) -PCR and electrophoresis.

Conclusions
Young children were much more likely to be admitted with acute gastroenteritis. In this study astrovirus was detected in 1.6% of samples. There is no relation between age, sex and type of virus. rotavirus was detected in 50% of astrovirus positive samples. rotavirus was detected with ELISA. astroviruses are an important viral agents of gastroenteritis in Iran. Because of co-infection of astrovirus, improved personal hygiene and rotavirus vaccination will reduce the incidence of co-infections.
Viral infections and host

DEAMINASE ACTIVITY OF APOBEC3 IS REQUIRED FOR EFFICIENT RESTRICTION OF PORCINE ENDOGENOUS RETROVIRUS

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Background

Porcine endogenous retroviruses (PERVs) present a unique concern associated with xenotransplantation because they have been shown to infect certain human cells in vitro. Human APOBEC3G (huA3G) is a single-strand DNA cytosine deaminase, which inactivates the coding capacity of the virus by deamination of cDNA cytosines to uracils. This reaction occurs within (-) DNA strand during reverse transcription, resulting in G-to-A mutation in the (+) strand.

Objectives

A recent report showed that huA3G also could inhibit viral replication by cytidine deaminase-independent mechanisms. Two contrasting results with regard to huA3G-dependent inhibition of PERV transmission have been reported. Some group suggests that the mechanism of inhibition does not require the DNA deaminase activity of huA3G. In contrast, other group report that PERV restriction is attributed to cytidine deamination.

Methods

To determine whether DNA deamination is required for huA3G-dependent inhibition of PERV transmission, we established a new set of 293-PERV-PK-CIRCE (human 293 cells infected with PK15-derived PERVs) clones stably expressing huA3G. In addition to huA3G, we included porcine APOBEC3G(poA3Z2-Z3) and murine APOBEC3(mA3). 293T cells were infected with virus-containing supernatant from 293-PERV-PK-CIRCE clones expressing huA3G, poA3Z2-Z3 and mA3. To search for antiviral mechanism of APOBEC3, genomic DNA from 293T cells was prepared at 10h postinfection and the PERV gag gene from 3D-PCR(differential DNA denaturation PCR) product was sequenced.

Conclusions

huA3G, poA3Z2-Z3 and mA3 induced G-to-A hypermutations in the PERV genome effectively. Based on these results, we conclude that the APOBEC3 restricts PERV replication by deaminase-dependent mechanism.
IS THERE ANY ROLE OF ADENOVIRUS 36 AS A RISK FACTOR IN GYNECOMASTIA? THE FIRST CLINICAL STUDY WORLDWIDE FOR PATIENTS WITH GYNECOMASTIA

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Background

Gynecomastia was indicated as a diseases which develops due to multifactorial causes.

Objectives

Based on adipose tissue affinity of Ad-36, we hypothesized, Ad-36 may also play a role in the development of gynecomastia by possibly accompanying increased regional adiposity.

Methods

In this cross-sectional and case-control based study, 33 adult male (BMI ≤ 25 kg/m²) and 15 adult male (BMI ≤ 25 kg/m²) with anatomic disorders without gynecomastia pathology were included in this study as patient group (PG) and patient control group (PCG), respectively. Breast reduction samples obtained by the lipoaspiration/subcutaneous mastectomy method, were studied by PCR methods.
Simultaneously, the presence of Ad-36 antibodies and serum leptin and adiponectin levels were assessed by serum neutralization assay (SNA) and ELISA, respectively.

**Conclusions**

Ad-36 antibody was detected in 8 (24.2%) out of 33 PG by SNA and was statistically significant (p<0.05). Mean leptin levels were higher in the Ad-36 positive group, while adiponectin levels were found to be lower in the Ad-36 positive group. We couldn’t detect Ad-36 DNA in breast reduction samples; however, we detected significantly higher Ad-36 antibody levels in the PG compared to the PCG. Ad-36 antibody positivity stayed in models in multivariate logistic regression analysis (with backward modeling) and has been identified as a risk factor. We report firstly Ad-36 may be a risk factor in gynecomastia. Moreover, with regard to leptin, adiponectin and serum cholesterol levels, our results suggest that there might be a relationship between Ad-36 and gynecomastia.
FEMS-2735
Viral infections and host

THERAPEUTIC POTENTIAL OF A NEW KLEBSIELLA PNEUMONIAE BACTERIOPHAGE: ISOLATION, GENETIC CHARACTERIZATION AND PHAGE DEVELOPMENT IN SLOWLY GROWING BACTERIA

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Background

Klebsiella pneumoniae is a facultative anaerobic Gram-negative bacterium associated with pyoderma in animals. K. pneumoniae can acquire resistance to carbapenems, thus becoming insensitive to almost all available antimicrobial agents. Phage therapy has the potential to replace antibiotic treatment. Lytic bacteriophages and/or their gene products such as lysins, can be easily used as therapeutic agents against bacteria as they are host specific and generally show no side effects.

Objectives

We focused on isolation and characterization of new bacteriophages against Klebsiella pneumoniae, which can be potentially used in phage therapy in animals.

Methods

We collected K. pneumoniae swab samples from the infected animals that could not have been treated with a standard antibiotic therapy. Afterwards, new lytic phage PRA33 was isolated from sewage and, with the use of electron microscopy analysis, it was structurally defined as a member of the Siphoviridae family. Phage plaque morphology analysis (the plaque clarity and the ability to increase its diameter in time) and determination of phage host range were carried out. Then, analysis of kinetics of adsorption and one-step growth curves were determined in bacteria growing with slow growth rates in a chemostat system. We also performed phage stability tests in various liquid media. Phage genome was sequenced de novo and the identification of putative ORFs was undertaken.

Conclusions
Based on the results of our experiments, we propose PRA33 phage to be a promising candidate for application in phage therapy against *K. pneumoniae* causing infections in animals that are otherwise difficult to treat.
DEVELOPMENT AND EVALUATION OF A SEROLOGICAL CHIKUNGUNYA ANTIBODY DETECTION ASSAY

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Background

Chikungunya (also named breakbone fever) is a highly emerging disease in many tropical settings with great socioeconomic impact. Chikungunya-Viruses are transmitted to humans by bloodsucking mosquitoes (Aedes aegypti, Aedes albopictus). The symptoms of Chikungunya include fever which can reach 39°C (102.2°F) a petechial or maculopapular rash usually involving the limbs and trunk, and arthralgia or arthritis affecting multiple joints which can be debilitating. The fever typically lasts for two days and abruptly comes down. Alphaviruses can be noticed as import or travel associated infection.

Objectives

The aim of this work was to develop an serological assay to detect IgG and IgM antibodies against Chikungunya and to evaluate in endemic outbreak settings.

Methods

An IgG-capture and IgM-capture ELISA was developed. Both take advantage of native antigens produced with a proprietary technique. In house measurements as well as external evaluations in many endemic regions of the world conducted by well know tropical institutes revealed excellent clinical sensitivity and specificity as well as high positive and negative predictive values. Data from the current outbreak in the Caribbean will be discussed.

Conclusions

The newly developed ELISA seems to be a superior tool to diagnose past and acute Chikungunya infection in common and outbreak settings all over the world. It will assist diagnosis of travel returners with unknown fever as well as military in endemic operation area. To further improve Chikungunya diagnostic a Lineblot is currently under development as tool for conformation of ELISA results as well as for small labs with limited lab equipment.
VACCINE-TYPE ATTENUATING MUTATIONS IN HIGHLY PASSAGED STRAIN OF VARICELLA ZOSTER VIRUS

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Background: Varicella-zoster virus (VZV) is causative agent for chickenpox and zoster. Live attenuated vaccines have been developed based on Japanese Oka strain and Korean MAV/06 strain. Comparison of the complete sequences between vaccine and wild-type strains suggested at least 24 mutations important for attenuation in vaccine strains. Certain strains with high in vitro passage history such as the strains Ellen and 32p72 contained some of the vaccine-type mutations.

Objective: In this study, we attempted to understand vaccine-type attenuating mutations in highly passaged VZV strains.

Methods: Genome sequences of clinical strains with different passages were determined by next-generation sequencing and compared with those of other VZV strains.

Conclusion: Vaccine-type mutations were found at position 106262 (T→C) in the Korean clinical strain YC01 passage 32 and at positions 560 (T→C), 105169 (A→G), 106262 (T→C) and 107252 (T→C) in YC01 passage 61. Similar mutations were also found in high-passaged another clinical strain YC02 at positions 560, 106262 and 107252. Same mutations at 106262 and 107252 were found in the strains Ellen and 32p72. Thus, in the course of in vitro passaging of VZV, mutations at 106262 and 107252 seemed to occur first, and followed by mutation at 560. Direct PCR sequencing of these positions in YC01 and YC02 strains at various passages identified the passage numbers when the attenuating mutations occurred. Further studies of in vitro passaging under attenuating conditions such as low temperature or guinea pig cells will help to understand the mechanism of attenuating mutations in VZV.
Viral infections and host

LIPIDATED DENGUE-2 ENVELOPE PROTEIN DOMAIN III INDEPENDENTLY STIMULATES LONG-LASTING NEUTRALIZING ANTIBODIES AND REDUCES THE RISK OF ANTIBODY-DEPENDENT ENHANCEMENT

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Background
Dengue virus is a mosquito-transmitted virus that can cause self-limiting dengue fever, severe life-threatening dengue hemorrhagic fever and dengue shock syndrome. The existence of four serotypes of dengue virus has complicated the development of an effective and safe dengue vaccine. Recently, a clinical phase 2b trial of Sanofi Pasteur’s CYD tetravalent dengue vaccine revealed that the vaccine did not confer full protection against dengue-2 virus. New approaches to dengue vaccine development are urgently needed. Our approach represents a promising method of dengue vaccine development and may even complement the deficiencies of the CYD tetravalent dengue vaccine.

Objectives
To develop a vaccine with long-lasting neutralizing antibodies and reduces the risk of antibody-dependent enhancement

Methods
Two important components of a vaccine, the immunogen and immunopotentiator, were combined into a single construct to generate a new generation of vaccines. We selected dengue-2 envelope protein domain III (D2ED III) as the immunogen and expressed this protein in lipidated form in Escherichia coli, yielding an immunogen with intrinsic immunopotentiating activity.

Conclusions
The formulation containing lipidated D2ED III (LD2ED III) in the absence of exogenous adjuvant elicited higher D2ED III-specific antibody responses than those obtained from its nonlipidated counterpart, D2ED III, and dengue-2 virus. In addition, the avidity and neutralizing capacity of the antibodies induced by LD2ED III were higher than those elicited by D2ED III and dengue-2 virus. Importantly, we showed that after lipidation, the subunit candidate LD2ED III exhibited increased immunogenicity while reducing the potential risk of antibody-dependent enhancement of infection in mice.
Background:

Enterovirus 71 (EV71), a positive-stranded RNA virus, is the major cause of hand, foot and mouth disease (HFMD) with severe neurological symptoms. Seven-day-old toll-like receptor 9 knockout (TLR9KO) mice infected with a non-mouse adapted EV71 strain developed neurological lesion-related symptoms, including hindlimb paralysis, slowness, ataxia and lethargy. Histopathological examination revealed a massive damage in the limb muscles, brain, and spinal cord. The TLR9KO mouse could be useful as EV71 animal model to study anti-EV71 drugs and vaccines.

Objectives:

A novel nanoparticle-based EV71 VP1 peptide vaccine was evaluated in the EV71 mouse model. The clinical symptoms of mice after EV71 infection were monitored.

Methods:

Seven-day-old TLR9KO mice were infected with non-mouse adaptive EV71 and showed neurological diseases. We found that EV71 alone cannot activate toll-like receptor 9 (TLR9) signaling. On the contrast, supernatant from EV71-infected cells could activate TLR9. Furthermore, we identified the EV71-infected cells-derived danger signal activate TLR9 to protect mice from EV71 challenge. To investigate whether TLR9KO mice could be used as an animal model for EV71 vaccine development, EV71-infected TLR9KO mice were treated with sera from vaccinated mice. We designed a VP1 peptide of EV71 that formulated with emulsion type nanoparticles adjuvant and immunization with wild type mice. Sera from immunized mice were administered to EV71-infected TLR9KO mice, and clinical neurological symptoms of mice were observed.

Conclusions:

Sera from EV71 VP1 peptide immunized mice could reduce neurological symptoms of TLR9KO mice from challenged with EV71.
ALGAL VIRUS PRODUCTION IS NEGATIVELY AFFECTED BY PHOSPHORUS AND NITROGEN LIMITATION, BUT THIS MAY BE COUNTERACTED BY NUTRIENT ASSIMILATION DURING INFECTION

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Background
Viruses are a main cause of phytoplankton mortality, thereby controlling their community composition and driving biogeochemical cycling in the marine environment. Phytoplankton growth and production is often limited by nitrogen (N) and/or phosphorus (P), which can affect the virus growth characteristics upon infection. The influence of nutrient limitation on phytoplankton – virus interactions are, however, still poorly studied. As climate change will lead to an expansion of stratified and consequently nutrient limited oceanic regions, it is important to understand how phytoplankton mortality rates in the future marine environment will be affected by these changes.

Objectives
Our goal was to study the effects of N- and P-limitation on virus host interaction and the potential relieve of these effects by the uptake of nutrients during viral infection.

Methods
Therefore, cultures of the phytoplankton species Micromonas pusilla and Phaeocystis globosa were grown semicontinuously under limitation of either P, N, or NP until the moment of infection. Upon infection the abundances of host cells and viruses were monitored to obtain one-step growth curves.

Conclusions
Nutrient limitation led to elongated viral latent periods (up to 3-fold) and strongly reduced (up to 8-fold) burst sizes, whereby the outcome was dependent on both the species and the limiting nutrient involved. Furthermore, the negative effects of P-limitation on virus production in the picophytoplankter M. pusilla could be overcome by spiking the infected and P-limited cultures with small pulses of inorganic and organic P, simulating bacterial remineralization and supply by lysis of neighboring cells, respectively.
SAFFOLD VIRUSES IN PEDIATRIC PATIENTS WITH DIARRHEA IN THAILAND

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Background

In recent years, several new viruses associated with diarrheal diseases in human have been reported. Saffold virus (SAFV) is one of a newly identified human Cardiovirus reported for the first time in 2007.

Objectives

This study investigated the prevalence of SAFV and characterized the genotypes distributed in children with diarrhea in Chiang Mai, Thailand during 2012-2013.

Methods

A total of 608 fecal specimens collected during January 2012 to December 2013 from children with diarrhea in Chiang Mai, Thailand were investigated for SAFV by RT-nested PCR and performed sequence analysis.

Conclusions

Nine out of 608 (1.5%) were positive for 4 genotypes of SAFV, SAFV1, SAFV2, SAFV3, and SAFV4. SAFV mono-infection was found in 5 cases, while co-infection with other viruses that cause diarrhea was observed in 4 cases. This study provides more information about SAFV genotypes circulating in pediatric patients with diarrhea in Thailand.
FEMS-2782
Viral infections and host

THE CYTOKINES EXPRESSION IS MODULATED BY EPIGENETIC MECHANISMS IN RESPONSE TO INFECTION WITH IPNV

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Background

The massive salmon farming associated to aquaculture industry has brought a significant increase in infectious diseases, thereby generating economic losses. The infectious pancreatic necrosis virus (IPNV), agent of high mortalities in fish farmed, has capacity to activate the immune response mediated by interferon in their host; the aim of this study is clarify whether the activation of the antiviral response caused by IPNV is through epigenetic mechanisms.

Objectives

We observed that expression of cytokines, in salmonids cells infected with IPNV was associated to promoter methylation status. Moreover, DNA methyltransferase inhibitors treatments were able to modulate the expression of these cytokines.

Methods

Studies of methylation in CpG islands of the promoters of cytokines in infected salmonids cells with IPNV were conducted. Through bisulfite sequencing, change in methylation of the promoter affecting the expression of interferon was evaluated. Furthermore, treatments with DNA methyltransferase (DNMT) inhibitor 5-azacytidine and 5-Aza-2′-deoxycytidine were performed to evaluate the expression of cytokines.

Conclusions

These results show that cytokines expression is regulated by promoter methylation status which could be correlated with the modulation of DNMT activity when cells are infected with IPNV. Our work suggested that epigenetic mechanisms could participate in the regulation of host genes expression in IPNV infection, and provided a new insight into understanding the mechanisms of viral infection.
FEMS-2475
Viral infections and host

EPIDEMIOLOGICAL, CLINICAL MANIFESTATION AND GENOTYPE CHARACTERIZATION OF HEPATITIS-C VIRUS AMONG THE HIV PATIENTS IN KASHAN
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Background
Due to shared transmission routes, HCV infection is highly prevalent among people infected with human immunodeficiency virus (HIV). Little is known about the characteristics of HCV genotypes in among HIV/HCV coinfected patients in Kashan

Objectives
In order to provide valuable information for HCV management in this particular population, we investigated the HCV genotypes in HIV-infected In Kashan-Iran.

Methods
This study was conducted in 2013 in kashan, Iran. The study population consisted of all HIV infected having records in behavioural concelling center and prison in Kashan . Demographic information and HCV, HIV-related risk behaviours were obtained through an interviewer–assisted questionnaire. After taking consent form, 10 cc blood taken from HIV patients and serum samples were screened for HCV infection using enzyme-linked immunosorbent assay (ELISA), and detection of HCV RNA was performed by PCR amplification. HCV subtypes were determined by direct sequencing of amplicons

Conclusions
54(85%) out of the 63 HIV infected patients , were HCV positive which all of them were male..there was a significant correlation between HCV frequency and occupation (P<0.0001) and level of education (p<0/05). 100% of HIV/HCV coinfected cases had history of illicit drug use,92.6% history of prison and and 40.7% had high risk sexual behavior.There was found genotype 1 in 75.9% and genotype 3 in24.1% of HCV patients .94.4% of HCV patients had subtype a.The mean of AST,ALT,TLC,CD4 in HCV patients were 50.6,58.5,397.2,398.8. There was no clinical symptom of chronic Hepatitis c.
The majority of HIV infected persons in Kashan city are HCV positive and predominant genotype is 1 and predominant subtype is a.
PREVALENCE OF HEPATITIS C IN INJECTING DRUG USERS IN KASHAN -IRAN

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Background
Drug injection is a very important risk factor for viral hepatitis and human immunodeficiency virus (HIV) infection

Objectives
The present study was performed to evaluate the prevalence of hepatitis C among injection drug users (IDUs) and to identify the related risk factors for these infections in this group.

Methods
This descriptive-analytical study was conducted in 2011 in Kashan, Iran. The study population consisted of 300 IDUs in MMT, DIC and counselling centers. Demographic information and HCV related risk behaviors were obtained through an interviewer-assisted questionnaire. IDUs serum samples were screened HCV infection using enzyme-linked immunosorbent assay (ELISA). Data analyzed using SPSS.

Conclusions
Of the 300 IDUs, 288(96%) were male. The majority of IDUs 127(42.3%) were in 30-39 age group with mean age 34.9±9.7. The majority of IDUs 224(74.7%) had more than 10 years history of addiction. The most common age of onset addiction was 15-20 year 134(44.7%). The prevalence of HCV was 142(47.3%). It was found that there was a significant correlation between using shared syringe, age and times of prison and HCV infection.

There was high prevalence of HCV among IDU. High risk behaviors such as tattooing, unsafe sex, needle sharing are common so regular screening of IDU, education of personal Health about using sterile syringe, HBV vaccination and treatment of addiction and HCV infection is recommended.
Background
TTV has been linked to non A-G hepatitis

Objectives
The aim of this project was to investigate the rate of infection and to determine the genotypic features of TTV in Qatar

Methods
A total of 644 blood samples representing different nationalities, non-Qatari (526) and Qatari (118) nationals (mostly from Arab and South East Asia countries) were studied for the presence of TTV DNA by nested PCR. The majority (573) of the blood samples belonged to healthy blood donors, while 53 and 54 of the blood samples belonged to hepatitis C virus (HCV) and hepatitis B virus (HBV)-infected patients, respectively.

Conclusions
Results: we showed that the infection rates of TTV in blood donors, and those HCV or HBV-infected patients were 81.4%, 84.9% and 90.75%, respectively. Significant association found between TTV viremia and nationality, or age was detected.
Sequence analysis of PCR fragments amplified from the 5'-untranslated region (5'-UTR) of all (531) TTV positive samples showed that 65.5% (348/531) of the PCR fragment sequences were classified into main genotype 3, followed by genotype 5 (24%), 2 (5.8%) and 1 (4.7%). Interestingly, genotype 4 was not detected among the studied subjects. Phylogenetic and pairwise analyses using sequences from TTV viremic samples also showed an overall close similarity to the main genotype 3. In conclusion, there was no significant difference in the detection rates of TTV among Qatari nationals and non-nationals, and most TTV genotypes, particularly genotype 3, were detected.
PARTIAL CHARACTERIZATION OF A PROBABLE NOVEL SUBTYPE OF BETAPAPILLOMAVIRUS HPV 17

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Background
Human papillomavirus (HPV) have epithelial and mucosal tropism. HPV type 17 belongs to cluster b2 of Betapapillomavirus. Distinct subtypes with dissimilarities of 2 to 10% at the nucleotide level from their reference HPV can be identified. Molecular cloning experiments enabled to recognize two subtypes (17a and 17b) isolated from epidermodysplasia verruciformis (EV)

Objectives
The aim of this study was investigate a possible novel subtype of Betapapillomavirus HPV type 17.

Methods
The isolate was collected from an oral mucosa of an asymptomatic woman. Partial L1 region was amplified using My09/11 degenerated primers. The complete E6, E7, and LCR genes were amplified with specific primers. The Ethics Committee of the College of Medicine at the University approved the protocols for collection and informed consent.

Conclusions
The viral DNA was sequenced and partially characterized. Within these sets, the DNA sequence was altered at 38 positions (15 in L1, 13 in E6, 8 in E7, and 2 in LCR gene). Partial L1 analysis showed high dissimilarity facing to prototype, reaching 5% of nucleotide substitutions. The E6 oncoprotein presented the highest modification among the sequences studied. The amino acid modified at position 62 (S-T) affected one zinc binding domain (CxxC(C)²⁹ CxxC) of this protein. E2 binding factors, transcription factors, and TATA signals were seen in LCR region. Despite the EV-associated HPV 17 has cutaneous tropism, the oral cavity is an appropriated niche to host HPV which naturally infects other sites. Based on these findings, we believe to have found a novel subtype of this type.
FEMS-1090
Viral infections and host

IMMUNOBIO蒂CS MODULATE ANTIVIRAL IMMUNITY AGAINST ROTAVIRUS VIA TOLL-LIKE RECEPTORS PATHWAYS IN PORCINE INTESTINAL EPITHELIAL CELLS
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Background
Rotavirus (RV) is a primary cause of acute diarrhea in infants and young animals, and despite attenuated RV vaccines have been able to reduce the global disease burden associated with RV gastroenteritis, this infectious disease is still a problem in developing countries. Several reports indicate that immunobiotics have the ability to reduce severity of RV mediated diarrhea. Previously, we demonstrated that some immunobiotic strains were able to beneficially modulate Toll-like receptor (TLR)-3 mediated immune response in porcine intestinal epithelial (PIE) cells.

Objectives
To evaluate the capacity of immunobiotics to modulate RV-induced antiviral response in PIE cells.

Methods
(i) PIE and MA104 cells were inoculated with several strains of RVs to investigate the susceptibility and the ability to induce the inflammatory cytokines against RV infection, (ii) The capacity of immunobiotics to beneficially modulate RV-induced immune response in PIE cells was investigated, by evaluating inflammatory cytokines, type I interferons, TLR signaling pathways, and TLR negative regulators.

Conclusions
Challenge of PIE cells with RV reduced cell viability and increased the expression of IFN-β, MCP-1, IL-6, IL-8, and TLR3. Immunobiotics were able to improve cell viability, and to reduce RV-induced inflammatory cytokines expression. Those effects were related to the capacity to modulate TLR negative regulators single such as SIGIRR, Tollip, A20, Bcl-3, IRAK-M, and MKP-1. Our results demonstrated that immunobiotics can be used as immunomodulators to alleviate viral induced diarrhea in livestock animals such as the pig.
FEMS-1505
Viral infections and host

NON-VIABLE IMMUNOBIOТИC LACTОBАСILLUS RHАМNOSUS CRL1505 IMPROVES RESPIRATORY IMMUNITY AND REDUCES INFLUЕNZA VIRUS-ASSOCIATED LUNG INJURY
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Background
Symptoms of influenza infection are variable ranging from mild respiratory distress to massive organ failure resulting in death. Seasonal influenza is usually self-limiting but in susceptible patients it may progress to acute lung injury, which is characterized by augmented pulmonary microvascular permeability, hypoxemia and respiratory failure. The ongoing threat of influenza epidemics and pandemics has emphasized the importance of developing novel, safe and effective therapeutic or preventive strategies against infections from divergent influenza viruses.

Objectives
To examine the effect of nasal administration of heat-killed Lactobacillus rhamnosus 1505 (Lr1505) on immune response to influenza infection in mice.

Methods
Six-week old BALB/c mice were treated with 10⁸ cells of Lr1505 by the nasal route during two consecutive days. Lr1505-treated and untreated control mice were then nasally challenged with Influenza virus. Lung tissue damage and respiratory and systemic immune responses were studied at several time points after viral challenge. Results demonstrated that non-viable immunobiotic strain protected infected mice by reducing pulmonary injury and lung viral loads through several mechanisms: a) inflammatory cytokines were down-regulated diminishing inflammation (p<0.01), b) IFNs with direct antiviral activity were enhanced (p<0.01), c) regulatory cytokines IL-10 was up-regulated (p<0.05), d) maturation markers and co-stimulatory molecules were up-regulated in lung antigen presenting cells (p<0.01), e) IFN-γ and IL-10 producing CD4⁺ lymphocytes were recruited to the lungs.

Conclusions
Our results strongly suggest that administration of Lr1505 may represent an interesting alternative to modulate respiratory immune response and reduce influenza
virus-associated pulmonary damage.
Background. Freshwater cyanobacterium *Microcystis aeruginosa* frequently forms massive natural blooms where the cell densities are comparable to those in laboratory media (10^7 to 10^8 cells/ml). The representative genome of this species contains abundant and diverse potential anti-viral defense genes. Our culture analyses showed distinct CRISPR arrays in this species. This suggested the *Microcystis* cyanophage may have co-evolved with its phage in a frequency-dependent selection.

Objectives. The aim of this study is to understand the co-evolutionary dynamics of the *Microcystis* cyanophage in natural settings.

Methods. The CRISPR diversity of *Microcystis aeruginosa* was examined in a small eutrophic pond using amplicon analysis of the leader-end of CRISPR fragments with a MiSeq mini-gene sequencer.

Conclusions. During the sampling period, a CRISPR-related genotype (CT19) occurred frequently in the *Microcystis* population. A qPCR targeting CT19 showed the copy numbers were from 4.9×10^2 and 3.2×10^5 copies mL^-1 and accounted for 0.2% to 18% of the *Microcystis* population. We obtained 176,800 leader-end CRISPR fragments using amplicon sequencing. Of these, 116,466 were classified into CT19 consisting of two major variants (28% and 72%, respectively). These major CTs co-existed and their numbers oscillated throughout the sampling periods. A few new variants were found to acquire new spacers or the lack of spacers at the leader-end of the major variants. This indicates *Microcystis* cyanophage co-evolution creates a few new genetic variations and also maintains the genetic diversity in the population.
PRODUCTION IN YEAST OF HUMAN BOCAVIRUS 1-4 VP2 VIRUS-LIKE PARTICLES AND GENERATION OF VP2-SPECIFIC MONOCLONAL ANTIBODIES AS NOVEL TOOLS FOR BOCAVIRUS SEROLOGY

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Background

Human bocavirus 1 (HBoV1), first described in 2005, was considered a causative agent of previously unexplained respiratory tract diseases. Recently, 3 new members of genus Bocavirus, HBoV2-4 were described. HBoV2-4 occurs mainly in the gastrointestinal tract but rarely in the respiratory tract, contrary to HBoV1.

Objectives

Recombinant viral antigens have been proven useful for serologic diagnosis of viral infections. Production of HBoV1-4 antigens in yeast expression system has not yet been reported. In the current study, the capsid proteins VP2 of HBoV1-4 were expressed in yeast S.cerevisiae. Electron microscopy demonstrated that all purified recombinant proteins self assembled into virus-like particles (VLPs) exhibiting the typical icosahedral appearance of parvoviruses with a diameter of approximately 20 nm. HBoV1-4 VP2 VLPs were stable in yeast and were easily purified by caesium chloride gradient ultracentrifugation. Four monoclonal antibodies (MAbs) of IgG1 subtype were generated by immunization of mouse with recombinant HBoV1 VP2 VLPs. Three of them specifically recognized only HBoV1 VP2 protein; one MAb was cross-reactive with HBoV2 and HBoV4 VP2 proteins. Recombinant HBoV1-4 VP2 VLPs and VP2-specific MAbs were employed to develop serological assays to detect virus-specific IgG antibodies in human serum specimens.

Methods


Conclusions
Yeast expression system proved to be simple, efficient and cost-effective, suitable for high-level production of HBoV1-4 VP2 as VLPs, that resemble native virus in regards of morphology and antigenicity.
REMARKABLE DIVERSITY AMONG NOVEL DENSOVIRUSES FROM CRICKETS
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Background
Densoviruses are members of the Parvoviridae family infecting invertebrates. Since 2009, a cricket densovirus, AdDV, caused a severe epidemic in the $600-million cricket industry in North-America.

Objectives
During the last 18 months, we also received samples of diseased crickets from North America, Europe and Japan that were negative for AdDV. Electron microscopy demonstrated that these samples contained densovirus-like particles. Objectives of this study included characterization of these virus(es), cloning, sequencing, and X-ray crystallography.

Methods
A SISPA method was used to detect the pathogens (J Virol 88:12152). Sequencing of the amplicons identified those with identities to known viruses allowing by primer extension to obtain the complete genomes and cloning in suitable vectors to obtain infectious clones. Sequencing revealed that some were novel circoviruses and a novel ambisense densovirus (Genome Announc. 1(2):e00079-13, 1(3):e00328 and 131(6):e00914-13). However, the greatest surprise was a densovirus with a segmented genome (AdSDV). NS- and VP- coding sequences are in separate segments of about 3.3 kb. Both NS- and VP- segments are flanked by identical hairpin terminal sequences but lack ITRs. The two ORFs for VP arose through recombination between a mosquito brevidensovirus and AdDV. The downstream ORF with AdDV-VP1up-like phospholipase A2 could be spliced to the C-terminus of the major capsid protein. Currently, the capsid structure (X-ray crystallography) and transcription strategy of AdSDNV.

Conclusions
A brevidensovirus that normally infects only mosquitoes has adapted itself through recombination with AdDV to crickets. These findings necessitate a revision of the definition and taxonomy of parvoviruses.
Viral infections and host

TIME SINCE ONSET OF DISEASE AND INDIVIDUAL CLINICAL MARKERS ASSOCIATE WITH TRANSCRIPTIONAL CHANGES IN UNCOMPLICATED DENGUE

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Background
Dengue virus (DENV) infection causes viral haemorrhagic fever that is characterized by extensive activation of the immune system.

Objectives
The aim of this study is to investigate the kinetics of the transcriptome signature changes during the course of disease and the association of genes in these signatures with clinical parameters.

Methods
Sequential whole blood samples from DENV infected patients in Jakarta were profiled using affymetrix microarrays, which were analysed using principal component analysis, limma, gene set analysis, and weighted gene co-expression network analysis.

Conclusions
Time since onset of disease associates with the shift in transcriptome signatures from immunity and inflammation to cell cycle and repair mechanisms in patients with non-severe dengue. The strong association of time with blood transcriptome changes hampers both the discovery as well as the potential application of biomarkers in dengue. Clinical diagnosis (according to the WHO classification) does not associate with differential gene expression. However, network analysis did show that that key clinical markers, including platelet count, fibrinogen, albumin, IV fluid distributed per day and liver enzymes SGOT and SGPT, strongly correlate with gene modules that are enriched for genes involved in the immune response. The expression level of these gene modules may support earlier detection of disease progression as well as clinical management of dengue.

COMPLEXITY OF THE VIROME FROM THE KY ISOLATE OF PHOMOPSIS LONGICOLLA FUNGUS

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Background

\textit{Phomopsis longicolla} (teleomorph = \textit{Diaporthe}) is an ascomycete phytopathogenic fungus. Mycoviruses were reported to debilitate their hosts \cite{1}. The KY isolate and derived subcultures of \textit{P. longicolla} exhibited abnormal growth and reduced virulence. Initial examination revealed that the isolate harbors \textit{Phomopsis longicolla} hypovirus 1 (PIHV1) \cite{2}. This ssRNA virus belongs to a group of betahypoviruses that, unlike alphahypoviruses, were not reported to reduce virulence of their hosts.

Objectives

The objective of the study was to investigate potential viral (or other) determinants of hypovirulence and characterise structure and diversity of PIHV1 population in the KY isolate of \textit{P. longicolla}.

Methods

Illumina-based DNA and RNA deep sequencing was performed of a number of the KY subcultures and the type strain of \textit{Phomopsis longicolla} Hobbs, anamorph (ATCC\textsuperscript{®} 60325\textsuperscript{™}).

Conclusions

The analyses of the NGS data revealed that the virome of the KY isolate consists of PIHV1 recombinant with a previously undescribed hypovirus, population of deletion mutants of PIHV1, and a novel RNA virus related to members of \textit{Narnaviridae} family.
CHARACTERIZING THE VIROME OF THE ENTOMOPATHOGENIC FUNGUS BEAUVERIA BASSIANA

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Background

Entomopathogenic fungi are of scientific interest because they enable analysis of virus-host interactions and can be used as biocontrol agents of insect pests. The deuteromycetous fungus Beauveria bassiana has a widespread geographical distribution and a wide host range. Mycoviruses have been described mostly in phytopathogenic fungi and are occasionally associated with hypovirulence, while in entomopathogenic fungi presence of mycoviruses has been reported only rarely.

Objectives

The aim of the present study is to detect and analyse the viruses present in a collection of B. bassiana isolates sourced from worldwide locations.

Methods

Population studies, RT-PCR, cloning and sequencing, hybridization experiments, time-course studies and transmission electron microscopy (TEM) were used to characterize the virome of B. bassiana.

Conclusions

A population study revealed that 17/75 (22.7%) B. bassiana isolates harbor dsRNA elements. Two members of the Partitiviridae family, B. bassiana partitivirus-1 and -2 (BbPV-1 and BbPV-2), have been sequenced, while TEM revealed the presence of virus-like particles. Hybridization experiments revealed that BbPV-1 and BbPV-2 are present in seven B. bassiana isolates derived from different hosts and geographical origins. Additionally, four members of the Totiviridae family, originating from the Iberian Peninsula and the Canary Islands, have been partially sequenced. Furthermore, two previously uncharacterized viruses, B. bassiana polymycovirus-1 (BbPmV-1) and B. bassiana non-segmented virus-1 (BbNV-1), have been detected and fully sequenced and the prevalence of further PmV-like viruses in B. bassiana is currently under investigation. Finally, a time-course study revealed a strong negative
correlation between the copy numbers of viral dsRNA and the developmental stages of the fungus.
THE NEW MESOPHILIC FILAMENTOUS ANOXYGENIC PHOTOTROPHIC BACTERIUM FROM SULFIDE HOT SPRINGS.
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Background

Filamentous anoxygenic phototrophic bacteria (FAPB) belong to the deep-branching phylum Chloroflexi, and are considered to be the most ancient lineage of phototrophs. The unambiguous reconstruction of the evolution of FAPB is complicated by poor representativeness of mesophilic FAPB.

Objectives

The above mentioned pointed our efforts to the search and study of new mesophilic FAPB in mesothermal cyanobacterial mats from hot springs in Barguzin valley (Buryat Republic, Russia).

Methods

Mat samples were collected from mesothermal zones of Umkhey (36 – 40°C) and Kuchiger (35 - 39°C) sulfide hot springs. The enrichment culture containing mesophilic green filamentous anaerobic bacterium Um-2 was obtained from Umkhey mat sample at the Pfenning medium (pH 7.0) at 28°C under illumination. The Um-2 cells (diameter about 0.8 µm) were combined to trichome. The main cellular pigments was bacteriochlorophyll c (maximum absorption at 740 nm). Gas vacuoles were not found. According to the phylogenetic test results (by 16S), Um-2 formed the branch belonging to mesophilic FAPB cluster. The nearest neighbor of Um-2 was “Candidatus Chloroploca asiatica” Um-3 (KJ605349) (91% of similarity). The Um-2 phylotype representatives was detected in Umkhey and Kuchiger mat samples using PCR test with FAPB-specific pufLM primers. The sequences of rrna (16S rRNA coding) and pufLM genes were deposited in GenBank under accessions KP341999 and KP342000 respectively.

Conclusions

Thus the enrichment culture of new mesophilic FAPB belonging to the new genus and species of the Chloroflexales order was obtained. It was shown that the new bacterium is presented in the microbial community of sulfide springs of Barguzin valley.
FEMS-0578
Free subjects

ANTIFUNGAL ACTIVITY OF NONHYDROLYTIC SYNTHESIZED TiO2

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Background
Despite of many studies on antibacterial activity of titanium dioxide (TiO₂), its antifungal properties are not yet well evaluated.

Objectives
The aim of our study is to exam the activity of nanosized TiO₂ against yeast cells.

Methods

TiO₂ used was synthesized by sol-gel method and characterized as anatase with particles size about 20 nm. Its antifungal activity was examined against C. albicans, C. tropicalis, C. lusitaniae, C. glabrata and C. krusei, using the following experimental setup: concentration of TiO₂ powder – 1 mg/ml, 100 ml fungal suspension with cell density about 10⁴ CFU/ml and BLB lamp (50 Hz 8W T5 with emission 365 nm) situated at distance 10 cm. At regular intervals of time, samples were taken and serial dilutions were prepared. The number of viable cells was detected by spread plate method. For determination of the post-irradiation effect, the treated suspensions were tested after 48 h dark period.

Conclusions
The data obtained about antifungal activity of TiO₂ are in agreement with photocatalytic properties of metal oxide nanoparticles. After 120 min UV-A photocatalytic treatment with 1 mg/ml TiO₂, the number of colonies of C. albicans and C. tropicalis decrease from about 10⁴ CFU/ml to less than 5 CFU/ml with 100% reduction index. Final reduction index in C. lusitaniae and C. glabrata is 99.643% and 99.898%, respectively. These data indicate strong photocatalytic fungicidal effect of TiO₂ on the tested fungi. At the same conditions, the colonies of C. krusei are reduced with 98%. Our data show that the nonhydrolytic synthesized nanosized TiO₂ possesses antifungal activity against isolates of the genus Candida with some differences in the speed of photocatalytic process.
CIPROFLOXACIN EFFECT ON SERUM OXALIC ACID LEVEL

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Background
Ciprofloxacin ranks among the primary drugs used as treatment in urinary system infections in adults.
It is known that Oxalobacter formigenes colonization in intestines is disrupted by antibiotic intake.
Due to the Oxalobacter formigenes colonization deficiency, the fact that high level serum oxalic acid results in formation of renal calcium oxalate stones has been determined.
Additionally, serum oxalic acid level increase is thought to have negative effect on urinary and cardiovascular system.

Objectives

The purpose of our study is to search whether there is an increase in serum oxalic acid level because of intestinal flora disorder in the patients with urinary system infection and the individuals given ciprofloxacin as treatment.

Methods

In our study, 40 people who have urinary system infection and take ciprofloxacin treatment have been included. Patients serum oxalic acid levels were evaluated both in pre-treatment and post-treatment evaluation which was a month later. Serum oxalic acid was measured by a commercially available kit (SunRed Biological Technology, Serial number: 201-12-2153), using Human Oxalic Acid ELISA method.

Conclusions

In conclusion, when pre-treatment and post-treatment serum oxalic acid levels were compared, no statistical difference was found (p=0.47). It was found that ciprofloxacin treatment didn’t affect serum oxalic acid levels of the patients with urinary system infection.

In the light
of this data,
it is concluded that these patients don’t need to be followed up for any possible complications
because of high level serum oxalic acid.
Background

Anaerobic bacterium Clostridium difficile has been traditionally considered as a human nosocomial enteropathogen. Although its natural habitat is human and animal intestine, the ability to form dormant endospores enables C. difficile to survive also in aerobic environment. Recent emergence of community acquired C. difficile infections and infections in animals implies that environment might be a reservoir of C. difficile and a source of infection.

Objectives

Our aim was to investigate diversity and overlap of C. difficile PCR-ribotypes isolated from animals and different non-hospital environments in Slovenia.

Methods

Strains included in the comparison originated from domestic animals (calves, piglets, horses, poultry and birds, goats, sheep, dogs and cats) and different environments (puddles and soil from rural and urban areas, compost and water from rivers and ponds) in Slovenia. Isolates were characterized by PCR-ribotyping and toxinotyping.

Conclusions

This work is unique in presenting a large number of C. difficile strains from several rural and urban environments and compare them to animal and human isolates. Altogether 80 PCR-ribotypes (47 in animals and 58 in the environment) were identified and 25 were shared between animals and the environment. Forty-six of
these PCR-ribotypes were found also in humans in Slovenia. Twenty-seven PCR-ribotypes were identified as non-toxigenic with more than half found only in the environment, mainly puddles and soil, which seem to represent an independent environment. Our results show that C. difficile genotypes that are commonly isolated from animals can also be isolated from different environments, indicating that non-hospital environment could represent an important reservoir.
Background

When a high-voltage direct-current is applied to two beakers filled with water, a horizontal electrohydrodynamic (EHD) bridge forms between the two beakers. In this work we studied the transport and behavior of yeast cells and ThP1 human-monocytic-cell-line in this special environment.

Objectives

The behavior of *Escherichia coli* in an EHD-bridge set-up was recently investigated: *E.coli* cells survive the procedure; the main direction of cell transport is from the cathode to the anode beaker. The goal of this work was to test in how far these results are specific for *E.coli*, thus studies with yeast cells and ThP1 cells were conducted.

Methods

Cells were added to one or to both beakers, and the transport of the cells through the bridge was investigated using optical and microbiological techniques. One experimental series consisted of three different configurations: carrier solution in both beakers, only in the anode or only in the cathode beaker. As carrier solutions for yeast 5% glycerol and ThP1 cells 4.5% glucose were used.

Conclusions

The behavior of cells in an EHD-bridge depends largely on their size and their surface charge. The heavier the organism, the slower the transport; the higher the surface charge, the more pronounced the preferred flow direction.

*S.cerevisiae*: transport was observed in both directions. The absence of a preferred flow direction means that electric forces play a smaller role which is corroborated by the zeta potential.
ThP 1 cells survive the transport and showed similar behavior as the *E.coli* cells: pronounced transport towards the anode.
POST-ANTIBIOTIC AND POST-ANTIBIOTIC SUB-MINIMAL INHIBITORY CONCENTRATION EFFECTS OF CHLORHEXIDINE AGAINST ORAL BACTERIA

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Background
The post-antibiotic effect (PAE) is defined as the suppression of bacterial growth for a particular duration after a brief exposure to an antimicrobial agent. Chlorhexidine is a widely used biocide found in antiseptic products; however, its PAE remains unclear.

Objectives
In this study, PAE, post-antibiotic sub-minimum inhibitory concentration (MIC) effects (PA-SME), and sub-MIC effects (SME) of chlorhexidine on oral bacteria were investigated.

Methods
For PAE measurement, bacteria were exposed to 10x MIC chlorhexidine for 1 min, which was then eliminated by washing. For determining PA SME, bacteria were exposed to 0.1, 0.2, and 0.3x MIC chlorhexidine during the postantibiotic phase and to sub MIC chlorhexidine for measurement of SME. PAE, PA SME, and SME of chlorhexidine were observed.

Conclusions
The PAE lasted for 0.9 h for Streptococcus mutans, 0.1 h for Streptococcus gordonii, and 0.35 h for Lactobacillus acidophilus. The PA SME against oral bacteria lasted for longer duration as the chlorhexidine concentrations increased. The PA SME against oral bacteria lasted for substantially longer than SME did. The present study illustrates the existence of chlorhexidine-induced PAE, PA SME, and SME against oral bacteria, thereby extending the pharmacodynamic advantages of chlorhexidine.
Background

According to the demographic ageing of labor force participation in Taiwan, we imported massive foreign laborers to Taiwan for nursing or other labor force work since October, 1989. The purposes of this study was to evaluate the effectiveness of single examination to diagnose Enterobiasis by scotch tape perianal swab technique in non-constant time.

Objectives

The survey covered southeast asian laborers from four countries including philippines, thailand, Indonesia and vietnam. The data were collected from 391 valid questionnaires, 391 valid stool samples and 391 valid scotch tape perianal swab samples.

Methods

Single scotch tape perianal swab technique was performed by foreign laborers in non-constant time. Stool samples were also obtained after the scotch tape perianal swab. This was a cross-sectional study. Descriptive statistics were applied to the data.

Conclusions

Scotch tape perianal swab technique in non-constant time could increase the detection rate of Enterobius vermicularis from 0% to 2.0%(8/391). Occupational health personnel should educate the foreign laborers to get into the habit of washing hands before a meal in their daily practice. The employers should arrange the examination of scotch-tape anal swab technique for caretakers periodically.
SPECIFIC KILLING OF BACTERIA USING A TOXIN-INTEIN BASED AND-LOGIC GATE

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Background

Synthetic biology aims to reprogram bacteria to design and construct sophisticated new biological systems by gene regulatory networks. However, gene regulatory networks (gene expression, protein or molecule concentration) are continuous, thus fitting them into a deterministic model like Boolean approximation (0/1) is noisy.

Objectives

We would like to implement a model aiming at killing specific bacteria in mixed populations. We will design an AND-logic gate to solve Boolean satisfiability (SAT) problem. Our model is based on conjugation where plasmids will act as wires (plasmid inside cell is 1; no plasmid is 0).

Methods

To avoid the noise associated with transcriptional regulators, our system is based on half toxins (thus non-functional) coupled with split intein technology. We are designing two types of plasmids: input plasmids (A and B) and clause plasmid (CP). Only CP is conjugative. A population of bacteria could have already implemented the input plasmids in different combinations: A and B; only A; only B or no one. The output in our system is cell death. The CP will infect these bacteria by conjugation and it will evaluate the inputs and will specifically kill bacteria harboring both plasmids (A and B). When both plasmids are together in the same cell, the expression of the toxin-intein fusion triggers the reaction between two inteins domains, allowing the reconstitution of the toxin and killing the cell.

Conclusions

We have successfully completed the evaluation of Boolean-SAT problem by the construction of an AND-logic gate and using the inputs plasmids A and B, and the CP.
OUTBREAK OF BOTULISM IN MEMBERS OF A FAMILY FOLLOWING THE CONSUMPTION OF KUSHK

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Background
Botulism is a serious neuroparalitic disease caused by the Clostridium botulinum

Objectives
In this article, we report a outbreak of the foodborne botulism due to consumption of kushk in 5 members of a family

Methods
A 41 and 46 year old women presented with dylopia and blured vision from 4 days ago. She had progressive symptoms including dry mouth, dysphagia and swallowing disorder, constipation, dyspnea, general weakness of neck muscle and lower and upper limbs. Pupils were dilated and noreactive to light. There was no gag reflex. Deep tendon reflexes were normal and the muscle force of upper limb was 4/5.

The third case was a 13 year old boy presented some days after two above mentioned cases. His complains were petosis, dylopia, dry mouth, vertigo and weakness of lower and upper limbs but had no constipation, dysphagia and swallowing disorder and dyspnea. The forth case was a 12 year old boy with history of diabet and similar symptoms and sign and the fifth case was a 8 year old girl with neck rigidity and petosis.

They were members of a family and had history consumption of kushk. Interval between involvement of children was longer than adult and incubation period. Disease was more severe in adult than children. EMG was done in one of the woman and was suggestive of butolism. After clinical diagnosis, serum of patients was obtained and the patients received antitoxin. Their symptoms resolved gradually.

Conclusions
The diagnosis of botulism should be made based on the patient's initial sign and symptoms and history of consumption of suspected food...
BLOOD BORNE DISEASES IN INTRAVENOUS DRUG ABUSERS

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Background
Drug injection is responsible for numerous minor to life-threatening and fatal complications such as the transmission of human immunodeficiency virus (HIV), sexually transmitted diseases, and viral hepatitis

Objectives
This paper has tried to offer a brief look at the epidemiology of drug abuse, while focusing on blood born infections seen in IDUs.

Methods
The study was carried out using a cross sectional design. A questionnaire was distributed among 135 IDUs. Serological tests (ELISA Anti HCV, Anti HIV, HBs Ag) were taken to confirm the existence of blood borne diseases.

Conclusions
A total of 135 cases between the ages of 15-57 were studied. Opium was most frequently first drug used (58.5%), followed by cannabis (20.7%) and heroin (11.1%).

Nearly 68.9% of the population began abusing drugs before the age of 20. 61.4% of the cases had a history of sharing needles. Other results showed that 51.5% had an accompanying blood borne disease. Further analysis indicated a meaningful relationship (P< 0.001) between needle sharing and accompanying blood borne diseases. Of the 51.5% who had an accompanying blood borne disease, 17.1% were found to have either reactive ELISA of AntiHIV alone, or in combination with Hepatitis. In other words, it can be concluded that 8.1% of the total population had reactive ELISA of AntiHIV. Results also showed that 11.1% of the total population had HBs Ag positive while 47.4% had Positive ELISA Anti HCV.
MICROBIAL CAUSES OF WOUNDS AND SKIN INFECTIONS IN INJECTING DRUG USERS

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Background
60–80% of hospital admissions in IV drug abusers are due to infectious complications and skin and soft-tissue infections seem to be some of the most common infections among this group.

Objectives
The goal of this study was to identify the characteristics of wounds and skin Infections of this group to hopefully have early recognition and as a result better prevention and management strategies.

Methods
This descriptive cross sectional study was carried out on 135 IDUs. All current injection drug users who had at least one wound as a result of injecting drugs were enrolled in this study. The wounds were physically examined and smear samples were also cultivated on Blood Agar and EMB Agar environments for microbiological assessments.

Conclusions
A total of 135 cases between the ages of 15-57 were studied. The most wounds (45.2%) were located in the lower limbs such as the foot. The majority (44%) of wounds were as old as six months up to one year. Each individual had at least 4 wounds with an average size of 1.8 cm². 22.2% of the population had some form of infection such as abscess or cellulites. Thrombophlebitis was seen in 37% and. Only 10.4% showed necrotizing ulcers.

Microbiological assessments found that the majority of wounds (54.1%), had gram positive cocci. This included staphylococcus (36.3%), streptococcus (17.8%). 7.4% of the cases showed gram negative species. 30.4% of the bacterial cultures showed negative or nontypable results, since the culture environments used were very routine and non-specific. Fungi were seen in 8.1% of the cases.
MICROBIAL CAUSES OF WOUNDS AND SKIN INFECTIONS IN INJECTING DRUG USERS

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²Infectious disease, Kashan University of Medical Sciences, Kashan, Iran

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OUTCOME OF HIV/AIDS PATIENTS WITH PNEUMONIA

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Background
Pneumonia is also a major cause of prolonged hospitalization, mortality and morbidity among HIV infected people that costs millions each year.

Objectives
The goal of this study was determine of outcome and prognostic factors of pneumonia in HIV positive individuals.

Methods
In a descriptive method, 80 HIV positive patients that hospitalized in Loghman-hakim hospital between 2005 to 2011 with pneumonia have been studied and their demographic findings include age, sex, marriage, job, social history such as smoking, I.v drug using and imprisonment, medical history of co infections such as pulmonary TB, HCV and HBV, and also clinical features include their initial signs and symptoms, lab data and imaging, and at last their outcome gathered and record in questionnaire. Then gathered data enter SPSS#13 software, analyzed by descriptive tests.

Conclusions
80 HIV positive patients hospitalized with pneumonia in loghman-hakim hospital since 2005 to 2011 entered the study. Mean age was 37.4(±8.2) years. Death was the first with 43.8%, then recovery with 30% and pulmonary complications with 16.3% were the most common outcomes. Mortality was more common in the elderly. Most of the patients had history of smoking, injection drug use and imprisonment history and mortality rate in these patients were more common. The mortality rate was also more common in patients with anemia, high ESR levels, CRP +, high urea and high creatinine levels and low CD4 counts and pleural effusion.
SCREENING OF SIDEROPHORE BIOSYNTHESIS GENES FROM SIDEROPHORE PRODUCING ACTINOMYCETES

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Background

Iron is an essential trace element required for growth of all microorganisms. However, iron in environment is mostly insoluble form and cannot be readily utilized by microorganisms. In iron starvation condition, actinomycetes have developed specific mechanisms involving in the production of small organic metal chelator called siderophore.

Objectives

Actinomycetes were isolated from coastal and mangrove sediments in Thailand. Genomic DNA were extracted by phenol-chloroform method. Hydroxamates and catecholate siderophore biosynthetic genes were detected by PCR using desD and entF specific primers, respectively. Siderophore production was determined by CAS assay. Siderophore producing actinomycetes were identified by 16S rRNA gene sequence analysis.

Methods

A 1.2 kb fragment of desD gene that encodes for desferrioxamine, a hydroxamate siderophore was detected in 7 isolates. A 203 bp amplified fragment of entF gene encoding enterobactin, a catecholate siderophore was detected in 3 isolates. All isolates produced siderophore on Chrome Azurol S agar. High siderophore producing Streptomyces parvulus S2-SC3 was selected for sequence analysis of desD gene. A 1.2 kb amplified PCR fragment showed 90% sequence similarity with desferrioxamine synthetase from Streptomyces lividans strain 1326. Phylogenetic analysis revealed that these siderophore producing actinomycetes were members of the genus Micromonospora, Streptomyces and Verrucosispora.

Conclusions

PCR offers a fast way to screen for siderophore producing actinomycetes. However, some producing strains were failed to detect by our PCR screening, several actinomycetes strains may carry diverse siderophore biosynthesis gene which could not be detected by currently available primers. Our results suggested that novel siderophore may exist in these actinomycetes.
TAXONOMIC DIVERSITY OF ACTINOMYCETES FROM THAI MANGROVE SEDIMENTS

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Background

Actinomycetes are large group of high GC Gram positive bacteria that widely distributed in nature. They have proved to be an important source for industrially useful bioactive metabolites, notably antibiotics. In Thailand, there are very few studies on microbial diversity in mangrove forests. In this study, we report on the diversity of cultivable actinomycetes in mangrove sediments and provide taxonomic evidence of their novelty taxa.

Objectives

Actinomycetes were isolated from mangrove sediments and taxonomically characterized. Representative isolates were selected and identified based on 16S rRNA gene sequence analysis. Putative novel taxa were subjected to polyphasic taxonomic characterization using standard methods.

Methods

Over 100 strains of actinomycetes were isolated from mangrove sediments using media selective for actinomycetes. Phylogenetic analyses based on a 16S rRNA gene sequences showed that the majority of isolates were dominated by genus Streptomyces. In addition, members of the genera Amycolatopsis, Jiangella, Micromonospora, Nocardia and Verrucosispora were also found. Some isolates may represent a new species. For example, Jiangella strain 3SM4–07T was most closely related to Jiangella alkaliphila JCM 15620T (99.0%), However, they shared mean DNA–DNA hybridization values of only 31.5±2.6%.

Conclusions

Our results provide further evidence that mangrove sediments harbor taxonomically diverse actinomycetes. The diversity of these organisms and their novelty support the view that this under studied habitat is a rich source of novel actinomycetes which can be further exploited for biotechnological purposes.
ACTINOMYCETES FROM EUCALYPTUS AND THEIR ACTIVITY TO INHIBIT EUCALYPTUS LEAF AND SHOOT BLIGHT DISEASE

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Background

Eucalyptus are the most important raw material supply for pulp industries in Thailand. Leaf and shoot blight caused by Cryptosporiopsis eucalypti, Cylindrocladium sp. and Teratosphaeria destructans is a serious disease in eucalypt plantations. The search of actinomycetes for biological control agents is one of the promising approaches.

Objectives

In this study, we isolated actinomycetes capable of inhibiting the growth of fungal pathogens causing leaf and shoot blight on eucalyptus. The potential strain to use as biocontrol agent was selected for further study.

Methods

Actinomycetes were isolated from roots and rhizospheric soil of eucalyptus using starch casein agar and humic acid-vitamin agar supplemented with antimicrobial antibiotics. The isolates were screened for their ability to inhibit eucalyptus fungal pathogens using dual culture technique.

Conclusions

A total of 478 actinomycete strains were successfully isolated from roots (95 isolates) and rhizosphere soil (383 isolates) of eucalyptus. Among these, 439 isolates (91.8%) were streptomycetes and 39 isolates (8.16%) were non-streptomycetes. All isolates were evaluated for their in vitro antagonistic effect on plant pathogenic fungi. The result showed 273, 120 and 24 isolates were antagonistic to Cryptosporiopsis eucalypti, Cylindrocladium sp. and Teratosphaeria destructans, respectively. Among these, 48 isolates (10%) displayed activity against all three fungal pathogens. The isolate EUSKR2S82 showed the strongest inhibitory effects against all tested fungi.
The identification of this strain using 16S rRNA genes sequence revealed that isolate EUSKR2S82 shared 99.4% similarity with *Streptomyces ramulosus* NRRL B 27-14.
CULTIVABLE ACTINOMYCETES ASSOCIATED WITH GLOMUS MOSSEAE SPORES AND THEIR PLANT GROWTH PROMOTING ACTIVITY

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Background

Actinomycetes are prolific producers of bioactive compounds. Much of our research effort has been directed at assessing the diversity of actinomycetes in various environments. Currently, we are interested in mycorrhizal helper bacteria which may be a good source for novel taxa for bioprospecting.

Objectives

In this study, we report on the isolation of actinomycetes from spores of Glomus mosseae and provide preliminary evidence of their potential in agriculture as plant growth promoter.

Methods

Actinomycetes were isolated from spores of Glomus mosseae using the dilution plate technique and media designed for the selective isolation of members of specific actinomycete taxa. All isolates were identified by 16S rRNA gene analyses. The isolates were examined for their ability to produce indole-3-acetic acid, siderophores and solubilize phosphate in vitro.

Conclusions

Six strains were isolated from spores of Glomus mosseae using media selective for actinomycetes. Phylogenetic analyses based on a 16S rRNA gene sequences showed that the isolates belonged to the genera Lysibacter and Streptomyces. All isolate produced siderophore, four isolates produced IAA and two isolates solubilized phosphate at varying level. S. thermocarboxydus S3 produced 11.23±0.02 mg/ml IAA and high activity of phosphate solubilization and siderophore production. The results provide evidence that actinomycetes were associated with arbuscular mycorrhizal spores of Glomus mosseae. The ability of these organisms to produce plant growth promoting agents support the possibility of using these actinomycetes for agricultural purposes.
PROFERROROSAMINE A SECRETION BY ERWINIA RHAPONTICI P45 INHIBITS GROWTH OF THE FIRE BLIGHT PATHOGEN ERWINIA AMYLOVORA

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Background
The classic diagnosis of the fire blight pathogen E. amylovora (EA) takes the cell morphology on King’s B agar into account. Sometimes, an unusual pink coloration of EA colonies can be observed on this agar. This coloration is due to the co-cultivation of E. rhapontici or E. persicina and their production of the iron (Fe²⁺) chelator proferrorosamine A (pFRA), that accumulates in E. amylovora. Dependent on the proximity of a pFRA synthesizer, EA colonies can also exhibit an attenuated growth.

Objectives
To get a deeper insight into the mechanism of the Fe²⁺ chelator, we aimed to identify the genes involved in proferrorosamine A synthesis.

Methods
pFRA negative mutants of E. rhapontici P45 were isolated in a phenotypic transposon mutagenesis screen.

Conclusions
We identified a 9.3 kb cluster of seven genes that encode proteins involved in pFRA synthesis. Wild type E. rhapontici P45, but not pFRA negative mutants, showed a growth inhibition effect on EA in vitro and detached flower assay.
COMBINED EFFECT OF OSMOTIC AND UV-B IRRADIATION ON A N2 FIXING CYANOBACTERIUM FROM A RICE PADDY

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Background
Cyanobacteria play a crucial role in the productivity of terrestrial and aquatic systems and have developed a series of defense mechanisms against abiotic factors as UV-B light and osmotic stress.

Objectives
In order to study the protective strategies of Calothrix BI22 spp., this cyanobacterial isolate from a subtropical rice field was exposed to low doses of UV-B alone or in combination with osmotic stress.

Methods
After the stress exposure, oxidative damage was determined by TBARS and proline accumulation quantified. The activity of enzymes (superoxide dismutase and catalase) involved in the decrease of reactive oxygen species (ROS), \( \text{O}_2^- \) or \( \text{H}_2\text{O}_2 \), were determined and the \textit{in vivo} production of ROS was followed by confocal microscopy. Photosystem II yield and \( \text{O}_2 \) photoevolution were also studied as proxies of the effect of the stress on the cyanobacterium physiology.

Conclusions
Osmotic stress, in general, caused a more pronounced oxidative damage but in combination with UV-B the damage drastically increased. Among the enzymes involved in the antioxidant defense, CAT had a higher activity under osmotic stress but in the presence of both types of stresses its activity doubled. SOD, on the other hand, had higher rates only under UV-B light. The accumulation of ROS followed a similar pattern as the antioxidant enzymes studied. Photosynthesis was affected in all of the situations studied. This study showed that this cyanobacterium had a differential response when exposed alone to each type of stress than in combination.
Background
Agarose gel electrophoresis has been widely used as the standard method to separate, identify, and purify DNA/RNA fragments in the life sciences research, because it’s easy to perform, relatively inexpensive, and excellent in analytical performance characteristics. However, technologies to separate for short DNA fragments in high resolution are still lacking.

Objectives
In response to the above challenge, we hypothesized that the tailoring agarose molecules with various pendant groups would provide the rapid analysis of an array of small to large DNA fragments in the gel electrophoresis.

Methods
We examined this hypothesis by tailoring the agarose molecules with various pendant groups such as octadecyl chains, and aromatic groups, and then directly fabricate the three dimensional agarose gel for electrophoresis. This concept of the hydrophobically-modified 3D microporous gel can be regarded also as a novel microfluidic-based mold for being demonstrated and also used in the bio-microelectromechanical system (Bio-MEMS). The molecular structure (degree of substitution and molecular weight) of the modified agarose was confirmed by 1H-NMR, and MALDI-TOF (Matrix: CHCA) analysis. Interestingly, the tailoring agarose gel provided the distinctive separation of small DNA fragments efficiently and also enhanced the discriminatory power in the gel electrophoresis based on the pendant groups’ characteristics.

Conclusions
Overall, the results from this study may be useful in understanding the role of the hydrophobic/hydrophilic balance in the separation of DNA fragments and further in the extracellular microenvironments in tuning a variety of cellular activities.
TEMPERATURE- AND NITROGEN SOURCE-DEPENDENT GROWTH OF LISTERIA MONOCYTOGENES IMPLICATES AN IMPORTANT ROLE OF 2-OXOGLUTARATE AS INTERNAL SIGNAL DURING ADAPTATION TO CHANGING ENVIRONMENTS

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Background
The pathogen Listeria monocytogenes lives either saprophytically in the environment or within cells in the vertebrate host. Thus it has to adapt its lifestyle to its ecological niche and the availability of nutrients.

Objectives
The aim of this study was to investigate the impact of different nitrogen sources and temperatures on growth of L. monocytogenes.

Methods
Nitrogen source dependent growth analysis was conducted in defined minimal medium at 24°C or 37°C with either glutamine or ammonium as nitrogen source. Nitrogen source- and temperature-dependent gene transcription was validated by qPCR.

Conclusions
A preference for ammonium over glutamine was observed in growth experiments. This might be ascribed to the interrupted TCA cycle in L. monocytogenes, causing accumulation of the intermediate 2-oxoglutarate. 2-oxoglutarate has to be removed to allow a permanent glucose metabolism by continuous channeling of pyruvate in the TCA cycle. During growth on ammonium twice the amount of 2-oxoglutarate is removed compared to growth on glutamine.

Reduced growth on glutamine was more obvious at 24 °C than at 37 °C. When grown on the same nitrogen source, but at different temperatures, the removal of 2-oxoglutarate is mainly determined by the amount of glutamate used for the de novo biosynthesis of amino acids which is expected to be elevated at the optimal growth temperature of 37 °C.

It is assumed that in L. monocytogenes the intracellular 2-oxoglutarate concentration serves as internal signal that allows the adaptation of L. monocytogenes to two environmental parameters, temperature and nitrogen source, at the same time.
INVESTIGATION OF AUTOANTIBODIES IN CANCER PATIENTS

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Background

Malignant diseases are often related with the induction of autoimmunity, characterized by the generation of autoantibodies against a wide range of autoantigens. Serum autoantibodies have been identified in patients with solid tumors and in patients with hematological malignancies.

Objectives

The purpose of the study was to investigate the prevalence of antinuclear antibodies (ANA) in sera of the patients with various cancer types.

Methods

Serum samples were obtained from 54 patients with cancer (37 breast, 1 myxoid liposarcoma, 1 brain, 2 lung, 1 colon, 3 gastric, 1 pancreas, 2 lymphoma, 1 multiple myeloma, 1 nasopharynx, 1 renal, 1 prostate, 1 vulva, 1 endometrial) and with rheumatic compliant. A total of 100 healthy blood donors served as the study control group. ANA patterns were searched by using the HEp-2010/Liver (Monkey) indirect immunofluorescence assay (IFA) kit (Euroimmun AG, Germany). In addition these samples were further processed by line immunoassay (LIA), (Euroimmun AG, Germany).

Conclusions

Of 54 patients, 18 were ANA-positive by IFA and LIA. Homogen (8), granular/speckled (2), nucleolar (2), anticentromere (2) and mixed (4) patterns were detected by IFA. SS-A/Ro (8), SS-B/La (6), antidsDNA (2), antiScl (2) and CENP-B (2) positivity were observed by LIA. In healthy control group, ANA was not detected. There was statistically significant difference between the results of the patients and of the control group. Of 54 patients, 29 were diagnosed with cancer while 20 were diagnosed with rheumatic disease firstly; and 5 were diagnosed with both simultaneously. In
conclusion, the association between rheumatic diseases and malignant diseases need further investigation.
ANTIDFS70 ANTIBODIES IN 500 HEALTHY BLOOD DONORS
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Background
Autoantibodies against DFS70 (dense fine speckled 70) antigen have been recently identified among the antinuclear antibodies (ANAs) in patients with systemic autoimmune rheumatic diseases.

Objectives
The typical DFS has been described as indirect immunofluorescence (IIF) staining pattern which is uniformly distributed throughout the interphase nucleus and, most notably, are localized on metaphase chromosomes. We undertook this study to examine the frequency of anti-DFS70 antibodies in a large number of healthy blood donors and to evaluate the significance of these antibodies, accordingly.

Methods
Sera of 500 healthy blood donors (459 men, 41 women) were analyzed for ANAs and anti-DFS70 antibodies by IIF with HEp-2 cells as a substrate.

Conclusions
ANAs were determined in 0.8% of all blood donors by IIF. Positive results were detected in different serum dilutions: Three subjects (0.6%) 1:320 nucleolar pattern, and 1 subject (0.2%) 1:1000 homogeneous/nucleolar pattern. There were 3 anti-DFS70 antibody positive blood donors. In this study, anti-DFS70 autoantibody was detected 0.6% in healthy blood donors. Among anti-DFS70 antibody-positive subjects the percentage of male was 0.65% (3 of 459 subjects). The prevalence of anti-DFS70 antibody positivity decreased with increasing age. According to our results, anti-DFS70 autoantibody is detected more commonly in male than female healthy blood donors. More comprehensive studies are needed to be done to investigate the anti-DFS70 autoantibody specificity for healthy individuals.
ANTI-DFS70 ANTIBODIES IN SYSTEMIC AUTOIMMUNE RHEUMATIC DISEASES

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Background
Systemic autoimmune rheumatic diseases are major cause of morbidity and mortality. The presence of various autoantibodies, such as antinuclear antibodies produced against intracellular antigens, are characteristic of systemic autoimmune rheumatic disease (SARD).

Objectives
In this study, we aimed to investigate the prevalence of anti- DFS70 antibodies in SARD and to evaluate the significance of these antibodies, accordingly.

Methods
In our study we created test groups as; 107 healthy blood donors, 418 Rheumatoid arthritis (RA), 101 Systemic Lupus Erythamatosus (SLE) patients, 36 Systemic sclerosis patients, 71 Sjögren syndrome patients, 43 Ankylosing spondylitis and 2555 Connective tissue disease (CTD) pre-diagnosed patients. Serum samples were evaluated by indirect immunofluorescence assay (IFA). Anti-DFS70 positive samples were also tested by ELISA test.

Conclusions
Anti DFS70 antibody was detected 1.4% in RA, 2.97% in SLE, 1.4% in Sjögren syndrom and 1.05% in disease group with pre-diagnosed CTD. It was not detected in AS and SSc patient groups. Totally 39 out of 47 samples were also found to be anti DFS70-positive by ELISA method. According to our results, anti-DFS70 autoantibody is detected more commonly in certain disease sub-groups of SARD than healthy individuals.
Background

Clostridium difficile is a leading cause of infectious diarrhea and pseudomembranous colitis. Standard treatment for C. difficile infections (CDI) includes the administration of antibiotics. However, the rate of CDI recurrence after initial therapy is approximately 20% and even increases when antibiotic therapy is repeated.

Objectives

Therefore, alternatives or additional measures to current antibiotic therapy are required to achieve lower rates of CDI recurrence. Vaccination is one of prevention measures by which low recurrence rate is expected. In this study, we demonstrate the potential of nontoxigenic C. difficile membrane fraction (ntCDMF) as a vaccine candidate.

Methods

Mice immunized with ntCDMF showed serum IgG and intestinal IgA titers of 4,660 and 156, respectively, as compared to 30 and 13, respectively, for non-immunized controls. In Caco-2 cell adhesion assay for 9 different C. difficile strains, the number of adherent bacterial cells pretreated with the immune serum was 5.8 – 9.3 CFU per 100 Caco-2 cells, which was significantly lower than in the case treated with non-immunized serum (21.1 to 36.6). Similar findings were observed for intestinal fluid, where the number of adherent bacterial cells ranged from 19.6 - 29.1 and 25.9 - 42.0 CFU per 100 Caco-2 cells for immunized and control mice, respectively.

Conclusions

These results support the idea of ntCDMF-derived C. difficile vaccine capable of preventing the adhesion for C. difficile to the intestinal cells, as well as emphasize its potential as a preventive measure against CDI.
Background
This paper reviews the toxicity and antibacterial activity of ten cyanobacterial strains isolated from fresh waters in Serbia.

Objectives
All investigated cyanobacterial strains showed toxic effect to *A. salina*. The highest toxicity was observed in the case of strains belonging to *Microcystis* genera (*Microcystis* L1 and P1) with detected larval mortality of 98% and 96%, respectively. The observed LD50 values were from 0.24 mgml\(^{-1}\) (*Anabaena* P1) to 3.37 mgml\(^{-1}\) (*Oscillatoria* K3). The intracellular microcystin concentrations, calculated as microcystin-LR equivalents, were in the range of 0.1–3.96 µgmg\(^{-1}\) dry weight. All cyanobacterial strains revealed antibacterial activity and inhibited the growth of at least one bacterial species. The intracellular extract of *Aphanizomenon* K2 strain showed the most evident inhibitory effects to 6 bacteria. The MIC values of the tested strains were in the range of 0.00014 gL\(^{-1}\) - 0.5805 gL\(^{-1}\).

Methods
The toxicity of intracellular extracts of cyanobacterial strains was tested by *Artemia salina* bioassay and expressed as the percentage of larval mortality. The intracellular hepatotoxin (microcystin) content of cyanobacterial strains was determined by colorimetric protein phosphatase 1 inhibition assay (PP1). Methanolic intracellular extracts of cyanobacterial strains were tested for antibacterial activity using agar diffusion method and broth microdilution assay.

Conclusions
The results suggest that toxic effect on larvae *A. salina* found in the most tested strains is likely to be a consequence of their microcystin production. The obtained results indicate that toxins are probably not responsible for the expressed antibacterial activity of the tested strains.
Background

Eicosapentaenoic acid (EPA) is an essential omega-3 polyunsaturated fatty acid (PUFA) well known for its beneficial effects on human and animal health. EPA is present at relatively high levels in a number of marine animals with its primary dietary source being microalgae and some bacteria. This essential bioactive plays a critical role in marine life; however, its transfer between different trophic levels is yet to be fully understood.

Objectives

A simple and affordable method to study and trace the fate of EPA through the trophic food chain was developed.

Methods

*Shewanella sp.* is a marine bacterium, known to produce EPA [1, 2]. In this work, we describe a simple method to label the EPA present in the bacterial strain IRL 567 (*Shewanella sp.*) with the stable isotope $^{13}$C. Labelling assays were performed at small scale (100 ml shake flask) and bench scale (1lt stirred tank bioreactor). By incubating the bacteria with $^{13}$C-acetate in culture medium, we demonstrated that EPA is *de-novo* synthesized utilizing the simplest carbon source -acetate- as precursor. $^{13}$C incorporation into the EPA molecule was determined by mass spectrometry (ESI TOF MS), finding that 95.5% of synthetized EPA being labelled in the shake flask and 88.6% in the bioreactor, yielding a concentration of labelled EPA of 5.36 mg/l and 2.90 mg/l, respectively, with between 2 to 8 $^{13}$C atoms contained in EPA structure.

Conclusions

Labelled EPA could be a valuable research tool to understand the assimilation and metabolic routes of fatty acids, essential components in human and animal diets.
Background
The occurrence of candidemia is on a rise worldwide. Non-albicans Candida species have emerged as major causes of candidemia in many countries. Added to it is the problem of antifungal resistance in Candida isolates.

Objectives
The aim of our study was to investigate the isolation of Candida spp in blood cultures and to evaluate their antifungal susceptibility during a 8-year period (2007-2014) in a tertiary care hospital.

Methods
The blood cultures were incubated in the automated blood culture system BACTEC9240 (Becton Dickinson). Positive blood cultures were examined microscopically directly for yeast or pseudohyphae and subcultured on Sabouraud dextrose agar (Liofilchem Italy). Candida isolates were identified using automated VITEK 2 system (bioMerieux) or Api 20CAUX (bioMerieux). Antifungal susceptibility was carried out by automated VITEK 2 system (bioMerieux) using ASTM01 or AST Y07 test card.

Results: During the study period there were 67 candidemia cases. The 51.35% of candidemias occurred in the ICUs, the 40.54% in the medical wards and the rest of 8.11% in surgical wards. C. parapsilosis was the predominant species (35.15%), followed by C. albicans (32.43%), C. glabrata (10.81%), C. tropicalis (8.12%) and C. norvegensis, C. guilliermondii, C. kefyr, C. famata, Cryptococcus neoformans with (2.70%). All tested isolates were susceptible to amphotericin B and voriconasole. Among non-albicans strains increasing resistance to fluconasole was found: C.parapsilosis (8.33%), and C. tropicalis (8.33%).

Conclusions
Candidemia was more frequent in ICUs followed by medical and surgical wards. C. parapsilosis was the predominant cause of candidemias in our Hospital. Amphotericin B and voriconasole were active against all tested isolates.
ASSESSMENT OF IMMUNE PROTECTIVE CAPACITY OF THE RECOMBINANT IRON SUPEROXIDE DISMUTASE (FESOD) FROM BORDETELLA PERTUSSIS

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Background
Whooping cough (pertussis) is a highly contagious respiratory infection caused by Bordetella pertussis. Although availability of effective pertussis vaccines seems to decrease the incidence of the disease, B. pertussis circulation in population has not been eliminated. Thus, finding new protein candidates with high immune protective capacities is necessary to enhance the efficacy of current acellular pertussis (Pa) vaccines.

Objectives
The aim is to evaluate immune protective capacity of iron superoxide dismutase (FeSOD) protein from B. pertussis with the aim of developing new generation pertussis vaccines.

Methods
FeSOD gene was cloned, expressed in Escherichia coli and the recombinant protein was purified. FeSOD was formulated with Aluminum hydroxide (Alum) or Monophosphoryl Lipid A (MPLA) and mice were immunized intraperitoneally. IgG1, IgG2a and IFN-γ levels were determined and bacterial colonization on mice lungs were evaluated.

Conclusions
IgG1 and IgG2a responses were significantly increased in both mice groups immunized with FeSOD-Alum and FeSOD-MPLA, the level of IgG2a was relatively higher in mice vaccinated with FeSOD-MPLA. Immunization with FeSOD-MPLA formulation provided a significant decrease in bacterial count in mice. Moreover, antigen specific-IFN-γ response was significantly increased in the group vaccinated with FeSOD-MPLA. These findings, altogether, suggested that the recombinant FeSOD protein formulated with MPLA can be a possible acellular pertussis vaccine candidate.

References:
Background

MIRRI (Microbial Resources Research Infrastructure, www.mirri.org) is an initiative within the European Strategy Forum on Research Infrastructure (ESFRI) that includes 16 partners (14 public microbial Culture Collections / mBRCs), supported by 21 collaborators; the preparatory phase is currently funded by the European Commission.

MIRRI aims to construct a pan-European distributed infrastructure that boosts research and development in the field of biotechnology by improving access to the microbial resources present in European public collections, their associated data and expertise.

Objectives

One of the MIRRI objectives is to create clusters bringing together the expertise available at different partner and non-partner institutions to respond to concrete demands of the MIRRI stakeholders. These expert clusters will provide solutions in aspects such as regulatory framework (e.g. biosecurity, ABS, IPR), managerial best practices, taxonomy or microbial applications, among others.

Methods

To design the content, composition and rules of operation of the clusters, the relevant stakeholders have been consulted (face to face meetings, workshops and surveys). An analysis of the comparable RIs of the ESFRI 'Food and Health' group was performed to bridge the gap between information sharing and facilitating a collaboration landscape.

Conclusions

Here, we present the concept of the MIRRI Clusters of expertise as a tool to share and generate knowledge within the MIRRI stakeholder community (culture
collections, policy makers, scientists, bioindustry, etc.) to foster innovation within science, research and development. The authors gratefully acknowledge the contribution of the MIRRI consortium and the consulted persons to this work.

This project has received funding from EU FP7 (grant agreement 312251).
Background
The bacterial flagellum is a complex organelle requiring the coupling of gene expression to the assembly pathway. In *Escherichia coli* and *Salmonella enterica* serovar Typhimurium the *flhDC* operon encodes the essential FlhD6C2 flagellar-specific master transcriptional regulator. The activity of FlhD6C2 is influenced by many flagellar-specific and global regulatory stimuli. FlhD6C2 activity responds to cellular and environmental factors through regulators including OmpR, RcsB and CRP. Flagellar-specific signals known to influence FlhD6C2 activity are controlled by the action of FliT and FliZ. *flhDC* expression is also sensitive to cell growth.

Objectives
Our objective in this study was to investigate the impact of growth rate on flagellar formation in *E. coli* and *S. Typhimurium*

Methods
Our methods to study our objective included the use of steady-state chemostat cultures. Our measure was flagellar abundance using a functional FliM-GFP fusion protein. We will show that flagellar abundance correlates with growth rate, where faster growing cells produce more flagella in both species. We will present data showing that the impact of transcriptional, post-transcriptional and flagellar-specific regulation of *flhDC* with respect to the growth rate response of the flagellar system leads to a marked difference in flagellar abundance when the species are compared.

Conclusions
Our data suggests that even though both flagellar systems have a high degree of genetic similarity the way *E. coli* and *S. Typhimurium* assimilate signals during flagellar regulation may reflect the lifestyle of these two bacterial species.
Bacterial nanomachines

TYPE VI SECRETION IN THE SOIL BACTERIUM PSEUDOMONAS PUTIDA: ROLE IN BACTERIAL COMPETITION

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Background

Bacterial type VI secretion systems (T6SSs) are recently discovered nanomachines used to inject effectors into prokaryotic or eukaryotic cells. T6SSs are therefore involved in both inter-bacterial competition and bacterial pathogenesis.

Objectives

The aim of this work is the analysis of T6SS in the soil bacterium Pseudomonas putida. P. putida is known for its capacity to colonise the root of crop plants providing growth advantages to the plant and, importantly, protection against plant pathogens. Since T6SS is mainly used by environmental bacteria for interbacterial competition, we analysed whether this secretion system of P. putida might be involved in such protection.

Methods

- In silico analysis of P. putida KT2440 genome to identify potential T6SSs
- Competition assays to determine P. putida T6SS activity and bacterial targets.
- qRT-PCR and transcriptional fusions to study P. putida T6SS expression and regulation

Conclusions

P. putida contains three putative T6SSs named as H1, H2, and H3. These clusters contain the genes encoding the T6SS conserved core components and some accessory proteins, including regulatory proteins and toxins-immunity pairs. Additional T6SS-related genes are found scattered on the chromosome. At least the H1-T6SS is active and used by P. putida to kill other bacteria, including the phytopathogen Pseudomonas syringae. Expression of this system is maximal in stationary phase and it is controlled by the global regulators RetS and GacS-GacA, and by two alternative sigma factors, RpoS and RpoN.
A PSEUDOMONAS FLUORESCENS TYPE VI SECRETION SYSTEM IS INVOLVED IN ANTIBACTERIAL ACTIVITY

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Background

To persist in an ecological niche, bacteria have developed several strategies to enable them to resist in the environment. One of these mechanisms able to struggle with other bacterial species is Type VI Secretion System (T6SS). This allows the bacterium to inject toxins directly into prey's cytoplasm and could cause death. A mucoid environmental strain of *Pseudomonas fluorescens*, MFE01, has antibacterial activity carried by T6SS. MFE01 is not virulent against eukaryotic cell models (amoebas, plant or animal cell models). Currently, genomic analysis reveals only one T6SS core component cluster and three *hcp* genes were found (named *hcp1*, *hcp2* and *hcp3*).

Objectives

Aims of this study are to assign a role to each Hcp protein and to test the potential application of this secretion system to prevent infection by nosocomial strain (like *Pseudomonas aeruginosa*) in burn wound.

Methods

Mutations of *hcp1*, *hcp2* and *hcp3* were performed. Furthermore, T6SS apparatus was also inactivated by *tssE* gene mutagenesis. Competitor activity were established in different conditions: immobilized experiment, swarming or swimming conditions. Hereafter, we will use Hacat keratinocyte model to study the ability of MFE01 to protect against infection by etiologic agents.

Conclusions

MFE01 T6SS plays a crucial role in bacterial competition in ecological niche. Hcp2 was involved in antibacterial activity whereas Hcp1 immobilized prey cell. Hcp1 and Hcp2 proteins had a synergic effect which Hcp1 inhibits prey cell mobility and then Hcp2 can killed it. Presently, no role is attributed to Hcp3.
Background
Selenium oxyanions can be reduced by diverse microorganisms resulting in the formation of selenium nanoparticles (SeNP). This process has been mostly investigated using pure bacterial cultures. However, using pure cultures for treatment of wastewater containing selenium oxyanions is not recommended. Anaerobic granular sludge, which consists of aggregates of mixed microbial communities, may also reduce selenium oxyanions and would thus be appropriate for developing treatment systems.

Objectives
The aims were: (i) to investigate which microorganisms prevailed in a granular sludge that was exposed to selenite and reduced this oxyanion to SeNP, (ii) to investigate at the macro- and micro-scale the location of the produced SeNP, and (iii) to identify proteins associated to the SeNP.

Methods
High-throughput pyrosequencing, electron microscopy and proteomic approaches were employed.

Conclusions
A large change in the microbial community occurred when the inoculum granular sludge was exposed to and reduced selenite during a 20 day period. Most abundant microorganisms in the selenite reducing granular sludge were affiliated to Veillonellaceae and Pseudomonadaceae families. Most of selenium bioreduction occurred in the outer layer of the granular sludge, where most biomass contents are localized. At the micro-scale, electron tomography showed that SeNP were produced inside the cells. A proteomic analysis of extracted SeNP revealed that most of their associated proteins were proteins found in the outer and inner membranes. The used granular sludge readily reduced selenite and the combined results allowed proposing mechanisms for the bacterial synthesis of SeNP in these mixed microbial aggregates.
Bacterial nanomachines

YERSINIA PSEUDOTUBERCULOSIS TYPE III SECRETION IS RELIANT UPON AN AUTHENTIC DUAL FUNCTIONAL N-TERMINAL YSCX SECRETOR DOMAIN.

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Background
Various Gram negative bacteria employ a type III secretion system (T3SS) to deliver effector proteins into eukaryotic cells to form mutualistic or pathogenic interactions with their host. A T3SS is comprised of about 25 different proteins many of whose functions still remain obscure. In \textit{Yersinia} Ysc-Yop T3SS, YscX and YscY are two proteins which have not been well characterised. However, a Ysc-Yop T3SS is non-functional in the absence of either \textit{yscX} or \textit{yscY}, suggesting that both YscX and YscY are crucial Ysc-Yop T3SS constituents.

Objectives
An attempt to understand the role of YscX and YscY was initially investigated by studying the functional interchangeability between genetically conserved members of the YscX-YscY protein families. It suggested that YscX might be functionally unique to \textit{Yersinia} despite reciprocal binding with YscY family members. Intrigued by the specificity of YscX function, we scrutinized the role of YscX N-terminus in secretion of itself and other T3S proteins.

Methods
Site directed mutagenesis and defined domain swapping revealed YscX N-terminus to be a critical aspect of \textit{Yersinia} T3S as it prevented the polymerisation of surface localized YscF needle. This was neither due to a defect in YscX secretion potential nor any observable defect in forming a bipartite interaction with YscY or a tripartite interaction with YscV as measured by standard protein interaction assays.

Conclusions
Therefore, the YscX N-terminus must perform dual functions; on the one hand it is a secretion recognition motif and on the other, a non-redundant recognition signal important for the correct assembly of Ysc-Yop T3SS.
NEW INSIGHTS ON THE ROLE OF SPAS AUTOCLEAVAGE IN SALMONELLA TYPE III SECRETION SYSTEM

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Background

The type III secretion system (T3SS also known as ‘injectisome’) is organized in three main structures: a multi-ring basal body that houses the export apparatus, a needle, and a translocon in the host cell membrane. To assemble a functional T3SS, specific substrates must be targeted to the apparatus in the correct order. The substrate specificity switch from early to intermediate substrates occurs once the needle has reached its full length.

The Salmonella T3SS export apparatus protein SpaS, is a multifunctional component involved in the switch. It consists of a transmembrane region and a C-terminal cytoplasmic domain that undergoes autocleavage at a conserved NPTH motif.

Objectives

Elucidating the timing of autocleavage and why it is critical for SpaS function.

Methods

In this study we test expression and secretion levels of T3 proteins by immunoblotting and blue-native gel electrophoresis to study the composition of the assembled T3SS. Our data are complemented by injectisome purification and its visualization by electron microscopy.

Conclusions

Studying autocleavage in its physiological context we could demonstrate that the kinetics of SpaS autocleavage are not regulated by injectisome state of assembly or secretion and that the autocleavage mechanism is independent from needle length control. Rather, autocleavage is a fast process occurring before SpaS incorporation into the injectisome base, which is demonstrated by the fact that a functional injectisomes can assemble on pre-expressed fully cleaved SpaS. We also found that autocleavage at the NPTH motif is not a prerequisite for switching since also SpaS cleavage by an endogenous site-specific protease leads to functional switching.
Background

Magnetotactic bacteria are motile prokaryotes with the ability to swim along the Earth’s geomagnetic field lines due to the presence of intracellular structures called magnetosomes. Magnetosomes are composed of a mineral core of magnetite or greigite surrounded by a 2-4 nm thick lipid bilayer membrane (1). In recent years, the magnetosomes have attracted great attention because of their potential usefulness in biotechnological and biomedical applications (2).

Objectives

In this work, we explore different methods to isolate magnetosomes from Magnetospirillum gryphiswaldense and check their biocompatibility on eukaryotic cells.

Methods

Two mechanic cell lysis procedures are compared based on the amount and quality of the magnetosomes obtained. The isolated magnetosomes were structurally characterized by TEM as cuboctahedral magnetite particles with an average diameter of 45(±3) nm (3). The aggregation behaviour of magnetosomes in solution is followed by Dynamic Light Scattering technique. The Zeta potential was always lower than -30mV indicating moderately stability. Infrared spectroscopy (FT-IR) revealed the presence of poly-β-hydroxybutyrate under some extraction conditions.

The biocompatibility is evaluated in vitro in cytotoxicity assays on murine macrophage ANA-1 cell line. Cells are incubated with 30 pg of magnetite per cell and viability is measured using flow cytometer by annexin-propidium staining. Magnetosomes labeled with fluorescein isothiocyanate are used to track the location of the internalized particles in the cells.

Conclusions
No evident signs of cytotoxicity were found after 48 hour of incubation.


FEMS-1737
Bacterial pathogenicity

DETERMINATION OF VIRULENCE OF FOOD-BORNE SALMONELLA SEROTYPES BY USING CAENORHABDITIS ELEGANS AS A MODEL SYSTEM
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Background
Salmonellosis caused by S.enterica is an important food-borne disease worldwide and pathogenicity and antibiotic resistance of these bacteria is an important health concern. C.elegans is a current nematode model to test pathogenicity of Salmonella.

Objectives
We determined pathogenicity, antibiotic resistance, serotypes and plasmid profiles of 32 non-typhoidal food-borne Salmonella isolates and searched for a correlation between these characteristics among the isolates.

Methods
Pathogenicity of isolates was tested in C.elegans model. Nematodes (20/group) were fed with Salmonella isolates grown on NGM agar (0.6 at OD600). The time required for 50% of nematodes in each group to die (TD50) was calculated. S. Typhimurium ATCC14028 and E.coli OP50 were used as positive and negative controls, respectively. Plasmids were isolated by Kado-Liu method and examined by agarose gel electrophoresis (0.7%). Serotyping of isolates was done by slide agglutination test and their antibiotic resistance was investigated by disc diffusion method.

Conclusions
26 of Salmonella isolates belonged to Infantis, 4 to Enteritidis, 1 to Kentucky and 1 to Telaviv serotypes. 6 isolates harboured 1-3 plasmids ranging from 1.2 to 35.8 kb. TD50 values were 4.2±0.5 days for S. Typhimurium ATCC 14028 fed, 8.0±0.02 days for E.coli OP50 fed, and between 3.5-7.2 days for food-borne Salmonella fed nematodes. The significance of differences in TD50 values were tested by t test (p<0.05). 22 of isolates were pathogenic in C.elegans; 20 of these isolates were multi-drug resistant, 2 were ciprofloxacin resistant and 21 were nalidixic acid resistant. Multi-drug resistant isolates identified as pathogens in nematodes may pose serious health risk for consumers.
Bacterial pathogenicity

LECTIN PRODUCTION BY ACINETOBACTER BAUMANNII CLINICAL AND ENVIRONMENTAL ISOLATES

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Background
Acinetobacter baumannii is an emerging opportunistic pathogen which cause severe infections in immunocompromised patients and exhibit resistance to many conventional antibiotics. Factors responsible for A. baumannii pathogenesis are numerous; some of them are biofilm formation and hemagglutination, which are dependent on lectin production.

Objectives
The aims were to determine the lectin production ability of the A. baumannii, to determine the lectin specificity to different blood types, and to semi-quantify lectin produced by A. baumannii.

Methods
The study was carried out with 3 reference strains (ATCC BAA-747, ATCC 19606, NCTC 13423), 8 environmental and 20 MDR A. baumannii wound isolates. Lectin production screening was performed using the hemagglutination assay. The supernatant of overnight cultures were examined using 3 % erythrocyte suspension of different blood types (O Rh+, A Rh+ i B Rh+) and results were recorded after 30 min. The semiquantitative determination of lectin production was examined by ex vivo modified hemagglutination assay with erythrocyte of A blood type in microplates using two-fold culture supernatant dilutions (from 1/2 to 1/2048).

Conclusions
All examined isolates were able to produce lectins in vitro and expressed hemagglutination activity to all examined blood types. The semiquantitative metod showed that titer is in a range from 1/4 to 1/64. The clinical strains produed lectines in slightly higher amount comparig to the environmental isolates, but without significant statistical difference (P>0.05). The results indicate that lectine production can be a new target for development of novel anti-Acinetobacter agents.
MOLECULAR CHARACTERIZATION OF KLEBSIELLA OXYTOCA STRAINS ISOLATED FROM PATIENTS WITH ANTIBIOTIC-ASSOCIATED DIARRHEA

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Background

The main complication of colitis would be seen after antibiotic treatment by β lactams, quinolone and aminoglycosides antibiotics. Recently, Klebsiella oxytoca has been known to cause this type of diarrhea.

Objectives

Investigating the prevalence and characterizations of K. oxytoca isolated from patients with antibiotic associated diarrhea. K. oxytoca isolates were also tested for cytotoxin production.

Methods

This study was carried out from May 2011 to Dec 2013. Fecal samples were collected from hospitalized patients who received antibiotics. Initial cultivation was performed on specific media. Clinical isolates were confirmed by PCR using the specific K. oxytoca polygalacturonase (pehX) gene. The double disk diffusion test was used to detect ESBLs producing strains. Tracking ESBLs encoding genes were performed by PCR. The bacteria`s cytotoxin production was conducted using cell culture on Hep2 cell lines.

Conclusions

Five (12%) isolates had cytotoxin activity less than 30%, 12 (30%) strains had moderate cytotoxin activity between 30 and 60% and 23 (58%) strain had cytotoxin activity equal to/or greater than 60%. The cytotoxin producing K. oxytoca can be described as one of the causes of antibiotic induced colitis. Drug discontinuation and creating opportunities for the establishment of intestinal normal flora or appropriate medication after antibiogram would perform better chance earlier in patients with hemorrhagic colitis caused by antibiotics.
Background

*Haemophilus parasuis* is the etiological agent of porcine Glässer's disease.

**Objectives**

Based on previous studies, three recombinant outer membrane proteins of *H. parasuis* serovar 5 (P2, P5 and D15) were designed and produced for studying their immunogenicity and potential use as vaccine antigens.

**Methods**

Each gene was amplified and cloned into a pBAD/TOPO vector, this plasmid being enable to express each protein with a tag of histidines for facilitating the subsequent purification. *Escherichia coli* was transformed with the vector and then, it was cultured with arabinose to induce protein expression. As the three proteins were expressed at insoluble fraction, it was necessary to develop a denaturation process using urea, in order to solubilize and refold them. Refolding was performed at the same time as purification by means of immobilized nickel-histidine affinity chromatography. Each of these proteins was retained in the stationary phase and washed with buffers containing decreasing urea concentrations until removing totally urea; the proteins were then eluted with imidazole. Rabbits were used to evaluate the capacity of these proteins to induce a humoral response before testing them in Glässer's disease natural host. Groups of animals were immunized thrice with each of these proteins, and blood was collected after each inoculation. Sera were used to perform indirect ELISAs, in which purified proteins were used as antigen for coating plates.

**Conclusions**

An increasing in antibody titres with seroconversion can be found after immunizations for recombinant P2, P5 and D15 proteins.
Background

Crohn's disease (CD) is a chronic gastrointestinal inflammatory disease. Adherent invasive Escherichia coli (AIEC) strains, which have the capacity to attach and invade intestinal epithelial cells, and also survive within macrophages; have been associated with CD.

Objectives

To identify immunogenic proteins in the outer membrane of AIEC isolates obtained from Chilean patients with CD, and determine their role in pathogenicity.

Methods

AIECs were isolated from biopsies of Chilean patients with CD. AIEC outer membrane proteins (OMPs) were obtained from cultures grown at 20 °C and 37 °C, and separated by SDS-PAGE. Immunogenic proteins were identified by immunoblot using sera from patients with CD and MALDI-TOF/TOF. Genes coding identified proteins were knocked out and mutant strains were characterized according to: 1) Capacity to adhere and invade intestinal cells, and to survive within macrophages. 2) Differential expression of outer membrane proteins by (2D SDS-PAGE and MALDI-TOF/TOF. 3) Capacity to stimulate production of inflammatory cytokines.

Conclusions

Siderophore receptors IutA and ChuA were identified as immunogenic OMPs. No effect in adherence and invasion capacity was observed after knocking out their genes, however, mutant strains did not survive within macrophages. Additionally, mutation altered the outer membrane profile by apparently affecting expression of virulence-associated proteins. A decrease in IL-1, IL-2, IL-12, IL-4 and IFN-gamma levels was detected in supernatants of macrophages infected with the mutant, while TNF- alpha level did not change. Therefore, iutA and chuA would be required for AIEC intracellular survival in the host, in order to capture iron and express virulence factors. Finally mutating these receptors the strain becomes unstable and avirulent bacteria, which can not survive oxidative stress because other virulence genes are repressed by iron deficiency.
Background
Klebsiella pneumoniae (K. pneumoniae) is one of the main pathogens in nosocomial infections. Type 3 pilus (T3P) is a virulence factor required for biofilm formation and adherence to epithelial cells. Three promoters control the expression of T3P: mrkA (codes for the pilin) mrkHI (code for two activator proteins) and mrkJ (a repressor protein). H-NS nucleoid protein structures the bacterial genome, mainly Gamma-proteobacteria. Hypermucoviscosity that is attributed to the capsule formation, is a hallmark of pathogenic strains. No reports have been described for the role of H-NS protein in regulating T3P and capsule in K. pneumoniae.

Objectives
1. Determine the role of H-NS protein in transcription of T3P.
2. Analyze both the biofilm and capsule formation in the absence of H-NS.

Methods
Transcriptional expression of mrkA, mrkH, mrkI and mrkJ genes was carried out by qRT-PCR. Using EMSA, promoter regions (PCR) of mrkA, mrkJ and mrkHI were incubated with purified H-NS protein and DNA-proteins complexes were observed. Biofilm formation was quantified by the method of 96-wells polystyrene microplates. The abundance of capsule was determined by measuring spectrophotometrically the supernatants of bacterial cultures at exponential phase.

Conclusions
K. pneumoniae hns mutant affected the growth at exponential phase and resulted in a hypermucoviscosity phenotype. The absence of H-NS affected the biofilm formation and differentially regulated the mrk promoters: derepressed and repressed to mrkHI-mrkJ and mrkA, respectively. H-NS directly bound to the three promoters of T3P. The H-NS protein is a crucial regulator of the main virulence determinants of K. pneumoniae.
EXOGENOUS SUPPLEMENTATION WITH BRANCHED-CHAIN AMINO ACID PROMOTES AEROBIC GROWTH OF SALMONELLA TYPHIMURIUM UNDER NITROSATIVE STRESS CONDITIONS
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Background
An intracellular pathogen *Salmonella enterica* serovar Typhimurium can detoxify cytotoxic nitric oxide (NO) produced in phagocytes of host animals by their metabolism depending largely on the flavohemoglobin Hmp. In *hmp* mutant *S. Typhimurium*, NO causes deficiency in amino acids including branched-chain amino acids (BCAA). In the BCAA biosynthetic pathway, iron-sulfur cluster-containing enzymes dihydroxyacid dehydratase (IlvD) and isopropylmalate isomerase (LeuCD) have been implicated to be targets of NO, suggesting the importance of BCAA metabolism in the NO-induced amino acid auxotrophy.

Objectives
This study aimed to determine the effect of BCAA supplementation on *Salmonella* resistance to NO under different oxygen conditions, and to test physiological roles of IlvD and LeuCD in the intramacrophage survival of *Salmonella* in the presence or absence of BCAA supplementation.

Methods
Gene mutations deficient in *ilvD* and *leuCD* were constructed in *S. Typhimurium*. With mutant *S. Typhimurium* having combinations of these gene mutations and *hmp* mutation, we measured the effect of BCAA on the susceptibility to nitrosative stress, the NO consumption rate, and the intramacrophage growth.

Conclusions
Under nitrosative stress conditions, BCAA supplementation restored the growth of *hmp* mutant and mutants further lacking IlvD and LeuCD in aerobic and semiaerobic cultures, but not in anaerobic cultures. Intracellular survival of *ilvD* and *leuCD* mutants in macrophages was also promoted by BCAA supplementation. Results suggest that the NO-induced BCAA auxotrophy of pathogenic bacteria due to inactivation of iron-sulfur enzymes in the BCAA biosynthetic pathway could be rescued by bacterial taking up exogenous BCAA available in oxic environments of host animals.
Pasteurella multocida toxin (PMT) is mitogenic for many cell lines. It acts via modification of three of the four families of heterotrimeric G-proteins which are Gq, G12/13 and Gi. PMT has been shown to induce anchorage independent cell growth, enhance anti-apoptotic proteins, increase β-catenin signalling in a differentiating adipocyte system, and up-regulate JAK/STAT and MAPK signalling pathway, all of which are implicated in various cancers. These different effects of PMT suggest it may have a carcinogenic potential.

Objectives
Here we are studying PMT as a potential carcinogen by chronically treating cells in vitro with PMT to analyse its involvement in cell transformation.

Methods
The following end points are being monitored to assess the carcinogenic potential of PMT: cell proliferation, stress fibre formation, cellular morphology, levels of normal G-proteins, levels of PMT-modified G-proteins, and anchorage independent growth. Results achieved so far show that PMT-treated cells have an increased cell number, show morphological changes, and differences in protein levels compared to the untreated cells.

Conclusions
Together the data suggests that prolonged exposure to G-proteins modified by chronic treatment with PMT can set cells along a transformation pathway.
SINC, A TYPE III SECRETED EFFECTOR OF CHLAMYDIA PSITTACI, TARGETS EMERIN AND THE INNER NUCLEAR MEMBRANE OF INFECTED CELLS AND UNINFECTED NEIGHBORS

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Background

Chlamydia species encode a type III secretion (T3S) system that enables the translocation of effector proteins into the host cell cytosol, where they target host pathways to benefit the pathogen. We have identified SINC, a novel T3S effector of Chlamydia psittaci, a highly infectious avian pathogen, and medically significant zoonotic pathogen of humans.

Objectives

To identify and characterize the eukaryotic target(s) of SINC.

Methods

Immunofluorescence, immunoelectron microscopy, differential digitonin permeabilization, inhibitors of nuclear transport, BioID (in situ biotinlylation analysis of effector-BirA-proximal eukaryotic targets, streptavidin affinity purification, mass spectrometry).

Conclusions

SINC uniquely targets the nuclear envelope (NE) of C. psittaci-infected cells and uninfected neighboring cells. Digitonin-permeabilization studies of infected or SINC-GFP-transfected HeLa cells indicate SINC targets the inner nuclear membrane (INM). SINC localization at the NE was blocked by importazole, confirming SINC import into the nucleus. Candidate partners were identified by proximity to biotin ligase-fused SINC in HeLa cells and mass spectrometry. Several candidates were identified with high confidence including the nucleoporin ELYS, lamin B1 and four INM proteins (emerin, MAN1, LAP1 and LBR), suggesting SINC interacts with host proteins that control nuclear structure, signaling, chromatin organization and gene silencing. GFP-
SINC association with the native LEM-domain protein emerin, a conserved component of nuclear ‘lamina’ structure, was confirmed by GFP pull-down. We conclude that SINC is a novel bacterial effector with the capability of globally altering nuclear envelope functions in the infected host cell and neighboring uninfected cells. These properties may contribute to the aggressive virulence of *C. psittaci*. 
Background

Colorectal cancer (CRC) remains one of the most prevalent cancers in the Western world. However, the factors leading to the development of colon cancer have not been determined, with genetics, diet and other environmental factors all likely to play a role. Mechanisms by which Bacteroides fragilis and Fusobacterium nucleatum may drive colonic tumour development have been identified. Some B. fragilis isolates contain a metalloprotease toxin known as BFT or fragilysin. BFT has been demonstrated to promote colonic hyperplasia in animal models. F. nucleatum is of relevance due to the consistent finding across several studies that it is enriched on CRC tumours. Its FadA adhesin can mediate invasion into the cell with a resultant increase in expression of host oncogenes.

Objectives

Advances in next generation sequencing technologies have led to a high resolution of the CRC microbial community. The aim of this study is to use a hypothesis led approach to focus on potential key isolates and genes in the development of colonic tumours.

Methods

We are compiling a comprehensive collection of human colonic tissue from patient cohorts representing different stages of disease. These samples are being analysed using a PCR-based assay to determine the prevalence of selected bacterial taxa and genes. Results show primer specificity, PCR optimization and sensitivity of detection that have been experimentally determined.

Conclusions
Overall this work describes the design and validation of a panel of primers with the intention of analysing a fresh collection of human tissue. Preliminary findings using a collection of pilot samples will be presented.
Background

Extraintestinal pathogenic *E. coli* (ExPEC) are a major source of urinary tract infection, newborn meningitis and sepsis in humans. Avian pathogenic *E. coli* (APEC) cause colibacillosis, a complex systemic infection in poultry. Certain human ExPEC and APEC cannot be clearly distinguished based on molecular epidemiology and their virulence-associated genome content. APEC are considered as a reservoir of virulence- and resistance genes for human ExPEC and a zoonotic risk cannot be excluded.

Objectives

To characterize potential differences between APEC and chicken commensal *E. coli*, we compared the virulence- and antibiotic resistance-associated genome content of commensal *E. coli* isolates from healthy chicken with APEC strains.

Methods

*E. coli* isolates from tracheal and fecal samples of healthy chicken were collected and subjected to virulence gene and antibiotic resistance profiling. The genomes of 127 commensal and APEC isolates were sequenced.

Conclusions

The virulence gene content of commensal chicken and APEC isolates was similar. Comprehensive genome comparison of APEC and chicken commensals will extend our knowledge of bacterial traits that could contribute to pathogenesis and antibiotic resistance. Interestingly, animal husbandry has a great impact on the spread of antibiotic resistance determinants among chicken *E. coli* isolates. As APEC have a zoonotic potential and serve as potential gene pool for human ExPEC, the increased presence of antibiotic resistance determinants in *E. coli* from conventionally raised...
chicken can facilitate the spread of multiresistant human ExPEC.
PROPERTIES OF THE DSBA PROTEIN (CLA0559) FROM CAMPYLOBACTER LARI RM2100.

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Background

*Campylobacter* spp. are known to be one of a major cause of gastrointestinal illness throughout the world. Most prevalent species responsible for human infections are *C. jejuni* and *C. coli*. Human infections by *Campylobacter lari* are also associated with gastrointestinal diseases but mostly in developing countries. So far, little is known about *C. lari* molecular attributes. CLA0559 protein shows high homology to members of DsbA family. As Dsb enzymes control the formation and rearrangement of disulfide bonds during the folding of extracytoplasmic proteins they play an important role in pathogenesis. *In silico* analysis indicate that *C. lari* Dsb system might be a novel, different from the ones described for those operating in *E. coli* or *C. jejuni* cells.

Objectives

*C. lari* CLA0559 mutant remains motile but becomes sensitive to DTT. *E. coli* complementation tests (cadmium resistance, motility test and alkaline phosphatase assay) clearly showed that CLA0559 substitutes for EcDsbA activity and acts in cooperation with EcDsbB. CLA0559 functions *in vitro* as an oxidase. It what was proven by *in vitro* RNase activity assay. CLA0559 doesn’t cooperate with ClDsbI1/DsbI2, homologs of DsbB).

Methods

All genetic manipulations were performed using standard molecular biology procedures. For complementation purposes in *E. coli* cells ClDsbA was expressed from low copy vector. *In vitro* tests were performed with ClDsbA purified from *E. coli* using affinity and size exclusion chromatography

Conclusions

Our results showed that *cla0559* gene encodes for ClDsbA protein functioning in an oxidizing pathway in *C. lari* cells. Its redox partner still remains unknown.
IN VITRO PROPERTIES OF THE TRUNCATED FORM OF HELICOBACTER PYLORI HP0231 PROTEIN.

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Background
Dsb proteins control the formation and rearrangement of disulfide bonds during the folding of membrane and exported proteins. The mechanism of disulfide bond formation in microorganisms is extremely diverse. The H. pylori Dsb system seems to be novel and different from the Dsb system in E. coli. Dsb system plays an important role in H. pylori pathogenesis as H. pylori mutant impaired in disulfide bond formation revealed greatly reduced ability to colonize mice gastric mucosa.

Objectives
Our recent work led to the characterization of the HP0231 functioning mainly in an oxidizing pathway in H. pylori cells. It is intriguing that HP0231 acts as periplasmic oxidase, as EcDsbA, despite its structural resemblance to EcDsbG. Additionally HP0231 acts as chaperone and is involved in the cytochrome c biogenesis. To assess relations between HP0231 structure and its oxidizing activity we tested biochemical activities of the truncated form of HP0231 (HP0231m) containing only catalytic domain.

Methods
All genetic manipulations was performed using standard molecular biology procedures. Correctness of the obtained constructs were verified by sequencing. HP0231m was overexpressed in E. coli Rosetta strain and purified using NGC™ Medium-Pressure Chromatography (Bio-Rad). Purified protein was used to perform insulin reduction assay, chaperone activity assay and izomerization/oxidation RNase activity test.

Conclusions
Our results show that HP0231m mutated protein possess chaperone activity slightly higher than native HP0231 form and at the same time it lost ability to reduce insulin. It displays the same level of oxidation activity as HP0231 and similarly to native HP0231 does not have izomerization activity.
NOD1+32656 T>G FUNCTIONAL POLYMORPHISM AFFECTS SUSCEPTIBILITY TO CHLAMYDIA TRACHOMATIS (CT) INFECTION AND RISK OF TUBAL FACTOR INFERTILITY IN WOMEN

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Background

Genital CT infections cause tubal factor infertility (TFI) in some women. Bacterial motif recognition by intracellular pattern-recognition receptors NOD1 and NOD2 can trigger immune response. Whether their functional polymorphisms impact CT infection has not been researched.

Objectives

Our objective was to test the effects NOD1+32656 T>G and NOD2 1007fs on susceptibility and severity of CT infection in Dutch Caucasian women.

Methods

Susceptibility cohort: We selected 737 women visiting the Amsterdam STD outpatient clinic. Questionnaires were collected in regards to urogenital complaints.

Severity cohort: 490 Dutch Caucasian female patients visiting the UMC Groningen Fertility clinic were selected. Laparoscopy was used to grade the tubal pathology status as TFI grade 0-4. Controls were TFI grade 0 as assessed by either laparoscopy or hysterosalpingography (HSG).

Conclusions

The NOD1+32656 GG insertion appears protective against CT infection, but predisposes to TFI and a higher occurrence of symptoms in women with a past CT infection. The GG variant might be enhancing successful clearing while acting deleterious in the upper genital tract, causing tubal pathology.
Background
Atypical Enteropathogenic Escherichia coli (aEPEC) and Enterohemorrhagic E. coli (EHEC) are human pathogens that cause attaching and effacing lesions (A/E) in the intestinal mucosa. EHEC differs from aEPEC because it carries stx genes.

Objectives
Characterizing genotypic and phenotypic samples of aEPEC and EHEC.

Methods
E. coli was isolated from 130 healthy sheep and characterized by PCR technique as aEPEC (eae+/bfpA/stx-) and EHEC (eae+/stx+). 25 samples of aEPEC and 14 samples of EHEC were analyzed by PCR for presence of ehx, tirY-P (Tir phosphorylated), tirS (no phosphorylated), tccp e tccp2. PCR-triplex was used for phylogenetic classification with chuA, vjaA genes and fragments of TspE4.C2. Samples were also tested for sorbitol fermentation (SOR), enzymatic activity of β-D-glucuronidase (GUD), hemolysins production and HEp-2 cell adhesion.

Conclusions
The stx gene subtypes analysis by PCR and cytotoxic effect in Vero cells were realized only in EHEC. The ehx, tirY-P, tirS e tccp2 genes frequency was 40, 60, 16 and 16% in aEPEC and 78, 64, 7 and 28% in EHEC. The tccp gene was present only in aEPEC and only two aEPEC samples were SOR-/GUD+, being the another samples characterized as SOR+/GUD+. Enterohemolysin expression was analyzed in 32% of aEPEC and 66% of EHEC. A diversity patterns adhesion were observed. For the phylogeny, both patotypes were classified as belonging to B1 group. A 100% of EHEC samples presented cytotoxic effect in Vero cells, eight (57%) presented stx1c and seven (60%) stx2d. We concluded that aEPEC and EHEC, isolated from ovine, are two heterogenic groups of bacteria.
DOMINANT GENOTYPE OF STAPHYLOCOCCUS AUREUS IN A WELFARE CENTER
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Background
*Staphylococcus aureus* is associated with various diseases both in hospitals and in the community. The MRSA strains had manifested primarily in hospitals, but in recent years have appeared in community outbreaks in healthy people, so is called MRSA strains community-acquired (CA-MRSA), and now is important its detection.

Objectives
Identify MRSA strains in vulnerable adults admitted to a Welfare Center

Methods
Pharyngeal, nose and hands swabs were taken, of 100 volunteers in two samples of 50 people each; one in November 2012 and another in May 2014. The presence of *S. aureus* was determined by microbiological methods and biochemical tests. MIC for oxacillin was performed. Detection of genes for resistance to methicillin, *mecA* and Panton-Valentine Leukocidin, *lukS-PV / lukF-PV*, was made by PCR. The strains were typed by *spa*-typing and the clonal determination was performed by PFGE.

Conclusions
44 strains of *S. aureus* were isolated: 15 in throat, 13 in nose, and in 16 hands. Only two strains were resistant to methicillin. The dendrogram obtained from PFGE banding pattern shows that almost all strains are identical, which was corroborated by the *spa*-type where 83% of the strains tested showed the same *spa*-type t003. One dominant genotype of *S. aureus* in the population studied was found: *spa*-type t003. The MRSA clone of this type is dominant in Europe, but was not reported in Mexico in addition the type found here is methicillin-susceptible (MSSA). It could be a health problem in this community center.
DETECTION OF COMMUNITY-ACQUIRED METHICILLIN-RESISTANT STAPHYLOCOCCUS SPP IN ICU AND IN AREA HOSPITAL.
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Background
Nosocomial infections are an important cause of morbidity and mortality worldwide. At present, Staphylococcus aureus and Staphylococcus epidermidis are important etiologic agents of hospital infections. S. aureus is common in surgical areas of adult intensive care (ICU), sepsis being the most common process involved. Currently strains of community-acquired MRSA (CA-MRSA) have entered in the hospitals.

Objectives
The aim of this study was to detect the strains of community-acquired methicillin-resistant Staphylococcus in a General Hospital

Methods
Swabs of oral and nasal cavity of 52 patients, both male and female with different pathologies were taken, aged between 28 and 72 years, 14 patients were in the ICU and 38 the hospitalization area. The presence of S. epidermidis or S. aureus was determined by microbiological methods. MIC for oxacillin was measured. The presence of the mecA gene, the Panton-Valentine leukocidin gene, and the arginine catabolic mobile (ACME) gene, were detected by PCR. The strains were typed by spa-type.

Conclusions
We found that 100% of the samples analyzed in the area of hospitalization were strains of S. aureus, of which 37% were resistant to methicillin (MRSA). While 93% of the strains tested in ICU were S. epidermidis, and 31% of these were methicillin resistant (MRSE). All methicillin-resistant strains showed the presence of the mecA gene. One CA-MRSA strain and two CA-MRSE strains were found.
We detected the presence of community-acquired methicillin-resistant Staphylococcus in the hospital analyzed, so it is important to conduct periodic sampling to prevent outbreaks caused by this pathogen.
FEMS-2256
Bacterial pathogenicity

DETECTION OF COMMUNITY-ACQUIRED METHICILLIN-RESISTANT STAPHYLOCOCCUS AUREUS IN A NURSERY
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Background
Staphylococcus aureus is a microorganism of great medical importance. For many years it has been recognized as one of the major pathogens for humans. Infections with methicillin-resistant S. aureus (MRSA), are usually acquired in hospitals (strains HA-MRSA). However, in the late 90’s, MRSA strains emerged in healthy adults and children in communities. These strains cause infections in the community. The prevalence of these infections has increased significantly in recent years. Strains of S. aureus that cause these infections are called strains MRSA community-acquired (CA-MRSA).

Objectives
The aim of this work was to make the identification and molecular characterization of strains of Staphylococcus aureus isolated in a daycare of Mexico City.

Methods
Throat swabs samples from 87 children, between 2 and 6 years old were taken. S. aureus was identified by microbiological methods. The mecA and Panton-Valentine leukocidin (PVL) genes were detected by PCR. The SCCmec type and spa-type were determined.

Conclusions
25% of the population had S. aureus in the throat. From the strains isolated, we found that 22% (five strains) were MRSA, of which only one present the mecA gene, the gene of PVL and SCCmec type IV, therefore we found a CA-MRSA strain. The results show there may be strains CA-MRSA in healthy carriers in Mexico City, so strains CA-MRSA are present circulating in the community.
PRESENCE OF COMMUNITY-ACQUIRED METHICILLIN-RESISTANT STAPHYLOCCUS AUREUS IN AN ORPHANAGE
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Background
Staphylococcus aureus is a pathogenic bacterium that is associated with various diseases both in hospitals and in the community. Between 20% and 35% of the adult population are carriers of this microorganism in the nasal vestibule, but can also be found in the pharynx of people. The MRSA strains had manifested primarily in hospitals, but in recent years have appeared in community outbreaks in healthy children and adults, with no history of hospitalization. These strains were called community-acquired methicillin-resistant Staphylococcus aureus (CA-MRSA).

Objectives
The objective of this work was to detect the presence of strains CA-MRSA in an orphanage in Mexico City.

Methods
Pharyngeal and nasal swabs of 80 people between 3 and 19 years old of both genders were taken. We identified S. aureus by microbiological methods. The mecA and Panton-Valentine leukocidin (PVL) genes were detected by PCR. The SCCmec type and spa-type were determined.

Conclusions
56% of the population had S. aureus; 12.5% only in the pharynx, 17.5% only in the nose and 26% at both sites. We detected only four MRSA strains (5%), one in the pharynx and three in the nose, which presented the mecA, the PVL genes, and have the SCCmec type IV, so they are CA-MRSA strains. The results show that there are CA-MRSA strains in healthy children and adolescents in an orphanage in Mexico City.
Background

Citrobacter rodentium is a natural mouse bacterial pathogen that has been extensively used as a surrogate model for studying the human pathogens enteropathogenic and enterohemorrhagic Escherichia coli. During infection these pathogens employ surface structures called fimbria to adhere and colonize the host intestinal epithelium; however, for C. rodentium only a small number of its fimbrial operons have been studied.

Objectives

To analyze the regulation and role in colonization of the C. rodentium fimbrial operon gcfFGABCDE.

Methods

The role in colonization of the Gcf fimbriae was evaluated in mouse colonization assays, while the regulation of the gcf operon was assessed using transcriptional fusions and western blot. The promoter of gcf was identified by in silico analysis, primer extension assays and site-directed mutagenesis. Mutant strains and electrophoretic mobility shift assays were used to determine the role of H-NS as a repressor of gcf expression.

Conclusions

Here we report the characterization of Gcf (Gut colonization fimbriae) as an important C. rodentium colonization determinant of the mouse gastrointestinal tract. We demonstrated that the promoter of the gcf fimbrial operon is highly repressed under several in vitro growth conditions by H-NS1, one of the five H-NS paralogs encoded in the C. rodentium genome. H-NS binds to the regulatory region of gcf, further supporting its direct role as a repressor. The gcf operon possesses novel and interesting features that open future opportunities to expand our knowledge of these
essential bacterial structures during infection. Work supported by grants IN209713 (DGAPA) and 154287 (CONACyT).
FEMS-2245

Bacterial pathogenicity

SALMONELLA TYPHIMURIUM STRAINS WITH DISTINCT CLINICAL PHENOTYPES CAN BE DIFFERENTIATED AT THE TRANSCRIPTOMIC LEVEL

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Background

Salmonella Typhimurium infects a wide range of animal hosts, and generally causes a self-limiting gastroenteritis in humans. However, some variants of this serovar, sequence-type ST313, have been associated with an emergent invasive Salmonella disease in sub-Saharan Africa, which usually affects susceptible HIV+ or malarial individuals. A genomic comparison between an ST313 isolate, D23580, and the well-characterized 4/74 strain (sequence-type ST19) shows that both strains share 96% coding genes. Genetic differences include 1000 SNPs, D23580-specific prophages and the presence of pseudogenes.

Objectives

We tested the hypothesis that the two strains had different pathogenic mechanisms that are reflected by altered gene expression patterns and gene fitness in environmental conditions that reflect the infectious process.

Methods

RNA-seq-based transcriptomic data were obtained for strains 4/74 [1] and D23580 grown under seventeen infection-relevant in vitro conditions. Transcriptomic data were validated with a proteomic approach. In addition, a transposon library was generated in the D23580 strain and sequenced after passaging in different media.

Conclusions

Comparative transcriptomics between the two strains revealed that the proportion of differentially-expressed genes varied between 1% to 9% of all genes in various stress conditions. Proteomics for the early stationary phase condition confirmed 60% and 54% of the transcriptomic data in 4/74 and D23580, respectively. We are currently analysing the D23580 transposon library data. The differences observed in the expression and fitness of virulence-associated genes under specific environmental conditions may reflect altered regulatory mechanisms of these Salmonella strains.

ADHERENCE PROFILE OF ATYPICAL ENTROPATHOGENIC ESCHERICHIA COLI (AEPEC) STRAINS ISOLATED FROM DOGS IN BRAZIL
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Background
Atypical Enteropathogenic E. coli (aEPEC) is a human pathotype implicated in endemic infantile diarrhea. aEPEC with an identical virulence gene profile has been isolated from several animals including dogs, suggesting a zoonotic behavior. Adherence to host enterocytes is a key step of aEPEC virulence, being mediated through the expression of LEE locus genes, leading to formation of attaching-effacing (A/E) lesions. However, similar to LEE-positive Shiga Toxin-producing E. coli (STEC), aEPEC may contain distinct non LEE-associated effectors molecules and adhesins.

Objectives
Investigate the adhesion profile of aEPEC strains through search of non LEE-associated virulence markers and interaction assays with epithelial cells.

Methods
We analyzed by PCR the distribution of several virulence genes, including 16 nle genes, among aEPEC strains isolated from children (n=5) and dogs (n=5). Adherence phenotypes were determined through in vitro assays with HEp-2, Caco-2 and T84 cells.

Conclusions
The occurrence of virulence genes was: ecpA (100%), fimH (80%), lpfA (60%), hcpA (50%), toxB (30%), iha (10%). Fifteen nle genes were detected in all strains with different combinations. aEPEC of O51:H40 and O4:H16 serotypes carried the nleB and nleE genes, both located in PAI-122, an important pathogenicity island of STEC O157:H7. All strains showed a localized-like adherence phenotype (LLA) in HEp-2, Caco-2 and T84 cells. Although aEPEC strains are a very heterogeneous group, these findings shows a similar virulence profile between human and canine isolates belonging to the same serotype, suggesting an important role for dogs in the maintenance of the virulence gene pool of aEPEC.
Background
L-asparaginase II (AnsB) enzyme of E. coli, known for their use in treatment of children with acute lymphoblastic leukemia (ALL), and has been described as a bacterial periplasmic protein. However, we detected its presence in outer membrane proteins (OMP) extracts of different STEC serotypes, by Western blot using sera from STEC infected-patients and diagnosed with HUS. Recent reports using a mouse T cells co-cultured with wild type S. Typhimurium strain, showed that AnsB is sufficient to suppress T lymphocyte blastogenesis.

Objectives
Determine the AsnB presence in outer membrane protein extract of different STEC serotypes, its secretion and effect on T lymphocyte proliferation

Methods
The OMP and soluble proteins fractions were separated by SDS-PAGE 12% and western blot directed to AnsB detection was performed with commercial specific antibody anti-AnsB. A electron microscopy using a secondary antibody labeled with gold particles for visualization by immunogold was done. Peripheral blood mononuclear cells, from healthy individuals were incubated in the absence or presence of STEC O157: H7, O157:H7ΔansB, O157:H7ΔansB/pVB1_ansB, STEC O113:H21, commensal E. coli HS strain and S. Typhimurium, as a negative and positive control for AsnB effect, respectively. The suppression of proliferation was measured by flow cytometry. Study protocols for peripheral-blood mononuclear cells were approved by the Institutional Review Board of the Faculty of Medicine, University of Chile.

Conclusions
We observed by immunogold that the AsnB enzyme is secreted and some molecules remain associated with the outer membrane in STEC. AnsB expressed in O157: H7 contribute to its pathogenicity by inhibiting T lymphocyte proliferation.
Bacterial pathogenicity

INSIGHTS INTO THE AUTOTRANSPORT PROCESS OF TRIMERIC AUTOTRANSPORTER YADA.
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Background

YadA is a trimeric autotransporter adhesin present in Yersinia enterocolitica and Yersinia pseudotuberculosis. It has a transmembrane domain which forms a highly stable β-barrel anchoring it to the outer membrane and an N-terminal passenger domain which translocates through the β-barrel forming a coiled-coil stalk followed by a sticky head which mediates attachment of bacteria to host cell ECM components (1). The mechanism by which the N-terminal domain is translocated is still under debate. A relatively flexible region has been found near the C-terminus of the passenger domain known as the ASSA region which has been proposed to act as a hairpin during translocation (2).

Objectives

The idea is to mutate residues in the ‘ASSA’ region to obtain stalled translocation intermediates. The intermediates will be analyzed with various methods, followed by structural analysis using solid state NMR experiments.

Methods

Heat stability assays of the mutants indicated a different tertiary structure as well as less stability compared to wildtype. Infrared Spectroscopy indicated loss of α-helical content of mutant YadA by 33%. Electron Microscopy images showed that mutant YadA does not form fibre-like projections on cell surface and cells do not autoaggregate which was proved in Autoaggregation assays.

Conclusions

Preliminary results indicate a translocation intermediate which needs to be further analysed by ssNMR.

References

FUNCTIONAL GENETIC ANALYSIS OF THE ENVZ-OMPR TWO-COMPONENT SYSTEM IN ENTEROHEMORRHAGIC E. COLI PATHOGENICITY

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Background

Enterohemorrhagic Escherichia coli O157:H7 (EHEC) causes severe diarrhea, hemorrhagic colitis, and can also lead to life-threatening diseases like hemolytic uremic syndrome (HUS). Several EHEC virulence factors have been reported and encoded in the virulence plasmid pO157 and the O157 pathogenicity islands (O-Islands), including the locus of enterocyte effacement (LEE) island (148 O-Island encodes the type three secretion system and effector proteins) and the 93 and 45 O-Islands (encode the two Shiga-like toxins).

Objectives

In our previous studies, we utilized the model animal C. elegans for studying EHEC infection in vivo. Using this EHEC-C. elegans model to perform an EHEC transposon library screen, we found that mutations in the ompR gene confer the virulence-attenuated phenotype of EHEC against C. elegans animals. Here, we aimed to test whether the EnvZ-OmpR two-component system can regulate these reported EHEC virulence genes to infect C. elegans.

Methods

We identified several potential OmpR binding sites in the promoter regions of these EHEC virulence genes through in silico analysis. Furthermore, the mRNA transcript levels of these EHEC virulence genes were significantly down regulated in the ompR deletion mutant.

Conclusions

Taken all together, our current data suggested that OmpR is one of the EHEC virulence master regulators to manipulate the expression of virulence factors during infection in vivo.
FUNCTIONAL ANALYSIS OF BIOCONTROL BACTERIUM PSEUDOMONAS TAIWANENSIS

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Background
Pseudomonas taiwanensis is a board host range Gram-negative bacterium. Recently, we found that P. taiwanensis displayed strong antagonistic activity against rice pathogen Xanthomonas oryzae pv. oryzae (Xoo). Rice bacterial blight caused by Xoo is one of the most destructive diseases of rice worldwide.

Objectives
Understanding which genes are required by P. taiwanensis to resist the leaf blight of rice pathogen Xanthomonas oryzae pv. oryzae (Xoo)

Methods
Here we combined whole genome sequencing and Tn5-transposon mutagenesis to identify anti-Xoo toxin factors and related regulatory pathway. A high quality complete sequencing was accomplished by a combination of Roche 454, Illumina Solexa, Sanger sequencing and Optical mapping.

Conclusions
The complete sequence of the 5.08-Mb genome sequence and 4666 CDS were determined. We used transposon random insertion to identify genes involved in the production and regulation of anti-Xoo activity based on analyses of 6000 individual insertion-strains. Our results show that the siderophore pyoverdine biosynthetic gene (pvd), Type VI secretion system (T6SS), and EnvZ/OmpR two-component system have important roles in antagonistic activity against Xoo. On the other hand, the toxicity of P. taiwanensis was negatively regulated by the RpoS sigma factor. We further used MALDI-imaging mass spectrometry (MALDI-IMS) to track pyoverdine in P. taiwanensis and mutants. The results showed that pyoverdine was positively regulated by EnvZ/OmpR two-component system and secreted by T6SS. In contrast, pyoverdine was negatively regulated by RpoS. To the best of our knowledge, this is the first report that Pyoverdine has toxicity toward Xoo and T6SS can secrete small compounds.
Background

IL-6 is a key proinflammatory cytokine which has been considered to be important in the pathogenesis of periodontal disease. Host modulatory agents directed at inhibiting IL-6, therefore, appear to be beneficial in terms of attenuating periodontal disease progression and potentially improving disease susceptibility.

Objectives

In the current study, we investigated the effect of flavonoid isorhamnetin on the production of IL-6 in murine macrophages stimulated with LPS from P. intermedia, a pathogen implicated in inflammatory periodontal disease, and its mechanisms of action.

Methods

LPS from P. intermedia ATCC 25611 was isolated by using the standard hot phenol-water method. Culture supernatants were collected and assayed for IL-6. We used real-time PCR to quantify IL-6 and HO-1 mRNA expression. HO-1 protein expression and levels of signaling proteins were monitored by immunoblot analysis. DNA-binding activity of NF-κB was analysed by using the ELISA-based assay kits.

Conclusions

Isorhamnetin significantly down-regulated P. intermedia LPS-induced production of IL-6 as well as its mRNA expression in RAW264.7 cells. Isorhamnetin upregulated HO-1 expression at both gene transcription and translation levels in P. intermedia LPS-activated cells. In addition, inhibition of HO-1 activity by SnPP blocked the inhibitory effect of isorhamnetin on IL-6 production. Isorhamnetin failed to prevent LPS from activating either JNK or p38 pathways. Although further research is required to clarify the detailed mechanism of action, we propose that isorhamnetin may contribute to blockade of the host-destructive
processes mediated by IL-6, and could be a highly efficient modulator of host response in the treatment of inflammatory periodontal disease.
Background

*Klebsiella pneumoniae* is an opportunistic pathogen that affects immunocompromised patients and commonly associated with nosocomial infections. In the past two decades, the capsular serotype K1 has emerged as the predominant pathogen in causing liver abscess, affecting primarily Asians. While many studies have been reported for K1 and K2 which has comparable virulence to K1, very few of the 82 capsular serotypes have been studied.

Objectives

To determine the association between *K. pneumoniae* serotypes K1, K11, K19 and K31, and liver abscess and complications in kidney, spleen and lungs

Methods

Bacterial suspensions of the four serotypes of *K. pneumoniae* were inoculated intraperitoneally into healthy mice models. After 72 hours, the organs were aseptically removed, and histopathological examination and bacterial enumeration were carried out. The presence of liver abscess and complications in the other organs were characterized by the observation of necrosis and severity of inflammation, respectively.

Conclusions

From the histopathology examination, serotypes K1, K19 and K31 resulted in liver abscess, unlike K11. The spleen was severely inflamed with K1 and mildly inflamed with the other three serotypes. Moderate inflammation was observed in the lungs infected with K1 while the other serotypes caused mild inflammation. The immune system of the mice was able to clear the bacteria in all the organs within 72 hours when infected with all four serotypes. Our preliminary data supports a statistical association between non-K1 *K. pneumoniae* serotypes and liver abscess and organ complications in healthy mice models.
THE EFFECTOR PROTEIN SDHA FROM PISCIRICKETTSIA SALMONIS IS OVEREXPRESSED DURING AN INFECTIOUS PROCESS IN CELL LINE SHK-1.

Background

_Piscirickettsia salmonis_ is the etiological agent of Piscirickettsiosis disease in Chilean Salmon farming and cause significant economic losses in this industry. For similar pathogens like _Legionella pneumophila_ and _Coxiella burnetii_ some effector proteins were described that modify cellular processes of the host to establish intracellular infection. Nowadays there is no evidence for similar effector proteins of _P. salmonis_. Only one report described a type 4b secretion system (SST4b) in this pathogen.

Objectives

Evaluate gene expression levels of _dotH_, _dotG_ and _sdhA_ genes during an infectious process induced by _Piscirickettsia salmonis_ on SHK-1 cell line.

Methods

RNA extraction was performed combining Trizol with commercial kit for total RNA. RT reaction was performed with M-MLV and Real Time PCR was performed by commercial SYBR Green master mix kit.

Conclusions

The expression levels of _dotG_, _dotH_ and _sdhA_ genes are increased in a correlated manner between them during an infectious process induced by _Piscirickettsia salmonis_. At early times such 2 hours after infection the expression levels are increased and then decrease a few fold-change to increase again at the final days of infection but not as much as in the beginning of the experiment. From this pattern of gene expression we propose that the SST4b is involved in traslocating the effector protein SdhA to the cytoplasm of the infected cell.

Funds

Conicyt Scholarship GO 21110631; FONDAP project 15110027.
Background
Leptospirosis is a zoonosis with global distribution recognized as a re-emergent disease caused by pathogenic bacteria of the genus *Leptospira*. LipL21 is major leptospiral surface protein identified by proteomics.

Objectives
The goal of this work is to obtain the recombinant protein LipL21 expressed in *E. coli* and to evaluate its binding activity with extracellular matrix components (ECM), integrins cell receptors (αLβ2, αMβ2) and C3b and C4b complement factors.

Methods
The gene LIC10011 encoding for LipL21 was cloned and expressed in *Escherichia coli*. Protein attachment to individual ECM components, αLβ2, αMβ2 integrins, and C3b and C4b was screened by ELISA, and the binding was evaluated by probing the reaction with anti-LipL21 serum. Binding of LipL21 recombinant protein to ECM and integrins was compared with their binding to gelatin, BSA and fetuin negative controls, by using Student's two-tailed t-test.

Conclusions
LipL21 was expressed in *E. coli* with a 6X HIS sequence tag at N-terminal. The protein was present in its soluble form, and was successfully purified in Ni²⁺-charged resin, as assessed by SDS-PAGE. Binding of LipL21 was statistically significant with cellular fibronectin, collagen IV, laminin and αLβ2, αMβ2 integrins, which may be justified by the occurrence of *DGEA* integrin binding domain in LipL21 protein. The binding of LipL21 with C3b and C4b was significantly enhanced when the protein was associated with αMβ2 integrin, probably due the presence of binding sites for these components in the ligand. These results suggest that this protein has the potential to cooperate in the bacterial immune evasion and dissemination in the hosts.
CHARACTERIZATION OF VIRULENCE FACTORS IN ESCHERICHIA COLI ISOLATES FROM HAEMOCULTURES

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Background

Extraintestinal pathogenic Escherichia coli (ExPEC) is one of the main etiological agents of Gram-negative bloodstream infections. Capacity of ExPEC to cause systemic infections is due to genes encoding virulence factors, which include adhesins, toxins, polysaccharide coatings, siderophores or iron acquisition systems.

Objectives

The aim of our study was to analyse the presence of virulence genes fimA, bfpA, pap, sfa, eae, afa, cnf1, α-hly, aer, iai, Lt, st, ipaH, iucC, cdtl-V and to determine the phylogenetic origin of E. coli strains isolated from septicemic patients.

Methods

This study included 133 E. coli strains isolated by haemoculture from adult patients with bacteriemia. Automated blood culture system (Bactec 9050) was used in haemocultivation. E. coli strains of positive haemoculture were identified on a Microflex MALDI Biotyper (Bruker Daltonik) and confirmed by standard biochemical methods ENTEROtest 16, ENTERO-Rapid 24 (Lachema, Czech Republic). Polymerase chain reaction was used to detect virulence genes and to determine ECOR phylogenetic classification of E. coli strains.

Conclusions

We found that fim A (96%), aer (75%) and iucC (68%) represent most common genes encoding virulence properties. Toxin studies revealed a relatively high incidence of cnf1 (20%) and α-hly (19%). Phylogenetic classification showed that 80% of E. coli strains tested fall into phylogenetic groups B2 and D, representing main groups of virulent E. coli.
FEMS-2019

Bacterial pathogenicity

GENOMIC ANALYSIS TO FIND NOVEL USHER-CHAPERONE ASSEMBLED FIMBRIAE IN ENTEROTOXIGENIC ESCHERICHIA COLI STRAINS

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Background

Enterotoxigenic Escherichia coli is leading cause of diarrhea, mainly in children and travelers to endemic regions. Currently, there is no effective vaccine to prevent ETEC-caused diarrhea. ETEC colonizes the small intestine by using a diverse set of adhesins including about twenty different pili. However, between 15-50% of the isolates obtained from diarrhea cases worldwide are negative in detection of the known adhesins, suggesting presence of additional unidentified adherence determinants and, at the same time, presenting an obstacle in the development of adhesin-based vaccine candidates.

Objectives
To identify novel fimbrial loci in ETEC strains negative for detection of the known adhesins

Methods

Genomes of 35 ETEC strains isolated from diarrhea cases in Chile (14 isolates), Kenya (5), Mozambique (2), The Gambia (1), Mali (3), Bangladesh (3), India (2) and Pakistan (5), which were negative in detection of 23 adhesin genes, were partially sequenced by a paired-end protocol (Illumina, HiSeq 2000). Genomic sequences were screened for 161 fimbrial usher genes belonging to nine families of fimbriae (alpha, beta, gamma-1, gamma-2, gamma-3, gamma-4, kappa, sigma and pi), in order to locate homologous genes and find putative novel loci encoding usher/chaperone assembled adhesins. Genomic sequences of non-pathogenic E.
coli strains (13) and other ETEC strains (H10407, E24377A and B7A) were included as controls.

Conclusions
Loci encoding putative novel gamma-2 fimbriae were found in 23 strains, being frequent among ETEC strains negative for detection of known adhesins.
FEMS-1018
Bacterial pathogenicity

COORDINATED REGULATION OF THE MAJOR VIRULENCE GENE PELD OF THE PHYTOPATHOGEN BACTERIUM Dickeya dadantii BY FIS, CRP AND H-NS

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Background
Pathogenic bacteria tightly regulate their virulence genes in response to the variations of their environment. Many transcription factors have been identified for such genes, but their interactions remain poorly understood. The pel genes of the phytopathogen bacterium D. dadantii are a good example of this situation, with a dozen identified regulators, yet an integrative view of their action was lacking.

Objectives
As a first step towards an integrative model, this work aims at taking into account the interactions between the major global regulators of virulence (Fis, CRP and H-NS) in the regulation of the pelD virulence gene. In particular, Fis has 2 sites overlapping the promoter (-34 and -10 from the transcription start site) and an upstream site (-126).

The relationships between those sites are investigated here as well.

Methods
Binding sites were inactivated individually by site-directed mutagenesis and the impact of the modifications was evaluated both in vivo by gfp gene fusion measurements, and in vitro by quantitative DNase I, potassium permanganate footprinting and in vitro transcription.

Conclusions
D. dadantii has established 3 redundant mechanisms to ensure a full repression of pelD in exponential phase: 2 of the Fis binding sites (-34 and -10) directly compete with RNA polymerase, 1 Fis site (-34) prevents the activator CRP from binding and a Fis-regulated (involving the -126 Fis site) reverse promoter represses the pelD promoter. On top of this, H-NS modulates the action of Fis. These results show that the importance of interactions in transcriptional regulation should not be underestimated.
FEMS-1316
Bacterial pathogenicity

YQIC IS REQUIRED FOR COLONIZATION, MOTILITY, AND FLAGELLATION OF SALMONELLA TYPHIMURIUM, AND INTERLEUKIN-8 AND HUMAN BETA-DEFENSIN-3 PRODUCTION IN HUMAN INTESTINAL EPITHELIAL CELLS
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Background

The phenotype of Salmonella yqiC gene and its impacts on host responses are little known.

Objectives

To investigate whether yqiC is responsible for Salmonella colonization/invasion, bacterial motility/adherence, and interleukin (IL)-8/human β-defensin-3 (hBD3) production in human intestinal epithelium.
Methods
HeLa, Caco-2, LS174T, and THP-1 cells were infected with *Salmonella Typhimurium*
wild-type SL1344, yqiC-depleted mutant ΔyqiC, its complemented strain ΔyqiC', and filC-deleted mutant ΔfilC (MOI=5) for 2 hours, and treated with plain medium (output pool A: colonizing bacteria) or gentamicin (output pool B: invading bacteria) for 1 hour. At these times cell-associated bacterial numbers were calculated. Next, motility and adherence of these strains were examined by soft agar motility assays and yeast agglutination tests. Last, LS174T cells were treated with these strains, flagellin, and IL-1β for 2 hours, incubated in gentamicin for 1 hour, and the infections were continued for 15 hours. At the end-point, the supernatants were collected to measure IL-8 and hBD3 using ELISA.

**Conclusions**

Compared with SL1344, ΔyqiC was attenuated in bacterial colonization and invasion in 4 cells (Fig.1), lost swimming ability (Fig.2A), and constitutively expressed type 1 fimbriae (Fig.2B). IL-8 production in ΔyqiC-infected cells was lower than that in SL1344-infected cells, with lower levels than those in ΔfilC-infected cells (Fig.3A). Furthermore, ΔyqiC and ΔfilC induced lower hBD3 secretion than SL1344 did in LS174T cells (Fig.3B).

In conclusion, yqiC is required for *Salmonella* Typhimurium colonization, and host IL-8 and hBD3 production. This is mediated by its modulation in *Salmonella* motility and flagellation via downregulation of type 1 fimbriae.
ROLE OF FIBRONECTIN IN THE ADHERENCE OF DIARRHEAGENIC E. COLI TO INTESTINAL CELLS AND THE INDUCTION OF SECRETION OF PRO-INFLAMMATORY CYTOKINES
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Background
Fibronectin has been shown to be important for the adherence of diarrheagenic E. coli (DEC), but the biological relevance of the DEC-fibronectin-mediated interaction to intestinal cells remains unclear

Objectives
We evaluated the impact of the DEC-fibronectin-mediated interaction on the induction of secretion of pro-inflammatory cytokines

Methods
We assessed the ability of EAEC, ETEC and EHEC to bind to HEp-2 cells in the presence or absence of purified fibronectin protein and we quantified the amount of IL-8 secreted by infected cells. Additionally, we quantified the expression of FOS, NFkB, IL8, CCL20, IL1A and TNFalpha genes. Alternatively, we quantified the IL-8 secretion of HEp-2 cells transfected with small hairpin RNA (shRNA) for fibronectin or scrambled shRNA infected with EAEC strain.

Conclusions
We found a significant increase in the adherence to HEp-2 cells pre-incubated with fibronectin compared with cells not incubated with this protein for all DEC assayed. However, the IL-8 secretion was significantly reduced in the presence of fibronectin. Real-time PCR assay indicated that of all genes assayed only IL8 gene expression was reduced in HEp-2 cells pre-incubated with fibronectin. Experiments using fibronectin shRNA revealed a decrease in the EAEC adherence to cells, but no changes in the IL-8 secretion.

Overall, all data suggested that fibronectin participates exclusively in the adherence of DEC to epithelial cells, but not in the inflammation originated as a consequence of the infection.
Bacterial pathogenicity

AROA DEFICIENT SALMONELLA – MORE THAN AN AUXOTROPHIC MUTANT?

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Background
Attenuated Salmonella employed as live vaccine carrier could provide protection against many infectious diseases. To guarantee successful vaccination, balance between attenuation and immunestimulation is required. Since aromatic amino acids are not freely available in the host, ΔaroA is commonly used as an attenuating factor. However, we observed that an aroA deficiency also dramatically modified the phenotype of Salmonella by increasing bacterial virulence.

Objectives
In the present work the molecular basis for the phenotypic changes in the strains should be determined by transcriptional profiling and gene deletion of apparently relevant genes.

Methods
We were able to demonstrate via transcriptome profiling that Salmonella is consuming excessive pyruvate by up-regulating sugar pathways like manXYZ or glpABCQT resulting most likely in an osmotic imbalance. Furthermore, aroA deficient mutants are more prone to express FljB flagella. These metabolic dysregulations could be a possible explanation for the observed phenotype of increased virulence. In accordance after intravenous application of ΔaroA Salmonella to mice (86/609/EEC), significantly increased induction of pro-inflammatory cytokines like TNF-α or IFN-β was observed emphasizing the importance of this deletion for Salmonella’s phenotype. We also employed such bacteria in bacteria-mediated tumor therapy. We could show that the ΔaroA strains display an increased tumor colonization and an improved anti-tumor response.

Conclusions
Disturbing the bacterial metabolism by mutating aroA in Salmonella leads to a modified phenotype that results on one site in attenuation but also improves Salmonella’s ability to induce an immune response. Therefore this mutation might be a preferable modification for vaccine and cancer research.
THE CAPSULE OF CAMPYLOBACTER JEJUNI PREVENTS BINDING TO SIGLEC-1 AND SIGLEC-7

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Background
Campylobacter jejuni (C. jejuni) is the leading cause of bacterial gastroenteritis worldwide. Besides diarrhoea, an infection with C. jejuni can precede the acute and severe polyneuropathy Guillain-Barré syndrome (GBS). Lipooligosaccharides (LOS), present in the outer membrane of C. jejuni, cause activation of the innate immune system and sialylation of LOS is critical for the induction of GBS. Sialylated LOS can be recognised by two host immune-receptors, Siglec-1 and Siglec-7. Recent research indicates that C. jejuni LOS are not always exposed to the external environment. We hypothesize that C. jejuni exploits the polysaccharide capsule as an evasion strategy to prevent immune recognition mediated by Siglec-1 and Siglec-7.

Objectives
To determine whether the polysaccharide capsule of C. jejuni prevents the specific binding of bacterial LOS to Siglec-1 and Siglec-7.

Methods
Capsule and sialic acid transferase knock-out mutants were generated. Bacteria were FITC-labelled, incubated with Siglec-1 or Siglec-7 transfected cells and binding of live or heat-inactivated bacteria was measured using flow cytometry.

Conclusions
Absence of the capsule of C. jejuni enhanced binding of C. jejuni sialylated LOS to Siglecs. For Siglec-1, this was observed for both live and heat-inactivated bacteria. In contrast, for Siglec-7 the difference in binding between wild-type and non-capsulated C. jejuni was only observed for live bacteria. Sialic acid mutants showed lower binding to Siglecs. Our results give insight in how the immune system recognizes C. jejuni and define the capsule as a bacterial factor that may influence the onset of GBS.
Bacterial pathogenicity

CTRA-DEPENDENT BRUCELLA ABORTUS CELL CYCLE REGULATION IN CULTURE AND DURING INFECTION
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Background
Brucella abortus is a facultative intracellular pathogen for mammals. Many aspects of its intracellular trafficking have been investigated in HeLa cells and RAW264.7 macrophages. However very little is known about its cell cycle regulation, except that bacteria are blocked in G1 phase during early hours of infection.

Objectives
The main goal of this project is to investigate the regulon of CtrA, a central transcription factor. We also decided to investigate a two-component system (TCS) located upstream of a signalling pathway regulating CtrA phosphorylation status. This TCS involves a histidine kinase, PdhS, and a single-domain response regulator, DivK (Hallez).

Methods
We used RSA-Tools (Van Helden) to look for the consensus sequence bound by CtrA in B. abortus genome. This in silico approach allowed the prediction of a long list of putative CtrA targets. A ChIP-seq analysis confirmed part of these targets. The activity of some of these promoters was monitored by fusing them to a gene coding for an unstable GFP.
We performed in vitro kinase assays to test PdhS autophosphorylation and its ability to transfert the phosphate group to DivK.

Conclusions
Promoters bound by CtrA are predicted to control genes involved in cell cycle regulation, such as division, chromosome replication and segregation. The reporter system showed that the activity of some promoters varies in bacteria grown in rich culture medium according to their cell size, as well as during B. abortus intracellular trafficking in HeLa cells.
In vitro kinase assays show that PdhS can autophosphorylate and quickly transfer its phosphate group to DivK.
THE DUAL BEHAVIOUR OF CARBON STARVATION GENES PROVIDES A LINK BETWEEN METABOLISM AND PATHOGENESIS OF SALMONELLA

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Background

The ability of the enteric pathogen Salmonella to utilize a diverse set of nutrients, allows it to survive in various nutrient limiting conditions, rendering it one of the most successful pathogens. Salmonella employs sophisticated machinery to either acquire a specific nutrient from the surrounding environment or modify the host’s ecosystem for its own metabolic benefit. Carbon starvation (cst) genes represent one such class that are proposed to be expressed during carbon starvation, the most common stress encountered by Salmonella. Two cst genes, cstA and yjiY, predicted to mediate peptide utilization, were previously reported to be regulated by the global regulator CsrA and YehU/YehT two-component system in E. coli.

Objectives

In Salmonella, cstA was shown to be required for virulence in C. elegans, however, the underlying mechanism remains unknown. The objectives of this study are to establish the role of cst genes in metabolism and to understand how they affect Salmonella pathogenesis.

Methods

We generated knockout for the genes cstA and yjiY in Salmonella Typhimurium. Phenotype microarray on various nutrient sources along with transcriptome analysis was carried out for wild-type and knockout strains. Virulence was assessed using cell culture and animal models.

Conclusions

Phenotype microarray confirmed the importance of cst genes in metabolism of Salmonella. Despite the high similarity in their sequences, cstA and yjiY displayed different effects on motility, adhesion, biofilm forming ability and virulence in the animal model of infection. This study brings forward pleiotropic role of cst genes and highlights the crucial cross-talk between metabolism and pathogenesis of Salmonella.
TRANSCRIPTOMIC ANALYSIS IN SALMONELLA ENTERICA REVEALS PRODUCTS OF THE STD FIMBRIAL OPERON AS GLOBAL REGULATORS OF GENE EXPRESSION

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Background
Fimbriae are virulence factors in Salmonella enterica, playing diverse roles in bacterial adhesion (Humphries et al. FEMS microbiology letters, 201:121-125, 2001). Synthesis of Std fimbriae, that are not expressed under laboratory conditions, has been shown to occur in the animal intestine, but the molecular mechanisms involved remain unknown (Weening et al. Infection and immunity, 73:3358-3366, 2005). It was previously described that derepression of the std operon occurs in DNA adenine methylase (dam) mutants and that HdfR, a poorly known LysR-like factor, is necessary for std transcription. However, HdfR-mediated activation of std transcription occurs only in a fraction of the bacterial population, suggesting the occurrence of either bistable expression or phase variation (Jakomin et al. Journal of bacteriology, 190:7406-7413, 2008).

Objectives
This work aims to further understand the molecular mechanisms that might allow expression of the std operon inside the animal and to analyze the effect of its expression in S.enterica.

Methods
A genetic screen for additional std regulators revealed that the std operon itself is a positive regulator, and the downstream genes of the operon, stdE and stdF, are crucial for this control. Surprisingly, transcriptomic analysis revealed that StdE and StdF control a plethora of S. enterica loci including genes involved in motility, chemotaxis, biofilm formation, conjugation, and virulence.

Conclusions
StdEF-mediated control may play a role in the crosstalk between motility, adhesion and invasion, considered essential for optimal Salmonella infection. If StdEF expression is subjected to phase variation during infection, subpopulation formation can be expected to occur in the animal intestine.
Background

Despite continuous efforts to reduce its incidence, mastitis remains a major disease in dairy cows. Main mastitis pathogens include Streptococcus uberis, Staphylococcus uberis and Escherichia coli. Induction of the innate immune response is a key mechanism in the initiation of the host response during infections of the mammary gland in cows by Escherichia coli.

E. coli P4 is a prototypical mastitis strain isolated from a case of clinical mastitis. Previous studied showed that E. coli P4 was virulent in a mouse model of mastitis. Yet, we have found that this strain induced a reduced pro-inflammatory response in mammary epithelial cells (MEC) compared to other E. coli isolates.

Objectives

Our objective was to identify genes responsible for the low pro-inflammatory response induced in MEC by E. coli P4 and determine if this reduced pro-inflammatory response contributed to the virulence of E. coli P4 in a murine mastitis model.

Methods

By screening a library of 2000 mutants for increased pro-inflammatory response on MEC, we identified a series of mutations that increase the response of MEC to strain P4. The impact of these mutations on the phenotype of E. coli P4 will be presented in details and discussed.

Conclusions

Altogether, results presented will allow a better understanding of the initial steps of the interaction between mammary epithelial cells and mastitis pathogens. This work will help understand if the low pro-inflammatory response triggered by E. coli P4 contributes to its virulence.
Background
Recurrent urinary tract infection (rUTI), most often caused by uropathogenic Escherichia coli (UPEC), is a life altering, potentially life threatening problem for many women. Asymptomatic carriage of UPEC in bladder reservoirs has long been believed to be a source for rUTI. However, the natural triggers of UPEC emergence from these reservoirs have remained mysterious. Women with bacterial vaginosis (BV), an imbalance of the vaginal microbiota, are at increased risk for UTI.

Objectives
We sought to address the long-standing idea that UPEC bladder reservoirs seed rUTI and to understand why women with BV are more prone to UTI. Translocation of BV bacteria to the urinary tract is likely to occur during sexual activity, which is often a prelude to rUTI. We tested the hypothesis that bladder exposure to BV-associated bacteria may initiate UPEC emergence from intracellular reservoirs.

Methods
Mice containing UPEC bladder reservoirs were challenged by transurethral inoculation with either PBS (control) or the BV-associated bacterium Gardnerella vaginalis. We monitored the consequences of bladder exposure to G. vaginalis by 1) enumerating urinary tract bacterial titers, 2) assessing epithelial exfoliation by immunofluorescence and scanning-electron microscopy and 3) RNAseq analysis.

Conclusions
Here we show that G. vaginalis triggers bladder epithelial exfoliation and emergence of E. coli from intracellular reservoirs. We observed phenotypes consistent with rUTI, including high UPEC titers and neutrophils in urine. These data strongly suggest that G. vaginalis presence within the urogenital microbiota may be a trigger for rUTI and likely contributes to the substantially increased risk of UTI in women with BV.
Background: The gene expression of microorganisms and environmental factors are closely related.

Objectives: To investigate possible changes in the gene expression levels of virulence genes [sfa/foc (S and F1C fimbriae), cnf 1 (cytotoxic necrotizing factor), and usp (uropathogenic-specific protein)] in a uropathogenic E. coli standard strain (UPEC C7) in the presence of insulin and glucose.

Methods: The UPEC strain was cultured in broth media (tryptic soya broth-TSB) under 5 different conditions: 0.1% glucose, 20µU insulin, 200µU insulin, 0.1 % glucose+200µU insulin, and control TSB. After incubation for 24 hours at 37°C, total RNA isolations were prepared from all five bacteria cultures. Gene expression levels were determined by quantitative PCR. Changes in gene expression were evaluated using the expression levels of 16sRNA as a “housekeeping gene,” according to the Pfaffl equation.

Conclusions: The medium containing 0.1 % glucose+200µU insulin had increased expression levels of all three genes in the E. coli C7 strain. In contrast, the other media (0.1% glucose, 20µU insulin, 200µU insulin) had a decrease in the expression levels of virulence genes, all compared to control medium.

This investigation shows that some environmental factors, such as glucose and insulin, could determine the pathogenicity of E. coli strains.
Bacterial pathogenicity

A THERMO-REGULATED PROLINE-RICH SURFACE PROTEIN OF ENTEROCOCCUS FAECIUM BINDS TO FIBRINOGEN, FIBRONECTIN AND PLATELETS AND CONTRIBUTES TO BIOFILM FORMATION

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Background

Enterococcus faecium is a commensal of the mammalian gastrointestinal tract. Recently, it has become an important nosocomial pathogen, causing infections that are difficult to treat. However, little is known about the mechanisms that E. faecium employs to colonize and infect mammals.

Objectives

We aimed to clarify the interactions of E. faecium with the host and we hypothesized that genes important for colonization and infection, could exhibit temperature-regulated expression control.

Methods

We performed a transcriptome analysis of E. faecium E1162, during mid-exponential growth at 25°C and 37°C. Furthermore, we functionally characterized a surface protein of E. faecium, that was found to be produced at higher levels during growth at 37°C than at 25°C.

Conclusions

We found thirty-three genes expressed significantly higher at 37°C compared to 25°C. One of the most highly upregulated genes (4.4-fold), is predicted to encode a peptidoglycan-anchored surface protein. The N-terminal domain of this protein is unique to E. faecium and closely related enterococci. The C-terminal domain contains
three proline-rich repeats, leading us to name the protein PrpA for proline-rich protein A. PrpA is a surface-exposed protein, that was maximally produced in exponentially growing cells at 37°C. PrpA is immunogenic as specific antibodies were observed in patients after they suffered from an *E. faecium* bacteremia. Heterologously expressed and purified PrpA was able to bind to fibrinogen, fibronectin and to platelets. Furthermore, a *prpA* deletion mutant was defective in the early stages of biofilm formation. Our data indicate that PrpA may contribute to the pathogenesis of *E. faecium* infections in hospitalized patients.
THE DENUDATION OF LATERAL PART PROMOTE CAMPYLOBACTER JEJUNI INVASION IN CACO-2 CELLS.

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Background
The intestinal epithelium provides a physical barrier to luminal bacteria. This barrier serves as the first line of defense against bacterial adhesion and invasion. Campylobacter jejuni is a Gram-negative commensal bacterium in domestic animal and cause gastrointestinal foodborne disease in human. C. jejuni is able to adhere and invade gut epithelium, which lead cause of acute gastroenteritis in human. It has believed that the function of tight junctions (TJs) in human intestinal epithelium is closely related with invasion of C. jejuni.

Objectives
The aim of this study was to investigate interaction between C. jejuni and host intestinal barrier functions. In this study, we used cultured islands (unpolarized) and sheets (polarized) form Caco-2cell.

Methods
Invasion of C. jejuni was measured by gentamycin protection assay in Caco-2 cells. Confocal microscopy was used to examine the localization of internalized bacteria. And the most of bacteria were observed on cellular lateral part in islands of cell. Interestingly, C. jejuni efficiently invaded from basolateral surface compared with apical surface and intracellular C. jejuni was significantly increased in disruption of TJs by the treatment with EGTA. Furthermore, C. jejuni infection induced the distribution of the TJs component.

Conclusions
These date indicated that the TJs formed by each of neighboring cells might act as physical barrier for prevention of C. jejuni invasion, and infection of C. jejuni might effect on formation of TJs component. These results suggested that the C. jejuni infection inducible TJs disruption might enhance C. jejuni invasion into host cell.
CAPNOCYTOPHAGA CANIMORSUS IS ADHERING TO HOST MEMBRANE GLYCOPROTEINS

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Background

*Capnocytophaga canimorsus* (Cc), member of the *Bacteroidetes* phylum, is an oral commensal from dogs which causes rare but severe infections in humans. *Cc* feeds on sugar moieties of glycoproteins expressed on host cell membranes. Removal of N-linked glycans is achieved by the glycoprotein deglycosylation (Gpd) complex encoded by a polysaccharide utilization locus, a hallmark of *Bacteroidetes*.

Objectives

The aim of this study is to determine whether *Cc* is adhering to the host cells it is feeding on and, if so, which mechanisms are implicated.

Methods

Cell lines were infected with strain *Cc5*. Mutant bacteria were constructed by site-directed gene replacements. Adhesion was monitored by microscopy and flow cytometry.

Conclusions

*Cc5* adheres to epithelial, endothelial and immune cells. In *Flavobacterium johnsoniae*, another member of the *Bacteroidetes* phylum, gliding motility and adhesion are associated. In *Cc5* adhesion was not abrogated in a mutant unable to glide. Adhesion of *Cc5* was mainly mediated by the Gpd complex. *Cc5* adhered to host cells via the membrane glycoproteins as inhibition of protein glycosylation by tunicamycin led to a decreased adhesion. Removal of serum from the assay, leaving cell glycoproteins as the only feeding source, increased adhesion. If fetuin, a glycoprotein deglycosylated by *Cc5*, was added to the serum-depleted medium, adhesion decreased. We conclude that adhesion of *Cc5* to cells is linked to feeding through the Gpd complex. Further work will be conducted to find out whether adhesion could be important during infection.
Background

Members of the *Mycobacterium tuberculosis* complex are the causative agents of tuberculosis, a major global health threat. The majority of individuals are thought to be infected latently whereby the bacteria are in a non-replicating dormant state, and the individual is asymptomatic. Many of the proteins secreted by this organism have been shown to be essential for the virulence, survival and dormancy, and function in a variety of different roles. The resuscitation promoting factors (Rpfs) are one such family of proteins whose peptidoglycan hydrolyzing activities have been correlated with the resuscitation of dormant bacteria; however, the precise molecular mechanisms underlying this process remain poorly understood.

Objectives

The aim of this work is to gain insight into the expression and localisation patterns of Rpfs during bacterial growth and macrophage infection.

Methods

We have generated a number of Rpf deletion mutants and Rpf specific polyclonal antibodies to monitor Rpf expression *in vitro* using flow cytometry, and during *Mycobacterium marinum* infection of murine macrophages by immunofluorescence and confocal microscopy.

Conclusions

Having observed Rpfs at the level of individual bacteria, we conclude that a subset of...
the mycobacterial population express Rpf at a detectable level during *in vitro* growth, and the infection process.
DISTRIBUTION OF GENES RESPONSIBLE FOR VIRULENCE IN CRONOBACTER STRAINS

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Background

Cronobacter spp. was proposed as a new genus in 2008. All the members of Cronobacter are considered to be potentially pathogenic; however recent epidemiological studies indicated differences in virulence. The most often isolated species from clinical samples were C. sakazakii and C. malonaticus. Some reports revealed the connection between sequence type and virulence of particular strain. It has been shown that sequence type 4 (ST4 – C. sakazakii) is prevalent for meningitides, ST7 (C. malonaticus) is often associated with adult infections. These sequence types are considered as the evidence of virulence.

Objectives

The objectives of this work were to implement PCR specific for virulent proteins and to reveal connection between sequence type (MLST) and virulence genes.

Methods

Several factors responsible for virulence were targeted: the system for iron acquisition, outer membrane protein A, haemolysin and metalloprotease placed on chromosome and type six secretion system, secretion system FHA and outer membrane protease placed on plasmids.

PCR protocols for the detection of corresponding genes were applied to all Cronobacter species and compared with determined sequence type.

Conclusions

The predominance of some virulence genes was observed within particular species, in some cases even with sequence type; for example, haemolysin was detected in all tested C. sakazakii strains and in C. malonaticus presenting sequence type 7.

This work was supported by the Czech Grant Agency (13-23509S).
Background

Haemophilus influenzae (Hinf) is a Gram-negative pathogen colonizing the upper respiratory tract mucosa. Hinf belongs to a group of human-restricted bacteria, which bind to carcinoembryonic antigen related cell adhesion molecules (CEACAMs) on epithelial cells. Adhesion to CEACAMs is thought to be mediated by the Hinf outer membrane protein (OMP) P5 promoting establishment of the pathogens in the human nasopharynx.

Objectives

Aiming at preventing Hinf colonization, we sought to identify the molecular requirements for Hinf binding to CEACAMs.

Methods

Binding assays with soluble receptor ectodomains, followed by flow cytometric analysis or Western Blotting, were used to characterize CEACAM-binding profiles of wildtype and mutant Hinf. OMP P1, and not OMP P5, was identified as the CEACAM-binding adhesin. Multiple amino acid sequence alignment of P1 combined to heterologous expression of wild-type, chimeric, or mutated P1 in E. coli depicted the molecular details of the P1-CEACAM interaction.

Conclusions

Surprisingly, Hinf P5 mutants still avidly bind CEACAMs and Hinf P5 expressed in E. coli fails to mediate CEACAM targeting. Instead, a genetic screen identifies Hinf P1 as the CEACAM-binding adhesin. Deletion of P1 in Hinf and heterologous expression in E. coli demonstrate that P1 is necessary and sufficient to bind several human CEACAMs. Concordantly, when expressed on the surface of E. coli, P1 promotes adhesion to and invasion into epithelial cells. Structure-activity relationship investigations with P1 mutants demonstrate that several flexible extracellular loops allow P1 to engage human CEACAMs. These results provide the first evidence for the involvement of the major outer membrane protein P1 of Hinf in pathogenesis.
LOCAL AND SYSTEMIC EFFECTS OF EXPERIMENTAL MURINE HELICOBACTER PYLORI INFECTION.

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Background

Despite its late-in-life pathogenicity (gastric cancer and peptic ulcer disease), due to its potential early-life benefits (protection against childhood-onset asthma, diarrheal disease, and reflux esophagitis), Helicobacter pylori (Hp) might be considered a commensal. The ongoing disappearance of Hp from the human microbiome is concerning as little is known about the influence of Hp on the host’s microbiome and immunity.

Objectives

In this study, we aimed to investigate the interactions of Hp with its host in a mouse model.

Methods

We infected C57/Bl6 mice with cagA+ Hp PMSS1, or not (control). Groups of mice were serially sacrificed over a period of 6 months and host responses analyzed.

Conclusions

All inoculated mice were Hp-positive when sacrificed, as verified with PCR and culture and displayed gastric histology typical for Hp colonization. All infected mice showed high levels of IgM/IgG to Hp but not to CagA. Nanostring® analysis, performed on stomach and lung, revealed a large number of immune genes in the stomach with increased expression. Immune gene expression was also altered in the lung and flow-cytometry confirmed significantly higher levels of Th17-cells in infected mice compared to controls. Extracted DNA from fecal pellets, gastric, ileal and cecal contents were subjected to 16S rRNA gene sequencing to determine changes in the microbiome due to Hp colonization. Overall, control and colonized mice displayed
different population structures in both their gastric and intestinal microbiota. These findings indicate that in this mouse model, *Hp* influences both the microbial population structure and local and distant host immune responses.
Background

Although *L. pneumophila* serogroup(sg)1 is the common disease causing serogroup, rare serogroups can also cause legionellosis.

Objectives

We aim to present six cases of legionellosis, of which four cases presented as Pontiac fever and the other two cases presented as Legionnaires’ disease. And than emphasize the regional epidemic and pandemic threats of infectious diseases that are transmitted between neighboring countries.

Methods

We reported 6 cases of legionellosis caused by rare sg of *L. pneumophila*. Their diagnosis was supported with clinical, Radiological and serological findings and serogrouping *L. pneumophila*. The analyses of IgG and IgM were performed in our unit using the Anti-Legionella Pneumophila Indirect Immunofluoresan IgM, IgG kit (Euroimmun AG, Leubeck,Germany). All six serum samples sent, Euroimmun AG Clinical Immunology Laboratory in Lubeck, Germany for serogrouping.

Conclusions

Two of them were diagnosed with Legionnaires’ disease caused by *L. pneumophila* sg12 (index case KY) and sg11 (his wife ZY) and 4 of the cases
were diagnosed with Pontiac fever caused by *L.pneumophila* sg14 (ZY), sg4(AK), sg4 with sg6 (BT) whereas the sg of *L.pneumophila* detected in ZC could not be identified. This study reemphasized that not only *L. pneumophila* serogroup 1, but other rare serogroups might cause also legionellosis which may increase in frequency and cause regional epidemics. It should be noted that these epidemics can also become a threat and a severe public health problem for all countries. Global-scale solutions should be developed for epidemics that could threaten this region
FEMS-1651
Bacterial pathogenicity

THE PATTERNS OF EPIYA MOTIFS AMONG CAGA POSITIVE HELICOBACTER PYLORI STRAINS: A PROSPECTIVE STUDY IN TURKISH POPULATION WITH EURASIAN GEOGRAPHICAL FEATURES

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Background
Geographical variation in the frequency of various gastroduodenal pathologies was shown to be related to the geographical diversity of H.pylori cagA/EPIYA patterns.

Objectives
We sought to determine the EPIYA patterns of cagA-positive H.pylori strains isolated from patients with different endoscopic diagnoses and from patients with gastric cancer (GC) who lived in Istanbul and surroundings areas, a Western region of Turkey located both in Europe and Asia.

Methods
One hundred and fifty-eight cagA-positive strains were included in the study; 84 were isolated from patients with gastritis (G), 38 were from patients with GC, 22 were from patients with duodenal ulcers (DU) and 14 were from patients diagnosed endoscopically with having normal gastrointestinal systems (NGIS). Specific primers were used for the detection of EPIYA patterns and representative bands were also confirmed by DNA sequencing.

Conclusions
EPIYA A, B, C or D were found in 142 of the 158 strains (89.9%). EPIYA-ABC was detected in 83 H. pylori 50 (59.5%) were isolated from patients with G, 14 (63.7%) were isolated from patients with DU, 9 (23.7%) were isolated from patients with GC and 10 (71.4%) were isolated from patients with NGIS. EPIYA-C with ≥2 repeats was detected in 34 (21.5%) cagA-positive H.pylori cases and 22(64.7%) of these cases involved GC. The most common EPIYA pattern isolated from cases with different endoscopic diagnoses was the Western type EPIYA-C. EPIYA-C with ≥2 repeats was
found to be more common in cases with GC; in other cases with different endoscopic diagnoses, EPIYA-C with one repeat was the most frequent.
Bacterial pathogenicity

IDENTIFICATION AND CHARACTERIZATION OF STRESS FIBER INDUCING EFFECTOR PROTEIN OF VIBRIO PARAHAEIMOLYTICUS T3SS2


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Background

Vibrio parahaemolyticus is an important pathogen that causes food-borne gastroenteritis in humans. The type III secretion system encoded on chromosome 2 (T3SS2) plays a critical role in the enterotoxic activity of this bacterium. Tissue culture analysis has shown that T3SS2 causes two dramatic changes in the actin cytoskeleton: the accumulation of F-actin beneath bacterial microcolonies and the induction of actin stress fibers. VopV, which is enterotoxic effector, exhibits F-actin binding activity and is responsible for the F-actin accumulation phenotype. However, the mechanisms that underlie T3SS2-dependent actin stress fiber formation and the main effector have not been elucidated.

Objectives

The aim of this study is to identify effector protein responsible for T3SS2-dependent stress fiber formation.

Methods

V. parahaemolyticus strain RIMD2210633 (KP-positive, serotype O3:K6) was used for parent strain. A four-primer polymerase chain reaction (PCR) technique was used to engineer an in-frame deletion mutation. Caco-2 cells were infected with V. parahaemolyticus for 3 h at a multiplicity of infection (MOI) of 10. Actin was detected with Alexa Fluor® 488-phalloidin. Cellular and bacterial DNAs were stained with Hoechst 33258.

Conclusions

After screening candidate ORFs encoded within the Vp-PAI region, a known pathogenicity island in pathogenic strains, we identified an effector candidate protein involved in stress fiber formation. A deletion of this gene did not affect to T3SS2-dependent enterotoxicity in rabbit ileal loop model, but this mutant caused a dramatic change in actin stress fiber formation, thereby suggesting that this protein is a T3SS2 effector involved in the induction of stress fiber formation.
STRUCTURE-FUNCTION ANALYSIS OF THE TYPE 3 SECRETION SYSTEM TRANSLOCATOR IpaB

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Background
The Type 3 Secretion System (T3SS) enables the invasion the human intestinal gut mucosa by *Shigella flexneri*. The T3SS consists of a membrane embedded basal body and an extracellular needle. At the distal end of the needle a tip complex senses and binds the host cell membrane. Upon binding secretion of effectors from the bacterial cytoplasm through the T3SS into the host cell is activated. The secretion of effectors follows a defined order and prior secretion unfolding of the effectors enables the transport through the needle. Early secreted effectors, known as translocators form pores in plasma membranes. The translocators such as the invasion plasmid antigen B (IpaB) are bound to chaperones (IpgC) in the bacterial cytoplasm. For successful infection the IpaB/IpgC complex disassembles and unfolded IpaB is secreted while IpgC remains in the bacterial cytoplasm. Thus, IpaB adopts different folding states before and after secretion which is studied by X-ray crystallography and other biophysical methods.

Objectives
Analysis and comparison of different IpaB conformations with the aim to gain insight into IpaB function.

Methods
Detergents for stabilizing the hydrophobic domains of full length IpaB were used for crystallization trials as well as full length IpaB in complex with IpgC without detergent. In another approach soluble fragments of IpaB were successfully crystallized and the structure determined. IpaB fragments were also cocrystallized with IpgC and the structural determination is in progress.

Conclusions
Based on this study, future design of small inhibitors preventing correct IpaB folding could be beneficial for therapeutic-clinical use against *Shigella* infections.

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Background
Aeromonas virulence remains poorly understood, virulence of any given strain being hardly predictable. In addition, Aeromonas human infections involve mixed Aeromonas isolates (5-10% cases) with an unknown impact on virulence.

Objectives
To evaluate virulence of strains recovered from mixed aeromonad infections, alone and in association.

Methods
Twelve Aeromonas isolates recovered in pairs from 6 mixed infections were tested against the Caenorhabditis elegans virulence model. Dead worms were counted each day during 15 days. Median time for killing worms (TD50) was determined for single isolates and for natural and experimental pairs. Experiments were repeated 3 times, independently. Genomes of 7 isolates were sequenced using an Illumina Myseq; virulence-associated genes were sought within the draft genomes.

Conclusions
Eight isolates were weak killers (TD50≥7 days) when they were administered alone, two pairs showed an enhanced virulence: the TD50 was significantly lowered from 7.8 and 8.3 days to 3.3 days (couple 76c+77c) and from 6 and 9 days to 4.5 days (couple 25a+25b). Synergy was also observed for 5 of the 14 experimental pairs tested, each including one strain from the natural synergistic couples. Only couples involving strains from distinct species showed synergy. The genome content of virulence-associated genes failed to explain virulence synergy and to identify pathotypes, although some virulence-associated genes present in some strains were absent from the companion strain (e.g., T3SS).

Synergistic virulence observed between infectious Aeromonas isolates stresses to consider Aeromonas infection process at the community level and not only according to virulence content as in pathotype concept.
Bacterial pathogenicity

STAPHYLOCOCCUS AUREUS-DERIVED MEMBRANE VESICLES EXACERBATE SKIN INFLAMMATION IN ATOPIC DERMATITIS

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Background
Skin colonization or infection with *Staphylococcus aureus* is known to trigger aggravation of atopic dermatitis (AD). However, the exact mechanisms by which *S. aureus* can worsen AD are unknown.

Objectives
We investigated whether and how *S. aureus*-derived membrane vesicles (MVs) contribute to worsening of AD.

Methods
HaCaT cells were treated with *S. aureus* MVs and were analyzed for the expression of pro-inflammatory cytokine genes. Immunopathology and cytokine gene profiles were analyzed after topical application of *S. aureus* MVs to AD-like skin lesions in a mouse model.

Conclusions
Intact MVs from *S. aureus* delivered their components to keratinocytes and stimulated pro-inflammatory cytokine gene expression. However, MVs with a disrupted membrane neither delivered their components to keratinocytes nor resulted in cytokine gene expression. A knockdown of nucleotide-binding oligomerization domain 2 by using small interfering RNAs completely suppressed IL-8 gene expression. Topical application of *S. aureus* MVs to AD-like skin lesions in the mouse model induced massive infiltration of inflammatory cells and the resulting eczematous dermatitis. This inflammatory reaction was associated with a mixed Th1/Th2 immune response and enhanced expression of chemokine genes in AD-like skin lesions. *S. aureus* MVs delivered effector molecules to host cells and triggered an inflammatory response both in vitro and in vivo. MVs produced by *S. aureus* colonizing or infecting AD skin lesions may be responsible for worsening of AD. Thus, *S. aureus* MVs are a new therapeutic target for the management of AD aggravation.
FEMS-0905
Bacterial pathogenicity

THE INTIMIN PERIPLASMIC DOMAIN MEDIATES DIMERISATION AND BINDING TO PEPTIDOGLYCAN
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Background
Intimin and Invasin are prototypical inverse (Type Ve) autotransporters and important virulence factors of enteropathogenic Escherichia coli and Yersinia spp., respectively. In addition to a C-terminal extracellular domain and a β-barrel transmembrane domain, both proteins also contain a short N-terminal periplasmic domain that, in Intimin, includes a lysin motif (LysM), which is thought to mediate binding to peptidoglycan.

Objectives
We investigated the functions of the Intimin periplasmic domain, particularly to study its role in peptidoglycan binding and dimerisation.

Methods
We used pull-down, solid phase binding and in vivo binding assays to demonstrate that the Intimin LysM binds to peptidoglycan. Dimerisation of the periplasmic domain was shown by analytical size exclusion chromatography and in vitro cross linking experiments. We further solved the structure of the Intimin LysM using NMR.

Conclusions
We show that the periplasmic domain of Intimin does bind to peptidoglycan both in vitro and in vivo, but only under acidic conditions. We were able to determine a dissociation constant of 0.8 μM for this interaction, whereas the Invasin periplasmic domain, which lacks a LysM, bound only weakly in vitro and failed to bind peptidoglycan in vivo. Furthermore, in contrast to previous reports, we show that the periplasmic domain mediates dimerisation.

We further show that dimerisation and peptidoglycan binding are general features of LysM-containing inverse autotransporters. Peptidoglycan binding by the periplasmic domain in the infection process may aid in resisting mechanical and chemical stress during transit through the gastrointestinal tract.
BRINE SHRIMP: A HOST MODEL SYSTEM TO STUDY BACTERIAL VIRULENCE AND SYMBIOSIS

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Background
Brine shrimp is an aquatic crustaceans belonging to a genus of Artemia. This organism is widely used for testing the toxicity of chemicals and for live food in the larviculture of economically important fishes and crustaceans.

Objectives
In this study, a brine shrimp, Artemia salina was evaluated for the bacterial infection and symbiotic host model.

Methods
Brine shrimp nauplii were incubated in petri dish containing 5 ml of autoclaved artificial seawater. Various numbers of bacterial cells were added to the seawater and incubated at 28°C for several days. The survival of shrimps was daily scored after the addition of the bacteria and the existence of bacteria in shrimp gut was observed by fluorescence microscope.

Conclusions
Pathogenic bacteria caused significant death of brine shrimps reflecting their virulence, but some shrimps surviving the infection were found to grow bigger and faster. Both E. coli and P. aeruginosa could survive in the brine shrimp gut, but E. coli was able to survive only for limited period whereas P. aeruginosa survived more and longer in the gut than E. coli. Furthermore, we found that the pre-infection of avirulent P. aeruginosa strain improves the survival of brine shrimp in the challenge of other pathogenic bacteria. These results strongly suggest that survival of P. aeruginosa as a symbiont has a beneficial effect on host and brine shrimps may be a valuable artificial model to study symbiosis.
DIRECT INVOLVEMENT OF CYCLIC AMP (CAMP) AND CAMP RECEPTOR PROTEIN (CRP) IN NATURAL COMPETENCE REGULATOR TFOX EXPRESSION

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Background
Vibrios are normal inhabitants in the estuarine and marine environments, where they exist both as free-living bacteria and in association with phytoplankton, zooplankton, and crustaceans. Chitin-induced competence found in several Vibrios represents a newly appreciated mode of horizontal gene transfer in these marine bacteria and undergoes tight regulation. Sxy (also called TfoX) and CRP are two important activator proteins in the competence development. Sxy functions synergistically with CRP to activate the competence genes regulon expression.

Objectives
Sxy/TfoX expression has been shown to be regulated by RNA second structure and sRNA tfoR at the post-transcriptional level. In this study, we investigated the transcriptional regulation of tfoX expression by cAMP-CRP complex.

Methods
qRT-PCR and complement assay were used to analyze the tfoX mRNA level. Promoter transcriptional fusion and site-directed mutagenesis of the putative CRP-binding sites were applied to determine the tfoX promoter activities. EMSA assay was employed to demonstrate the direct binding of CRP to the tfoX promoter. The transcriptional start site of tfoXVF was determined by 5’RACE.

Conclusions
CRP positively regulates the tfoX expression at the transcriptional level; there are two functional CRP binding-sites on tfoXVC promoter region and CRP directly binds to the promoter region to initiate its transcription; transcriptional start site of tfoXVF was mapped at -126 nucleotides upstream of translational start site; similar -10 motifs and putative CRP-binding sites on different species’ tfoX promoter indicate CRP regulation of tfoX is a conserved regulatory mechanism in Vibrio species.
REGULATION OF THE CYCLIC AMP SIGNALING PATHWAY OF KLEBSIELLA PNEUMONIAE IN TYPE 3 FIMBRIAE EXPRESSION

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Background
*Klebsiella pneumoniae* (Kp) is the predominant pathogen isolated from liver abscesses of diabetic patients in Asian countries. Although elevated blood glucose levels cause various immune problems, its effects on Kp virulence remains largely unknown. In bacteria, cyclic AMP (cAMP), a well-known second messenger, plays a fundamental role in global gene regulation in response to exogenous glucose levels. However, targets regulated by cAMP signaling pathway remain uncharacterized in Kp.

Objectives
To investigate the role of cAMP signaling pathway in Kp pathogenesis.

Methods
DNA microarray, promoter activity assay, qRT-PCR, western blotting assay, biofilm formation, electrophoresis mobility shift assay

Conclusions
Based on the analysis of DNA microarray results, we noted a serial of type 3 fimbrial genes was apparently increased in Δ*crp* strain. It indicates that CRP could repress the type 3 fimbriae expression. By using promoter activity assay, western blotting, and qRT-PCR to confirm that the MrkA expression, the major pilin of type 3 fimbriae, was regulated by cAMP signaling pathway. However, no typical CRP binding site was found in the sequence of *P*<sub>mrkA</sub>, suggesting CRP regulates the *mrkA* transcription via other mediator(s). According to previous studies, multiple proteins, which include Fur, MrkH, MrkI, MrkJ, and YjcC have been shown to mediate the type 3 fimbriae expression. The underlying mechanism of cAMP signaling pathway and the mediators as described above in regulating the type 3 fimbriae expression was investigated in this study.
IDENTIFICATION OF FITNESS DETERMINANTS IN PSEUDOMONAS AERUGINOSA INFECTION

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Background

Pseudomonas aeruginosa is a clinically highly relevant opportunistic pathogen known to cause acute and chronic infections. This bacterium can adapt to the host by environmentally driven alterations in gene expression as well as genomic alterations in response to selective pressures by the environment.

Objectives

The aim of this study is to identify genetic adaptations that are key to fitness and successful establishment of acute and chronic infections by P. aeruginosa.

Methods

We have developed an in vivo model for P. aeruginosa biofilm formation in CT26 tumor-bearing mice. This murine model offers the unique opportunity to screen the contribution of individual genes to the fitness of P. aeruginosa during in vivo biofilm formation. Furthermore, the inclusion of an acute lung infection model allows the comparison of fitness parameters in chronic to acute infections. These parameters will be investigated using (i) a PA14 transposon mutant library (Skurnik et al. 2013) as forward genetic approach and (ii) a reverse genetic approach by passaging bacteria in the animal host. To evaluate the contribution of individual genes onto bacterial fitness, isolated bacteria are quantified using high-throughput insertion and whole-genome sequencing, respectively.

All animal experiments were approved by the local regulatory board LAVES (Niedersächsisches Landesamt für Verbraucherschutz und Lebensmittelsicherheit) and the EU Directive 86/609/EEC, OJ L 358, p 1-28; 18.12.1986.

Conclusions

Here we present a dual approach to gain insights into fitness determinants important during P. aeruginosa infection in different host environments.
WILD-TYPE AND NONBINDING MUTANT OF TRANSFERRING BINDING PROTEIN B (TBPB) HAEMOPHILUS PARASUIS SEROVAR 5: IMMUNOENZYMATIC AND GENE EXPRESSION STUDIES

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Background
Glasser's disease, caused by Haemophilus parasuis (Hps), produces annually pig significant economic losses. It is broadly known the relevance of transferrin binding proteins (Tbps) as virulence factor for developing this disease in the host.

Objectives
Subunit vaccines based on the native, wild-type TbpB and on a mutant TbpB (TbpBY167) (losing its ability for binding porcine transferrin) from Hps serovar 5 were designed.

Methods
Immunization and challenge (with an intratracheal administration of 10⁸ CFU/ml of the Nagasaki strain of Hps 5) experiment was implemented in colostrum deprived pigs for comparing the protective response developed for wild-type and mutant proteins. Clinical symptoms were monitored after challenge and pathological findings were determined upon necropsy. All the pigs immunized with adjuvant alone and challenged did not survive more than 72 hours. On the other hand, most of the pigs immunized with wild-type TbpB survived the full two weeks after challenge but clinical signs and severe chronic lesions were observed. All the pigs immunized with the mutant also survived until the end of the study, without signs of infection. Were confirmed the results by indirect ELISA test, by immunohistochemistry and real-time quantitative PCR was used to determine the relative expression of 27 immune-related genes in the lungs of the pigs belonging to experimental groups, at the necropsy time.

Conclusions
The highest protective response was that afforded by the mutant. Genes associated with the inflammation are highly overexpressed in the group of only infected animals and this does not happen in the other two experimental groups.
Background
Uropathogenic *E. coli* (UPEC) strains are causative agents in the majority of human urinary tract infections (UTIs). UPEC strains contain several virulence factors allowing them to colonize human urinary tract (e.g. fimbriae type I). Moreover, higher prevalence of colicin E1 like plasmids and a possible role of colicin E1 protein as a potential important virulence factor of UPEC strains was described in UPEC strains.

Objectives
The main aim of the study focused on the genetic variability of colicin E1 and *fimH* gene (encoding the mannose-sensitive FimH adhesin of fimbriae type I) between human intestinal and extraintestinal *E. coli* strains (UPEC strain).

Methods
In the set of 58 human fecal *E. coli* strains and 55 UPEC strains, *fimH* gene and colicin E1 gene was sequenced and analyzed. All clinical samples were collected after patients gave informed consent. The study was approved by the ethics committee of the Faculty of Medicine, Masaryk University, Brno, CZ.

Conclusions
Two different colicin E1 genetic variants were described. Colicin E1-A genetic variant that prevailed among human fecal *E. coli* had 1566 bp and colicin E1-B genetic variant (prevailed in UPEC strains) had 1569 bp. The main genetic difference identified between these variants encoded colicin receptor domain. A significant increase of the substitution of residue Arg166 - His166 in fimbriae-associated pilin domain of FimH was identified in UPEC strains. It is possible that genetic variability identified in genes encoding colicin E1 and FimH modify virulence of UPEC strains.
FEMS-2996
Bacterial pathogenicity

DIFFERENCES AND SIMILARITIES BETWEEN S. ENTERICA SEROVARS TYPHIMURIUM AND ENTERITIDIS FIS MUTANTS UNDER IN VITRO CONDITIONS
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Background
Salmonella enterica is the causal agent of a widespread and neglected infectious disease in the world, that despite efforts, it lacks completely effective vaccine formulations. The non-typhoidal serovars Typhimurium and Enteritidis are frequently responsible for salmonellosis outbreaks, usually associated with gastrointestinal infections, but they can spread systemically, particularly in immunocompromised patients or those who suffering from malaria. In Africa, non-Typhi Salmonella comprises around 50% of reported cases of bacteremia and are associated with nearly 20% of the fatal cases of salmonellosis. It’s crucial to test the potential of new mutants and new serovars for use as live attenuated vaccines to contain the disease.

Objectives
We used recent clinical isolates of S. Typhimurium and S. Enteritidis to develop the strains LGBM01Δfis and SEnPT4Δfis, respectively, and evaluated their phenotype under in vitro conditions.

Methods
The phenotypic tests performed included analysis of growth in Luria Broth with measurements taken every hour, motility test on semi-solid agar plates and invasiveness in primary macrophages obtained from BALB/c mice.

Conclusions
The results showed no significant differences in growth rate in Luria Broth between the mutant strains and the respective wild strain, but the motility in semi-solid agar plates was significantly decreased, at least 50%, in the mutant strains. Interestingly, the ability in invade primary macrophage cells was significantly affected only in the Typhimurium mutant strain. Those results showed an important difference for the role of the same DNA binding protein in these serovars that needs to be considered for future use in vaccine development.
PATHOGENIC FEATURES OF BACTERIAL STRAINS ISOLATED FROM SURGICAL RINSE WATER AND MEDICAL INSTRUMENTS

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Background

One of the potential sources for bacterial contamination in hospital environments is represented by the inadequate manual cleaning of medical instruments.

Objectives

The purpose of this study was to investigate the pathogenic features of microbial strains isolated from sterile water use for rinsing the surgeons hands and surgical medical instruments, in order to establish potential contamination risks for patients.

Methods

A number of 200 water samples and 400 surgical instruments have been processed for routine microbiological analyses, following the existent ISO guidelines. The isolated strains have been identified using miniAPI galleries and investigated for antibiotic susceptibility profiles and for the production of cell-associated (adherence to inert and cellular substratum, biofilm development) and soluble, enzymatic (hemolysins, lecithinase, lipase, caseinase, gelatinase, amylase, esculin hydrolysis, DN-ase) virulence factors, using phenotypic (disk diffusion method, selective media for enzymatic factors production) and PCR-based methods.

Conclusions

Bacteria were isolated in 10% of the analyzed water samples and from 5% of the investigated instruments. The following bacteria were isolated in the frequency decreasing order: Pseudomonas aeruginosa, Brevundimonas sp., Micrococcus luteus, Staphylococcus haemolyticus, S. warneri, P. putida, P. stutzeri, Stenotrophomonas maltophilia, Moraxella sp. Pore-forming toxins and enzymes have been the most frequently produced, followed by caseinase and esculin hydrolysis. The genetic support of these virulence factors has been confirmed by PCR. The analyzed strains exhibited only natural resistance profiles, demonstrating the
environmental origin of the isolated strains. The highest frequency of pseudomonades suggests that biofilm formation within piping is the source of contamination of surgical rinse water.
Background
Typhoid fever is a systemic infection caused by Salmonella enterica serovar Typhi and transmitted from human to human via food or drinking water; therefore, hygiene and sanitary conditions mainly determine its spread. As a result of improved public sanitation, most of the cases in Japan have been sporadic and have come from abroad in recent years. In 2013, however, we observed an unusual increase of typhoid fever cases which seemed to be domestic infection.

Objectives
In this study, we assessed increase of the typhoid fever patients without histories traveling abroad and examined molecular epidemiological analysis of S. Typhi.

Methods
A total of 27 typhoid fever cases without histories traveling abroad were identified in 2013 and 26 S. Typhi isolates were collected. All isolates were examined with phage type and MLVA.

Conclusions
Of the 26 isolates, 15 strains were classified as phage type A or B1 and formed major cluster with 5 types of single locus variants by cluster analysis of the MLVA profiles. It seemed that closely related strains were causative agents of increased typhoid fever cases in Japan, 2013. These 15 patients were reported during 5 months (July to November) peaked at September. They ranged in age from 6 to 83 years (median age 32 years) and the male:female ratio was 1:1.5. Interviews by physicians or public health officers could not reveal specific information which leads to epidemiological link; therefore we have no identifiable source of infection.
Background
Burkholderia cepacia complex (Bcc) bacteria pose major threat to cystic fibrosis (CF) patients. Bcc infections can result in untreatable and fatal cepacia syndrome (CS) which is manifested as rapid deterioration of lung functions and sepsis. The mechanisms behind transition from Bcc chronic infection to CS remain unknown.

Objectives
To gain insight into processes involved in development of CS, we compared transcriptomes of a lung isolate (chronic infection) and a bloodstream isolate (CS) from the same patient. Both isolates were of the same sequence type (ST872) and belonged to the Bcc species B. contaminans which is predominant among CF infections in Argentina.

Methods
Both strains were cultivated in three growth media: sputum (natural habitat of lung isolate), heat-inactivated human serum (natural habitat of bloodstream isolate) and control mineral medium, each in biological triplicates. RNA was extracted in mid-log growth phase, sequenced using RNA-Seq technology (Illumina) and converted to normalized transcript level values.

Conclusions
The expression of approximately 1,300 (18%) genes differed more than 3-fold between lung and sputum isolate. The bloodstream isolate showed markedly increased expression of quorum sensing-regulated pathogenicity determinants (motility, extracellular proteases, AidA, lectins, Fip pilus), hypoxia-activated genes and two antifungal compound synthetic clusters (occidiofungin and pyrrolnitrin). Agar plate assays confirmed rapidly increased motility and proteolytic, hemolytic and antifungal activities of bloodstream isolate. Assessing these phenotypes might help to monitor the progress of infections by B. contaminans.
This work was funded by grant from Ministry of Health of the Czech Republic NT12405-5.
Background

*H. pylori* is the main cause of chronic gastritis, ulcer and gastric cancer. The balance between the bacterial virulence factors and the host immune response in the infection determines the different clinical outcomes. The association between Th1, Th17, Treg cells and *H. pylori* infection has been identified, but the effect of the nine major *H. pylori* specific virulence factors; cag A, vacA, oipA, babA, hpaA, napA, dupA, ureA, ureB on Th1, Th17 and Treg cells response in *H. pylori* infected patients has not been fully elucidated.

Objectives

To investigate the correlation of the *H. pylori* virulence factors with the manifestations of gastric disease by developing a multiplex–PCR and the relationship between these virulence factors with Th1, Th17 and Treg cells.

Methods

Multiplex and qRT-PCR were carried out to detect the *H.pylori* specific virulence factors and relationship between these virulence factors with T cells.

Conclusions

The multiplex-PCR was developed to rapid characterisation of nine *H. pylori* virulence genes with in a three PCR reactions. Due to the multiplex-PCR results there was no significant difference in prevalence of virulence factors in patients with gastritis and ulcer. However, prevalence of *napA* virulence factor was significantly higher in
patients with ulcer than gastritis. A positive correlation between the *dupA* virulence factor and IFN-g was observed in gastritis patients. Additionally, IL-17 expression was found significantly positively correlated with the *babA* virulence factor in ulcer patients. Furthermore, a novel expert derived model is developed to identify set of factors and rules distinguishing the ulcer patients from gastritis patients.
IMPACT AND ROLE OF THE RNA CHAPERONE HFQ IN VIRULENCE GENE REGULATION OF LEGIONELLA PNEUMOPHILA

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Background

Legionella pneumophila, the causative agent of the pneumonia-like Legionnaires’ disease, is commonly found in aquatic habitats worldwide where it multiplies within protozoa. To adapt between intra- and extracellular environments, L. pneumophila evolved a biphasic lifecycle wherein it alternates between a replicative (non-virulent) and a transmissive (virulent) phase. This switch is governed by a complex regulatory network. Expression of the Host Factor Q protein (Hfq), a hexameric, RNA-binding protein and chaperon of small RNAs (sRNA) is life cycle regulated and implicated in virulence. How Hfq expression is regulated in L. pneumophila is not known.

Objectives

Our aim was to understand how the growth-phase dependent expression of Hfq is regulated and whether a newly identified sRNA is implicated.

Methods

By Transcriptional Starting Site mapping of the L. pneumophila genome, a sRNA that is transcribed antisense to the hfq gene and is overlapping its 5’UTR region was identified. Thus, we postulate that this antisense RNA, named anti-hfq, might regulate the life cycle dependent expression of Hfq. We constructed and characterized the L. pneumophila hfq deletion mutant and anti-hfq overexpressing strains in vitro and in vivo.

Conclusions

Our results show that Hfq expression is growth-phase dependently regulated by this newly identified antisense RNA. Infection assays revealed that Hfq is a virulence factor necessary for efficient replication in amoeba. Overexpression of anti-hfq led to a similar virulence phenotype as hfq deletion. This suggests an important role of Hfq and anti-hfq in virulence and in the regulatory network governing the biphasic life cycle of L. pneumophila.
ISOLATION AND CHARACTERIZATION OF OUTER MEMBRANE VESICLES OF THE FISH PATHOGEN PISCIRICKETTSIA SALMONIS.

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Background

Piscirickettsia salmonis is a Gram-negative intracellular bacterium that causes Piscirickettsiosis in salmonids farms in Chile. It was recently reported that P. salmonis produces exotoxins that play a role in the pathogenesis. However a delivery system has not yet been identified. Outer membrane vesicles (OMVs) are 10 to 300 nm spherical-bilayer structures discharged from the surface of many Gram-negative bacteria which are able to deliver toxins and virulence factors. However, the production of OMVs by P. salmonis has not been described.

Objectives

The aim of this study was to investigate if P. salmonis is able to produce OMVs and to realize a microscopic characterization of these vesicles.

Methods

P. salmonis was grown in basal broth supplemented with Cysteine (3.18 mM) and ferric chloride (0.05 mM) at 18°C until early stationary phase. Bacteria were removed by centrifugation (5000 x g, 10 min at 4°C) and the supernatant were filtrated through a 0.22-μm-pore-size filter. Then, vesicles present in the bacterial free supernatant were isolated by ultracentrifugation (125,000 x g, 2h at 4°C) and analyzed by SDS-PAGE. Finally, CHSE-214 cells infected with P. salmonis and negative stained OMVs were visualized by TEM.

Conclusions
*P. salmonis* is able to produce OMVs. The purified OMVs appeared as spherical vesicles of different sizes between 27.3 and 145.5 nm. SDS-PAGE analysis showed similar protein profile between OMVs and outer membrane extracts. Finally, OMVs were found into *P. salmonis*-containing vacuoles in infected CHSE-214 cells, suggesting that OMVs may contribute to the pathogenesis of *P. salmonis*. 
Bacterial pathogenicity

MULTIPLE COLONIZATION OF HELICOBACTER PYLORI IN PATIENTS WITH CHRONIC GASTRITIS OF REGIONS OF HIGH AND LOW RISK OF GASTRIC CANCER IN COLOMBIA

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Background
Infection with virulent strains of Helicobacter pylori is necessary but not enough for the development of gastric cancer (GC) and its precursor lesions. In Nariño-Colombia, there are populations with high prevalence of H. pylori infection, but different GC risk. These results suggest different genetic characteristics of the circulating strains with varying degrees of pathogenicity.

Objectives
To evaluate multiple colonization of H. pylori isolated from patients with chronic gastritis from two populations with contrast in the risk of developing GC: Tumaco, low risk; Túquerres, high risk. Our hypothesis: there is a greater probability of multiple colonization of H. pylori in the population at highest risk for GC.

Methods
409 adult patients with dyspeptic symptoms were studied, of which gastric mucosa fragments from the antrum and body were used for the histopathological diagnosis, culture and genotyping (cagA, vacA and PCR-RAPD). 72 individuals were included, in whom H. pylori isolates was achieved in the three biopsies of gastric mucosa, (41/203) from Tumaco and (31/206) Tuquerres. Genetic similarity was calculated using the Nei’s coefficient.

Conclusions
Genetic diversity was higher among isolates from Tuquerres (0.13) than Tumaco (0.07). After adjusting for age, sex and diagnosis, multiple colonization was 1.7 times more frequent in Tuquerres than Tumaco, p = 0.05. In Túquerres: high risk of gastric cancer, there was more likely to have multiple colonization. Based on the analysis of the results of the PCR - RAPD, we found greater genetic similarity from isolates of H. pylori in the population with low risk for developing GC.
IDENTIFICATION OF GENES INVOLVED IN VIRULENCE OF PUM505 PLASMID FROM PSEUDOMONAS AERUGINOSA
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Background
pUM505 is a conjugative plasmid of 123 kilobases, originally isolated from a clinical strain of Pseudomonas aeruginosa. pUM505 possesses a pathogenicity island (PAI) of 78 genes, 64 of them have been found in the chromosomal PAPI-1 and PAPI-2 PAs of P. aeruginosa PA14, a virulent clinical isolate, however it is unknown whether pUM505 is involved in virulence.

Objectives
The aim of this study is to determine if the plasmid increases the virulence of the P. aeruginosa PAO1 standard strain and to identify the genes responsible for this property.

Methods
Using as models of virulence lettuce leaves and Dictyostelium discoideum cultures pUM505 showed increased virulence of the PAO1 strain. To identify genes involved in virulence a gene bank was constructed by digesting the pUM505 plasmid and ligating the fragments to the vector pUCP20. A library of 120 clones was obtained in the Top10 strain of E. coli and PAO1 of P. aeruginosa. Four clones of this library were found to increase virulence in both E. coli and P. aeruginosa. The cloned inserts of these transformants was sequenced, and several ORF’s of pUM505 were identified. The ORFs 2, 17 and 42 (hop) have homologues that have been reported involved in virulence.

Conclusions
In conclusion, pUM505 plasmid increases the virulence of its hosts due to some ORF’s outside the pathogenicity island. The ORF’s identified are currently under study.
Background

Many Gram-negative bacterial pathogens use a type III secretion system (T3SS) to deliver virulence factors into the cytosol of host cells in order to establish and maintain an infection. This system relies on the sequential secretion of first structural and later effector proteins for its correct assembly and function. The hierarchy of substrate secretion has been proposed to be ensured by a cytoplasmic protein complex, called the "sorting platform".

Objectives

We aimed to investigate and characterize the Salmonella Typhimurium "sorting platform" and its components, particularly SpaO, with regard to both their structure and function.

Methods

The two expression products of the spaO gene were analyzed and the gene mutated to investigate the mechanism by which the second, shorter product is generated. The two proteins were analyzed by SDS-PAGE, mass-spectrometry, size-exclusion chromatography and multi-angle light scattering. The SpaO protein was further co-purified with other "sorting platform" components and the ability of the complex to bind T3SS substrates tested by surface plasmon resonance.

Conclusions

The gene for the component SpaO gives rise to both the full-length protein (SpaO) and a shorter product SpaO'. SpaO' is produced from an internal translation initiation site and is required for the stability of full-length SpaO. Together these two proteins associate with other "sorting platform" components to form a complex that is capable of binding different T3SS substrates.

References
Background
Recent studies have been shown that Helicobacter pylori has an effective role in the migration of bone marrow derived mesenchymal stem cells (BMD-MSCs) throw to gastric tissue.

Objectives
This study aimed to find the effect of H.pylori on as bacterial microenvironment on BMD-MSCs transforming into cancer stem cells and metastasis of the tumor cells.

Methods
BMD-MSCs were followed for evaluation by flow cytometric analysis with the hAbs for positive and negative of surface markers and treated under the osteogenic and adipogenic differentiation medium. BMD-MSCs were co-cultured with H. pylori and Gastric epithelial cell line (AGS). The expressions of MMP-2, MMP-9, p53 and bcl2 were examined by qRT-PCR

Conclusions
When mesenchymal stem cells are attracted to chronic H.pylori infection tissue and gastric ulcer for their tissue healing function, they will be trapped under special microenvironment. This study demonstrated that H. pylori increased the anti-apoptosis factor bcl-2 to keep BMD-MSCs alive and lead them to cancer stem cells by changing the regulation of p53. H. pylori increased the metastatic proteins MMP2 and MMP9 that shows the role of H.pylori on metastasis action. Focusing on H.pylori-induced molecular pathogenesis and the impact of microenvironment in gastric progenitor cells or BMD-MSCs will be crucial to identify the molecular targets in tumor initiation and the origin of gastric cancer.
A NOVEL, UNSUSPECTED FEATURE OF SALMONELLA ENTERICA SPI-1 BISTABILITY

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Background
Salmonella enterica pathogenicity island 1 (SPI-1) is a gene cluster that encodes a type 3 secretion apparatus and effectors involved in invasion of epithelial cells. A well known trait of SPI-1 is bistable expression, which generates SPI-1 ON and SPI-1 OFF subpopulations. The biological significance of SPI-1 bistability has been addressed by previous, insightful studies. Bistability has been viewed, for instance, as a division of labour involving self-destructive altruism by the SPI-1 OFF subpopulation (Ackermann et al. Nature 454, 987-90, 2008). Another study, however, has envisaged that the SPI-1 OFF subpopulation might benefit from inflammation triggered by the SPI-1 subpopulation (Stecher et al. PLOS Biology 5:2177-89, 2007). Furthermore, enhanced tolerance to antibiotics has been detected in slow-growing SPI-1 ON cells (Arnoldini et al. PLOS Biology e1001928, 2014).

Objectives
In this communication we describe an additional, unsuspected feature of SPI-1 bistability.

Methods
Single-cell analyses, such as flow cytometry and cell sorting were used to study the expression of SPI-1 in Salmonella enterica and its ability to invade epithelial cells.

Conclusions
We show that a pure SPI-1 ON population obtained by bacterial cell sorting is non invasive, suggesting that the SPI-1 OFF subpopulation plays an active role in invasion. In support of this view, we also show that the invasion defect associated to unimodal expression of SPI-1 can be suppressed by mutations that permit formation of a SPI-1 OFF subpopulation.
THE INVOLVEMENT OF STREPTOCOCCUS ANGINOSUS INFECTION AND THE ABERRANT ACTIVATION-INDUCED CYTIDINE DEAMINASE EXPRESSION IN HUMAN ORAL SQUAMOUS CELL CARCINOMA

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Background
Streptococcus anginosus infection could be closely related with oral squamous cell carcinoma, although the mechanism underlying carcinogenesis is still unknown. Recent studies suggested that the aberrant expression of activation-induced cytidine deaminase (AID) in the epithelial cells could result in the generation of nucleotide alterations in tumor-related genes and possible malignant transformation of the AID-expressing cells.

Objectives

Using surgical tissue samples and cultured epithelial cells, a possible involvement of the aberrant AID expression in oral squamous cell carcinoma and S. anginosus infection was assessed.

Methods

The tissue specimens were obtained from the 17 patients with oral cancer after giving informed consent. S. anginosus infection and the aberrant AID expression were assessed by the species-specific PCR and RT-qPCR, respectively. Further, the aberrant AID expression and NF-κB activation were examined by a dual luciferase assay and RT-qPCR in three epithelial cell lines and the primary human gingival epithelial cells after stimulation with an S. anginosus antigen, SAA.

Conclusions

Both S. anginosus infection and the aberrant AID expressions were frequently observed in the tissue specimens (47% and 41%, respectively), and the infection was significantly correlated with the aberrant AID expression. The stimulation of the cultured cells with SAA could induce the NF-κB activation and aberrant AID expression in all the epithelial cells tested, and the addition of an inhibitor of NF-κB activation abrogated the aberrant AID expression. Thus, S. anginosus infection could be closely related with oral squamous cell carcinoma through the induction of the aberrant AID expression by S. anginosus antigen(s).
Background

*Cronobacter* spp. are Gram-negative, motile, non-sporeforming, facultative anaerobic bacteria that have been implicated in rare but severe cases of illnesses predominantly in premature and newborn infants. These organisms have been isolated from a wide range of environments, including humans, different animals as well as (raw and processed) food and food production facilities.

Objectives

Although progress has been made during the last years, there is still a lack of knowledge on the virulence-associated factors and processes involved during pathogenesis.

Methods

To survey the pathogenesis of *Cronobacter*, a *C. sakazakii* ES5 transposon mutant library was screened in several cell lines for genes that are involved in adhesion to and invasion into host cells as well as subsequent intracellular survival.

Conclusions

The majority of attenuated mutants carried insertions in genes involved in energy production and conservation, cell membrane biogenesis, biofilm formation, motility and metabolism. Several selected genes are being functionally analyzed such as genes for flagellar biosynthesis. Preliminary analyses showed that *C. sakazakii* flagella are necessary for adherence and efficient invasion of host cells. Using a specific monoclonal antibody against a living strain of *C. turicensis* 3032, flagellar motility could be inhibited and the ability to invade Caco-2 cells was reduced. Therefore, we present evidence that *Cronobacter* flagella play a crucial role in colonization of host cells.
FEMS-1138
Bacterial pathogenicity

TRANSCRIPTOME ANALYSES TO UNDERSTAND INFECTION MECHANISM OF BURKHOLDERIA GLUMAE IN TWO DIFFERENT INFECTION SITES OF RICE

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Background

Burkholderia glumae is a causal agent in grain and sheath rot, leading to severe damages in many rice-growing countries under the favorable conditions such as high temperature and humidity. However, its molecular mechanisms are not yet fully understood. For understating of infection mechanism, we analyzed B. glumae transcriptome from different infected rice tissues.

Objectives

We investigated molecular and physiological changes of B. glumae through comparative analyses of differentially expressed genes (DEGs) between in infected grains and in stems.

Methods

To gain a genome-wide gene expression profiling, we compared B. glumae transcriptome in stem and grain infection sites using the RNA sequencing. We then performed KEGG pathway and module enrichment of differentially expressed genes.

Conclusions

Most enriched metabolic and signaling pathways of B. glumae in infected stem tissues were bacterial chemotaxis-mediated motility, ascorbate metabolism, and sugar transporters including arabinose and xylose. However, we have confirmed different expression levels of genes involved in those pathways from the infected grains. For example, genes involved in flagellar assembly pathway were strongly down-regulated compared to those of stem tissues. Starch metabolisms were found to be highly enriched in infected grain tissues. Our study provides in vivo transcriptional profiling of B. glumae in two different tissues. Additionally, comparative analysis of B. glumae transcriptome obtained from the stem and grain will give clues on common and unique infection mechanisms in different tissues.
PREVALENCE OF COLIBACTIN GENES IN ESCHERICHIA COLI RECOVERED FROM URINARY TRACT INFECTION FROM PATIENT WITH GYNECOLOGICAL CANCER

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Background

Patients with gynecological cancer have increased risk for urinary tract infection (UTI) in which Escherichia coli is the main bacteria. The rates of antimicrobial resistance are increasing among these bacteria. Moreover, E. coli exhibits multiple accessory traits, including yersiniabactin. This virulence factor is associated with a peptide-polyketide compound, colibactin. This genotoxin, which damages DNA in eukaryotic cells, is related to the formation of colorectal cancer. However, little is known about colibactin production ratio in E. coli and its relation with antimicrobial resistance.

Objectives

To assess the prevalence of yersiniabactin and colibactin genes in a culture collection of E. coli isolated from patients with gynecological cancer with UTI and its relation with antibiotic resistance profile.

Methods

Two hundred eighteen E. coli UTI isolates were analyzed by PCR for detecting the phylogenetic group, fyuA, clbB and clbN genes. The isolate’s antibiotic resistance profile was performed previously. Statistical analysis was performed using Fisher’s test.

Conclusions

Virulence genes clbB and clbN were detected in 14.7% (32/218) of E. coli isolates. All strains in which these genes were detected belonged to the phylogenetic group B2 and had fyuA gene. Interestingly, multidrug resistance, defined as resistance to three or more antibiotic classes, showed to be a negative factor for the presence of colibactin production-related genes (p=0.0008). Colibactin genes were not detected in aminoglycosides, gentamicin and tobramycin, resistant strains (p=0.05) neither in fluoroquinolones, ciprofloxacin and norfloxacin, resistant strains (p<0.0001). Finally, 50.8% (32/63) of fluoroquinolone-sensitive strains belonging to the phylogenetic group B2 were positive for clb genes (p<0.0001).
Background

*Moraxella catarrhalis* is a human respiratory pathogen that causes acute otitis media in children and is a common cause of exacerbations in patients with chronic obstructive pulmonary disease. The first step for *M. catarrhalis* colonization is adherence to host tissues that is achieved by interaction with host extracellular matrix (ECM) proteins.

Objectives

Characterization of the interaction between *M. catarrhalis* with host collagens present in respiratory system.

Methods

Collagen I – VI were analysed for their interaction with *M. catarrhalis* clinical isolates by using ELISA and a radioactive direct binding assay. Collagen binding capacity of *M. catarrhalis* was also verified by coating collagens on the glass surface, followed by subsequent bacterial binding and detection by Gram staining. Molecular interaction between different human tissues/collagens and bacterial proteins were performed by EM.

Conclusions

Ubiquitous surface protein (Usp) A2 and UspA2 hybrid (UspA2H) of *M. catarrhalis* were identified as major collagen interacting proteins. The mutants of UspA2 and UspA2H exhibited defective collagen interaction and did not recognize collagen coated on glass surface. All clinical isolates showed binding capacity to fibrillar collagens type I, II and III, and network forming collagen IV and VI. Only 20% clinical isolates showed collagen IV binding capacity. Our data suggests that UspA2/A2H based interaction to collagens could be critical for adhesion of *M. catarrhalis* to host tissues and plays an important role in pathogenesis.
Background

Cronobacter spp. are opportunistic pathogens that cause severe infantile meningitis, septicemia, or necrotizing enterocolitis. Contaminated powdered infant formula has been implicated as the source of this pathogen in most cases, but questions still remain regarding the likely habitat and virulence potential for this strain.

Objectives

The motive of this study was to determine the occurrence of putative plasmid-harbouring virulence traits in C. sakazakii.

Methods

The isolation was done as per ISO 22964:2006 on chromogenic media followed by further confirmation by phenotypic and genotypic characterization. Two putative virulence genes Cronobacter plasminogen activator (cpa) and ferric-iron transporter eitABCD gene (eitA) were detected by PCR. The cpa gene is reported to provide serum resistance to C. sakazakii whereas eitA gene is generally associated with iron uptake mechanism especially from breast milk and infant formula by Cronobacter spp. which may enhance its spread and invasion in a host.

Conclusions

In our investigation, 15 C. sakazakii isolates were identified from 154 food and environmental samples based on phenotypic and genotypic characterization. PCR screening using eitA-specific primers revealed that the majority (67%) of the C. sakazakii isolates harbor the eitA gene while in contrast only 27% of the isolates were positive for the gene cpa. Taken together, these properties may contribute to the systemic survival of C. sakazakii and subsequent invasion of the central nervous system to cause disease. The current study clearly indicates that foods of plant origin are one of the most possible natural reservoirs of this pathogen and also provide important insight into this virulence plasmid from an emerging pathogen.
Background
Pathogenic Leptospira are the etiological agent of leptospirosis, a disease of human and veterinary concern. The ability of pathogenic leptospires to survive the bactericidal activity of human sera was demonstrated. Recently, it has been shown that leptospires acquire soluble regulators to evade the immune system.

Objectives
To assess the effect of the leptospiral adhesin Lsa23 on the activity of complement system.

Methods
The gene LIC11360 was cloned and the protein Lsa23 expressed in Escherichia coli. The effect of Lsa23 on classical pathway of complement was assessed by hemolytic assay using antibody-coated sheep erythrocytes, while the effect on alternative pathway was evaluated measuring the deposition of C5b9 on zymosan in presence of MgEGTA buffer. Interaction with factor H (FH), C4BP and plasminogen was evaluated by ELISA. Cofactor activity of complement regulators and proteolytic activity of plasmin were assessed by immunoblotting.

Conclusions
Lsa23 was expressed and purified successfully, as visualized by SDS-PAGE. It has been able to inhibit classical pathway-mediated hemolysis by 83%, and also blocked the alternative pathway reducing C5b-9 deposition by 90%. The interaction of Lsa23 with soluble C4BP, FH and plasminogen was dose-dependent. Competition assays suggest these components have distinct binding sites on Lsa23. C4BP and FH preserved their cofactor activity for factor I when bound to Lsa23, as demonstrated by cleavage of C4b and C3b, respectively. Plasminogen bound to Lsa23 could be converted into plasmin and degraded C3b and C4b. Thus Lsa23, a surface receptor for human plasma components, prevents the lytic activity of complement system and could contribute to leptospiral immune evasion process.
FEMS-0932
Bacterial pathogenicity

THE VIRULENCE EFFECT ON A HIGHLY CONSERVED REGION OF GENES RESPONSIBLE FOR CAPSULAR POLYSACCHARIDE SYNTHESIS IN KLEBSIELLA PNEUMONIAE
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Background
Capsular polysaccharides (CPS) are one of the major virulent determinant in Klebsiella pneumoniae (KP). Among different serotype of KP, CPS gene cluster could be distinguished into two regions, a conserved and a hyper-variable region.

Objectives
Virulence's study by genes' knockout, phagocytosis, serum resistance and mice lethality on highly conserved region of genes in CPS cluster was performed.

Methods
Serotype K20 KP was selected. Six genes including galF, acidPPc, wzi, wza, wzb or wzc in CPS conserved region were knocked out and assessed their effect on virulence. In comparing to parental K20 isolate, mutants showed a varied decline in mice lethality (LD₅₀) from 10 fold to > 10⁵ fold and could be categorized into low (L), moderator (M), and High (H) effect on virulence. For polycistronic mRNAs driven by P1 promoter consisted galF and acidPPc, low effect on serum resistance and anti-phagocytosis was observed. Only 10 fold of decreasing in mice lethality was achieved in the DgalF and DadicPPc mutants. For the rest of wzi, wza, Dwzb or Dwzc that were driven by P2 promoter, disruption of capsule surface assembly, Dwzi, had moderate effect with reduced mice lethality for 100 fold. The deletion of genes, Dwza, Dwzb or Dwzc which were involved in CPS polymerization, caused a significant decrease on virulence (LD₅₀ >10⁷) and became phagocytic susceptible indicating the importance of these genes in synthesis of capsule.

Conclusions
Although conserved genes in CPS cluster were all involved in KP-CPS synthesis, they contributed differently in virulence.
UNRAVELING THE FUNCTION OF A LEE-ENCODED ORF: ESCK (ORF4) IS A STRUCTURAL COMPONENT OF THE TYPE III SECRETION SYSTEM OF ENTEROPATHOGENIC ESCHERICHIA COLI

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Background
The type III secretion system (T3SS) is a molecular device employed by many pathogenic bacteria to translocate a set of proteins, called effectors, directly into the cytoplasm of eukaryotic cells in order to manipulate normal cellular processes to the benefit of the bacterium.

Enteropathogenic Escherichia coli (EPEC), an important causative agent of infant diarrhea, relies on a T3SS to deliver effectors into intestinal enterocytes allowing it to colonize the host gastrointestinal tract. Structural components assembling this biological machinery are encoded on a chromosomal pathogenicity island named the locus of enterocyte effacement (LEE).

Although the vast majority of LEE-encoded proteins have an assigned function, there are still few of them uncharacterized. One such example is the protein encoded by the escK gene (formerly orf4).

Objectives
To elucidate the role of the LEE-encoded protein EscK in T3SS biogenesis and function.

Methods
An EPEC ΔescK null mutant was generated to assess its type III secretion phenotype. An extensive search of EscK protein-protein interactions with components of EPEC T3SS was performed using the yeast two-hybrid system. Novel protein interactions were corroborated by pull-down experiments. Bacterial fractionation experiments and site directed mutagenesis were carried out in order to dissect the molecular function of this protein.

Conclusions
EscK is a critical component of the EPEC T3SS, it is localized to the inner membrane and its localization does not depend on other T3SS components. EscK interacts directly with the C-ring/sorting platform component EscQ. Essential residues for EscK function were identified.
Background

*Escherichia coli* is a commensal bacterium that can be found in the intestinal microbiota. However, pathogenic *E. coli* strains that are associated with numerous infections, possess virulence factor (VF) genes. *E. coli* is known to be an important cause of skin and soft-tissue infections (SSTI) as it is the 3rd most prevalent isolated species from SSTI nevertheless, *E. coli* from such infections are rarely investigated.

Objectives

The main goal of our research was to shed more light on certain iron uptake systems among SSTI *E. coli* strains. We determined their prevalence and analysed possible correlations between their genes and phylogenetic groups and some other VFs as well as bacteriocin genes.

Methods

A collection of 102 previously described SSTI *E. coli* strains was used. The methods employed were cultivation in LB medium, preparation of lysates, and polymerase chain reaction (PCR) with primers specific for the investigated target genes (*fyuA, iroCD, iucD, iha, ireA, picU* and *hbp*). The PCR products were detected and visualised with agarose gel electrophoresis. All results were statistically processed using Fischer’s exact test and Bonferroni correction.

Conclusions

The prevalence of iron up-take systems among the studied collection was 76%, 60%, 47%, 30%, 19%, 10% and 5% for *fyuA, iroCD, iucD, iha, ireA, picU* and *hbp*, respectively. The following correlations were observed: *fyuA* with B2 phylogenetic group and *cnf1, hlyA, kpsMTII, ompT* and *usp* genes; *hbp* with *ompT* and *microcin V*; *ireA* with *papGII*; *iha* with *papGII* and *iucD*; and *iroCD* with *microcinH* and *sfa*.
Background

Bacteria from the *Brucella* genus are gram negative intracellular pathogens responsible for Brucellosis, one of the most widespread anthropo-zoonosis worldwide. *Brucella* spp. are intracellular pathogens and despite their tremendous impact on world health and economics no human vaccine is currently available and little is known about molecular mechanisms underlying the infection process.

*Brucella* infection of cultured host cells (i.e. HeLa cells and RAW 264.7 macrophages) is biphasic. It is first characterized by a « lag phase » consisting in a very low but constant number of colony-forming units during the first hours of infection, and followed by a second « duplication phase » during which bacteria proliferate massively. This defines adhesion/invasion of host cells as a critical step for successful infection.

Recent data showed that infection is mainly carried out by a bacterial subpopulation composed of newly generated bacteria called « newborns », characterized by a single genome copy (Deghelt, Mullier *et al*. 2014)

Objectives

Identification of bacterial components mediating preferential adhesion to and/or invasion of cultured host cells by newborn *Brucella*.

Methods

'With *a priori*" gene deletion, 'without *a priori*" transpositional mutagenesis screen (Tn-seq), lectin staining.

Conclusions

So far, three adhesins (BmaC, BtaE, BtaF) have been shown not to act in the newborn selection when infecting HeLa cells and RAW 264.7 macrophages. Tn-seq is still ongoing. We have detected specific binding of the wheat germ agglutinin to the new pole of *B. abortus* when grown until stationary phase in rich medium.
INTERACTION WITH VITRONECTIN, LAMININ AND FIBRONECTIN: A NOVEL ROLE OF Haemophilus influenzae LIPOPROTEIN P4 IN SERUM RESISTANCE AND ADHERENCE TO PULMONARY EPITHELIAL CELLS

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Background Nontypeable Haemophilus influenzae (NTHi) is a human-specific Gram-negative species that primarily colonizes the nasopharyngeal and respiratory tract. Interaction with extracellular matrix (ECM) proteins is one of the colonization strategies described for NTHi, and involves several bacterial adhesins.

Objective We describe herein the identification and functionality characterization of a NTHi multifunctional ECM protein receptor.

Methods Outer membrane proteomic analysis revealed that Haemophilus outer membrane protein 4 (P4; an acid phosphatase involved in the uptake of nicotinamide adenine dinucleotide) interacted with multiple ECM proteins including vitronectin, laminin and fibronectin. This was further validated when an NTHi 3655 mutant devoid of P4 (NTHi3655Δhel) significantly lost binding to the ECM proteins. Binding was restored by a hel gene trans-complemented mutant. Analyses of protein-protein interactions by ELISA revealed that purified recombinant P4 has a high binding affinity for laminin (dissociation constant, KD=9.26 nM) and fibronectin (KD=10.19 nM), but slightly less to vitronectin (KD=16.51 nM). Interestingly, binding to the ECM proteins was optimal with dimeric P4, a structure that is dependent on the α-domain. Vitronectin acquisition conferred serum resistance to both P4-expressing NTHi 3655 and Escherichia coli transformants, but not to the P4-deficient strains. Importantly, NTHi3655Δhel also displayed decreased adherence to both type II alveolar epithelial and pharynx epithelial cells, and blocking with antibodies suggested the P4-mediated cell adherence was mainly attributed to fibronectin.

Conclusions In conclusion, our data provides new insight into the potential role of P4 as a multiple ECM protein receptor important for colonization and establishment of NTHi infection.
PASTEURIELLA PNEUMOTROPICA SEPTICEMIA IN AN ELDERLY WOMAN

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Background
Pasteurella pneumotropica (PP) is a small gram (-) coccobacillus which is a normal inhabitant of the oropharynx of small animals. Human infections due to PP rarely occur. We saw an elderly woman with septicemia due to PP, and describe herein.

Objectives
Human infections due to PP are rarely seen, and it is our intention to share the case experience with others.

Methods
A 76 yo woman with comorbidities including ESRD on hemodialysis, DM, CVA, and hypertension was admitted for evaluation of fever, chills, sore throat and fatigue. She also complained of increased cough with yellowish sputum production. She denied myalgia, arthralgia, and dyspnea. She owns a dog. On exam. temp., 38.9C, RR, 23/min, and PR, 112/min. Other positive findings were: erythematous pharynx without exudate, lung auscultation showed diffuse bilateral rhonchi. Lab. findings: leukocytes, WBC at 31.7K/cmm; BUN 43 mg/dl; and Cr 8.69 mg/dl. Sputum culture grew normal oral flora. Blood cultures (B/C) obtained on admission revealed positive (4/4) for small gram-negative coccobacilli which was confirmed to be PP. Antibiotic susceptibility testing: sensitive to penicillin containing preparations (i.e., ampicillin, amoxicillin/clavulanate), doxycycline, TPM/SMX, and moxifloxacin, while resistant to ceftriaxone and azithromycin. She was initially begun on amp./sulb. and she became afebrile and abacteremic on day 3. She remained abacteremic afterward and was discharged on amox/clav 875 mg po BID to complete a 2-wk course.

Conclusions
A patient with no known history of definite animal bite or scratch developed PP septicemia and she responded to penicillin preparation nicely without further complication.
Background

*Salmonella* infection is a major health concern and continues to have a serious economic impact worldwide. It is estimated that *Salmonella* serotypes cause 93.8 million human infections and 155,000 deaths annually through the world.

Objectives

The main objective of this study was to update the prevalence and antimicrobial resistance characteristics of *Salmonella* isolated from poultry and from humans experiencing gastroenteritis in N'Djamena and also to evaluate the phylogenetic relationships and to find epidemiological links between human and avian strains isolated in the same period of time and in the same region.

Methods

All samples (dropping, sterile Cloths, food and water) collected were analysed according to French Norm for *Salmonella* Spp. NF ISO 6579/2002.

A total of *Salmonella* isolates found in humans and poultry in this study were characterised by ERIC-PCR and IS200-PCR. The PCR analysis was followed by macrorestriction analysis (PFGE).

Conclusions

Diagnostic methods carried out during this study led to the isolation of one hundred and twenty seven *Salmonella* strains, belonging to forty two different serotypes. *Salmonella* Colindale was the most prevalent serovar (13.6 %), followed by *Salmonella* Minnesota (10.8 %), *Salmonella* Stanleyville (5.8 %).

PCR results demonstrated that, in the context of this epidemiological study, ERIC- and IS200-PCR methods can be used effectively to limit the number of isolates that have to be serotyped. The comparison of ERIC-PCR, IS200-PCR, PFGE and antimicrobial susceptibility profiles among isolates from human and avian origins has highlighted, the indirect evidence of human contamination sources by *Salmonella* serovars from poultry farms.
Background

Gram-positive bacteria release lipoteichoic acid (LTA) as an important virulence factor, which is structurally and immunologically similar to lipopolysaccharide (LPS). The knowledge about LTA actions in vivo remains unclear, as well as their participation in the etiology of pulp and periapical diseases, with no reported studies on the quantification of LTA in root canals.

Objectives

a) quantifying the levels of LTA in primary infections of root canals with necrotic pulp and apical periodontitis; b) verifying the effects of biomechanical preparation using sodium hypochlorite 2.5% in LTA reduction in root canals.

Methods

Ten single-rooted teeth of patients needing endodontic treatment were prepared with Reciproc rotation system and irrigated with 2.5% NaOCl. Root canals samples were obtained immediately after coronary opening (S1) and after biomechanical preparation (S2). LTA quantification was performed by enzyme immunoassay (ELISA), with specific anti-LTA antibodies. The optical density values were converted to µg/mL and analyzed by Student’s t test (significance level of 5%).

Conclusions

Results: All root canals showed LTA in the collection S1 with an average of 119.4 ± 73.2 µg/mL. After instrumentation, it was observed a reduction in the levels of LTA, with an average in the collection S2: 77.6 ± 22.4 µg/mL.

Conclusion: The root canals with necrotic pulp showed high levels of LTA and only instrumentation with sodium hypochlorite 2.5% was not enough for its entirely
elimination of the root canals. New studies with other endodontic treatment protocols should be performed for new findings.
Background
Leptospirosis is a worldwide zoonosis regarded as a major public health problem. Measures to control the disease are difficult to implement. The development of new strategies to prevent and control the spread of disease is urgently needed. Accordingly, vaccines emerge as strong candidates to solve the problem. For this reason, currently research has focused to identify conserved antigens that are involved in host-pathogen interactions.

Objectives
Evaluate the functional properties of the coding sequence LIC13479 and LIC10050 of L.interrogans serovar Copenhageni, identified by bioinformatics as putative outer membrane proteins.

Methods
The gene sequences were cloned into the expression vector pAE. Plasmids containing cloned DNA were introduced in E. coli strains for protein expression. After purification of the recominant proteins, mice were immunized for polyclonal antibodies production. Reactivity of the recombinant proteins was evaluated in serum samples of leptospirosis patients and of febrile unrelated diseases. The ability of recombinant proteins to interact with the host by adhering to extracellular matrix proteins or serum components was examined.

Conclusions
The coding sequences LIC13479 and LIC10050 were cloned and expressed successfully. Evaluation of the purified recombinant proteins showed that they are capable to stimulate antibody immune response in mice and, in addition, they are recognized by infected human serum samples. Both recombinant proteins exhibited adhesin properties and, in addition, interacted with plasminogen and can generate plasmin in the presence of activator. Our data indicate that these proteins could promote the attachment and contribute to the invasion processes within the hosts.
Background

STEC-associated HUS was first recognized in Georgia in 2009 following the diagnosis of a cluster of HUS cases in Georgia.

Objectives

In this study we genetically characterized two sporadic cases (Case A and B) of HUS syndrome occurred in 2012. Among them was one family case reported (Case B), where four individuals revealed positive on STEC, but only one developed HUS.

Methods

Clinical samples were enriched in broth and simultaneously plated on selective agar media. DNA was extracted and tested by two different conventional Multiplex PCR assays for final confirmation of STEC (stx1, stx2, eae, Ethyl) and detection O104 strain markers (stx2, terD, rfbO104, fliC H4). Furthermore, for molecular genotyping Pulsed Field Gel Electrophoresis (PFG E) was applied. Five STEC strains were isolated from both described cases. All revealed positive results by molecular tests on STEC but showed different genetic profiles based on combination of existing
toxigenic markers. Interestingly, it was observed that case A tested positive on all four O104 specific genes. Besides, PFGE typing showed two different genetic profiles as well.

Conclusions

Here described in both HUS cases, a diversity of STEC isolates was detected. However, one of the causative agents was considered to be STEC O104. It is remarkable that two O104 strains were isolated during STEC outbreak in Georgia in 2009 which are genetically related with the strains isolated during STEC outbreak in Germany in 2011. This finding suggests that O104 strain has spread and been circulating for several years in the country.
Background

The biosynthetic pathways of several siderophores and the genotoxin colibactin require the enzymatic activity of a phosphopantetheiny transferase (PPTase). We have shown that ClbA, the PPTase associated with the production of colibactin, can also contribute to the production of siderophores, whose production is dependent on the bioavailability of iron in the medium, through the regulation of the expression of EntD, the PPTase previously thought to be the unique PPTase involved in the production of siderophores in E. coli.

Objectives

We hypothesized that iron could also modulate the production of colibactin through the transcriptional regulation of clbA.

Methods

Mutant derivatives of E. coli were engineered. We investigated transcription of genes involved in the production of colibactin in iron depleted or repleted conditions using qRT-PCR and luciferase reporter gene assay. Quantification of total siderophores was assessed.

Conclusions

This study revealed an increased transcription of clbA in iron limiting conditions, and a decrease of clbA expression in iron supplemented media. Quantification of total siderophores revealed that the increased expression of clbA in iron-depleted conditions was associated to an increased synthesis of siderophores. Mutation of entD leads to an exacerbation of the induction of clbA expression, whereas overexpression of entD or clbA leads to a repression of clbA transcription. Analysis of the region upstream clbA shows the presence of two putative Fur-boxes, which could explain its regulation by iron. In conclusion clbA is tightly regulated by iron.
bioavailability leading to an increased production of siderophores that could explain the high pathogenicity of the strains synthetizing colibactin.
FEMS-0572

Bacterial pathogenicity

VIRULENCE FACTORS OF STREPTOCOCCUS PYOGENES ISOLATED FROM PATIENTS WITH TONSILLOPHARYNGITIS IN KONYA/TURKEY

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Background

*Streptococcus pyogenes* is an important bacterial pathogen that colonizes the throat and skin of a host and produces a wide variety of virulence factors such as toxins, proteases or DNases.

Objectives

The aim of this study was to investigate the antibiotic resistance and virulence factors (speA, B, C, G, H, I, J, K, L, M, smeZ, ssa, spd3, sdc, sdaB, sdaD, spyCEP, scpA, mac and sic) of *S. pyogenes* strains isolated from throat cultures of patients with symptomatic tonsillopharyngitis.

Methods

One hundred and fifty *S. pyogenes* isolates were identified by conventional methods and VITEK 2 automated system. Antibiotic susceptibility tests were performed by Kirby-Bauer disk diffusion method as recommended by Clinical and Laboratory Standards Institute. The virulence factors were determined by multiplex PCR.

Conclusions

All of the *S. pyogenes* isolates were susceptible to penicillin G, cefotaxime, ceftriaxone, chloramphenicol, clindamycin, erythromycin, levofloxacin, vancomycin and linezolid. Among streptococcal pyrogenic exotoxin genes the most frequent gene was speG (88.0%) following speC (59.3%), smeZ (46.7%), ssa (42.7%), speA (33.3%), speJ (24.0%), speK (18.7%), speH (14.0%), speI (13.3%), speL and speM (9.3%). From DNases, proteases and inhibitors sdaB, speB, spyCEP, scpA and mac were positive in all strains, and spd3, sdc, sdaD and sic were carried 64.7%, 36.0%, 24.7%, 2.0% of the isolates respectively. In conclusion, *S. pyogenes* isolates collected from throat cultures of patients with symptomatic tonsillopharyngitis in Konya/Turkey possess high virulence factors, and susceptible to antibiotics.
Background
Type 3 secretion system (T3SS), is major virulent factor in over twenty species of Gram-negative bacteria which export bacterial protein (effectors) to host cell to manipulate its cellular function for own's hope. Moreover, T3SS is a prospective useful tool with both experimental and therapeutic applications, including vaccine development.

Objectives
Upon analysis of the contribution of the T3SS injectisome and effector involved in this paradigm, it is still unknown that the number and timing of expression of the T3SS injectisome and effectors, and which factor have dominant role during infection are unclear. In the present study, we used fluorescence protein combined with single cell analysis to solve dynamics of the T3SS and effector during infection in Vibrio parahaemolyticus.

Methods
During infection, Vp1671 (T3SS component) but not the Vp1680 (effector) proteins showed to make spot like localization along the membrane of the cell. Although both percentages of spot positive cells and number of T3SS spots were increased depend on a time of infection, assemblies of the Vp1671 to the T3SS occur prior to increase fluorescence levels of bacterium in early time of infection. Cytotoxicity against the host cells were increased depend on the expression levels of Vp1680 and number of T3SS spot, if it is only one spot.

Conclusions
V. parahaemolyticus use a combination between a few T3SS injectisome regulated by localization and a lot effector regulated by expression for killing host cells.
Background

*Streptococcus agalactiae* causes urinary tract infection (UTI) including asymptomatic bacteriuria (ABU); however, growth of *S. agalactiae* in urine has not been reported and the role of this growth phenotype in human infection is unknown.

Objectives

We characterised the ability of different *S. agalactiae* UTI isolates to grow in urine, and defined the molecular mechanism(s) of growth and implications for human ABU.

Methods

We used growth assays, competition assays, metabolic phenotype arrays, whole-genome sequencing, and mutation analysis in *S. agalactiae*. We report robust growth of ABU *S. agalactiae* (ABSA) in human urine that was not seen among uropathogenic *S. agalactiae* (UPSA) isolated from patients with acute UTI. In competition assays using a prototype ABSA strain, designated ABSA 1014, and any one of several UPSA strains, we observed markedly superior fitness of ABSA 1014 for urine growth. Phenotype profiling of ABSA 1014 and UPSA 807, isolated from a patient with cystitis, using metabolic arrays revealed specific L-Malic acid catabolism in ABSA 1014 that was absent in UPSA 807. Whole-genome sequencing revealed divergence in malic enzyme-encoding genes between the strains predicted to impact the activity of malate metabolism. Urine growth assays comparing wild-type ABSA and mutants that were functionally inactivated for malate metabolism by disruption of *maeE* or *maeK* demonstrated attenuated growth of the mutants in urine and synthetic human urine containing malic acid.

Conclusions

We conclude that some ABU *S. agalactiae* can grow in urine, and this relates in part to malic acid metabolism, which may affect *S. agalactiae* UTI.
THE ROLE OF THE MYCOSIN PROTEASE IN TYPE VII SECRETION OF PATHOGENIC MYCOBACTERIA.

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Background

A promising new group of novel drug targets for Mycobacterium tuberculosis are the type VII secretion (T7S) systems. M. tuberculosis has five of these systems, ESX-1 to ESX-5, of which at least three are crucial for the virulence and/or viability of this pathogen. One of the conserved components of T7S systems is mycosin (MycP), a membrane-associated protease that is an essential part of the T7S system. Due to the proteolytic activity of the mycosins they are promising targets for novel drugs against TB.

Objectives

We set out to functionally dissect the role of the mycosin proteases in T7S, by analyzing the phenotype of MycP1 and MycP5 mutants.

Methods

Deletion strains of the mycosins of the ESX-1 and ESX-5 secretion systems were created in Mycobacterium marinum. Subsequently, proteolytic inactive versions of MycP1 or MycP5 were introduced in the deletion strains. Functional complementation by these variants was verified by the effect on the secretion by the ESX-1 and ESX-5 systems.

Conclusions

While deletion strains of MycP1 or MycP5 were defective in respectively ESX-1 or ESX-5 dependent secretion, the protease inactive variants were able to mediate secretion by ESX-1 or ESX-5. Thus the proteolytic activity of MycP is not essential for its respective ESX-system dependent secretion. These results indicate a dual function for mycosins, with a proteolytic role in substrate processing and a second, so-far unknown, role in the regulation of the secretion process. We are currently
unravelling this second function in T7S and pinpointing which domain of MycP is involved in this process.
FEMS-1868
Bacterial pathogenicity

EFFECT OF DIFFERENT FACTORS ON THE (TWITCHING) MOTILITY OF PHOTOBACTERIUM DAMSELAE SUBSP. PISCICIDA.

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Background

Photobacterium damselae subsp. piscicida, which is the etiological agent of photobacteriosis and affects different species of sea fish, is a non-motile Gram-negative bacterium. Notwithstanding, it is able to move on solid surfaces. Because of the presence of pili-like structures on the bacterial surface, discovered the last year by Remuzgo-Martínez et al., and being associated with twitching motility,

Objectives

the objective of this study was to modify some conditions (temperature, pH, concentration of nutrients, salinity, nature of the medium and others) to determine which ones promote motility.

Methods

A total of five bacterial strains were tested: C2, 94/99, DI21, PP3 and ATCC 17911. The results of motility assays were obtained after two and three days of inoculation in “twitching motility” medium and statistically analyzed with IBM SPSS Statistics 22.0 program.

Conclusions

Lower temperatures, alkaline and acidic environments and lacking salinity had a negative effect on cell motility, while nutrient limitation did not affect the bacterial response. The best twitching motility was observed when Phdp was inoculated in medium with 0.2% agar and pH 7.0. Also, interesting results were achieved after scraping petri dishes (surface where the bacteria moved) or the addition of cellular debris of SAF-1 cell line to the medium.
Bacterial pathogenicity

IDENTIFICATION AND DISTRIBUTION OF A NOVEL 22-KDA OUTER MEMBRANE PROTEIN RELATED TO SHIGA TOXIN–PRODUCING ESCHERICHIA COLI (STEC) STRAINS NEGATIVE FOR LOCUS ENTEROCYTE EFFACEMENT.

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Background
STEC strains are etiologic agents of acute diarrhea, bloody diarrhea and hemolytic uremic syndrome (HUS). Associated with STEC outbreaks and HUS, we may observe two main groups of strains, those that in their genomes is encoded the Locus Enterocyte Effacement (LEE-positive), including the serogroups, O157:H7, O26, O103 and O111, and bacteria without this Locus (LEE-negative) which include the serogroups, O113 and O91: H2.

Objectives
Describe a novel antigen present in the outer membrane (OM) of a LEE-negative STEC strain.

Methods
By western blot and immunoproteomic analysis (2D electrophoresis - MALDI-TOF/TOF) using sera from patients who developed HUS, we were able to identify an immunoreactive protein of 22 kDa with an isoelectric point of 5.0. By mean of bioinformatics tools, we found the encoded gene in the genome of STEC strain O91: H21 B2F1 LEE-negative (Accession: AFDQ0100026.1). Using specific primers, the gene was amplified and its presence was studied in a collection of 170 STEC strains and 11 commensal E.coli. Interestingly, the gene was detected only in LEE-negative STEC strains (63%; 32/51). Sequencing of PCR products showed that this gene is highly conserved. Preliminary results suggest that this protein could be involved in bacterial adhesion and hemagglutination.

Conclusions
Our results suggest that the identified antigen might be associated only with a subset of LEE-negative STEC strains. Additional studies are needed to determine the role of this protein in STEC pathogenicity.

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LEPTOSPIRA INTERROGANS STIMULATES PLASMINOGEN ACTIVATORS IN VIVO IN EXPERIMENTALLY INFECTED ANIMALS AND NATURALLY ACQUIRED HUMAN LEPTOSPIROSIS.

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Background
Leptospirosis is a globally important zoonosis, caused by pathogenic bacteria of genus Leptospira. The molecular mechanisms for leptospiral pathogenesis and virulence remain poorly understood. A pathogen invasion, dissemination and host tissue damaging depend on proteolytic enzymes both of the invading organism and of the host. Many pathogens express their own proteases or exploit hosts proteases to activate other protease-dependent cascade systems. We have described that leptospires capture plasminogen (PLG) on the outer surface, which is converted to plasmin (PLA) by exogenous activators. We also showed that leptospires induce the expression of PLG activators by human endothelial cells in vitro.

Objectives
Here, we further characterized the interactions of L. interrogans with human PLG/PLA system in vivo.

Methods
We studied the stimulation of PLG activators in different time points during leptospirosis in hamsters infected with lethal doses, and in human sera samples from the initial and convalescent phase of the disease.

Conclusions
Our data indicate that leptospirosis human patients have increased levels of circulating urokinase-type and tissue-type PLG activators when compared to normal sera, being especially augmented at the early phase of the disease. In the sera of experimentally infected hamsters, increasing levels of PLG activators were observed during the progression of the infection, until euthanasia. Additionally, hamsters’ organs extracts similarly showed increased PLG activators activity.

The results presented here further characterize the host response to the Leptospira infection, in the light of PLG/PLA system stimulation. Our data strengthen the importance of the fibrinolytic system to the leptospirosis infectious process.

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FEMS-1750
Bacterial pathogenicity

BINDING OF LEPTOSPIRA INTERROGANS TO HUMAN THROMBIN, BIOLOGICAL IMPLICATIONS AND IDENTIFICATION OF POSSIBLE BACTERIAL LIGANDS
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Background
Leptospirosis is a zoonosis caused by pathogenic spirochete of genus Leptospira. The symptoms range from a mild fever to a more complicated, severe form of the disease, called Weil’s syndrome. Since the 1980s, the incidence of severe pulmonary hemorrhage caused by Leptospira spp. infection has increased. The reasons behind the emergence of this syndrome are not known. Many pathogens can interfere in the coagulation cascade by binding to fibrinogen or by interaction with the enzyme thrombin.

Objectives
To characterize the interaction of leptospiral strains to thrombin, analyze the possible biological relevance and select possible proteins that could mediate these interactions.

Methods
The interaction of pathogenic (virulent and attenuated) and saprophytic strains with thrombin was performed by ELISA and western blotting. Characterization of the binding sites in thrombin molecule was accessed by co-incubation with different competitors. Reduction in thrombin activity was evaluated by fibrin clot formation assay. Identification of ligands was performed by ELISA by employing a set of recombinant leptospiral proteins.

Conclusions
We demonstrated that the pathogenic strains of Leptospira display a prominent binding to thrombin, particularly the virulent one. Competition assays indicate that virulent strain binds thrombin via exosite I. Virulent strain could reduce thrombin activity. Although we have identified some proteins that displayed binding to thrombin, none was capable to reduce the enzymatic activity. The interaction of virulent strains of Leptospira with the key enzyme of the coagulation cascade, thrombin, might constitute a novel mechanism of virulence, that could help understand the hemorrhagic features of leptospirosis.
RELATION BETWEEN THE SEQUENCE TYPE AND THE PROSPECTIVE VIRULENCE OF CRONOBACTER STRAINS

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Background

Bacteria from the genus \textit{Cronobacter} are pathogens endangering all age groups, with severe clinical complications especially in neonates and infants. A multilocus sequence typing (MLST) scheme based on 7 housekeeping genes was constructed for \textit{Cronobacter} spp. identification. The method also enables the recognition of sequence types (STs) and clonal complexes (CCs). Although 7 housekeeping genes (\textit{atpD, fusA, glnS, gltB, gyrB, infB, and ppsA}) used for MLST analysis are not virulence related, connection between infections and particular sequence types were described in the literature.

Objectives

The aim of this work was to evaluate the relation between sources of tested strains and their characteristics particularly their sequence type.

Methods

MLST aimed to 7 genes (\textit{atpD, fusA, glnS, gltB, gyrB, infB, and ppsA}) was performed on bacterial collection (over 80 strains). The amplification of genes \textit{gltB} and \textit{gyrB} showed to be difficult, and therefore new sets of primers were designed and tested.

Conclusions

The target genes were sequenced in the majority of tested strains; in the case of incomplete information putative sequence type was determined. The obtained results showed widespread distribution of STs within tested strains. The significant predominance of particular STs was not determined in tested strains. Perhaps, the amount of tested strains was not representative enough; the ST relation to virulence will be further studied together with other possible factors.
This work was supported by the Czech Grant Agency (13-23509S) and Specific University Research (MSMT No. /2015).
Background

*Staphylococcus aureus* is an opportunistic pathogen of humans and animals. Our studies have revealed a link between the genotype of poultry-isolated *S. aureus* strains and their virulence in chicken embryo but not nematode model, what may suggest the existence of factors responsible for host preference and virulence.

Objectives

Here we compare genomes and extracellular proteomes of four virulent and four non-virulent *S. aureus* strains in chicken embryo model. The open reading frames translation products for all strains were taken together and clustered according to the similarity level of 80%. Over three thousands clusters were obtained with only around 40% of clusters containing proteins with identical sequence across all strains. However, within the virulent and non-virulent group around 80% and 54% of clusters ORFs encoded identical proteins, respectively. The exoproteomes were highly heterogeneous irrespective virulence, however we were able to point alpha-hemolysin and bifunctional autolysin as indicators of virulence whereas glutamylendopeptidase as potential virulence attenuator.

Methods

The genomes were obtained using MiSeq Illumina sequencer and assembled with MIRA and CLC Main Workbench software. Extracellular proteins were precipitated, labeled with fluorescent dyes and subjected to two-dimensional difference gel electrophoresis in pairs virulent *versus* non-virulent. Differentiating protein spots were subjected to mass spectrometry analysis.

Conclusions

Deep sequencing confirmed genetic dissimilarity of virulent and non-virulent strains which finds its reflection in extracellular proteome. However, proteomics also points transcriptional and translational events as modulators of *S. aureus* virulence.

The study was supported by the grant UMO-2012/07/D/NZ2/04282 (to B.W.) from National Science Centre, Poland.
ADHESINS FROM YERSINIA RUCKERI

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Background

Yersinia ruckeri is a Gram-negative bacterium pathogen responsible for enteric redmouth disease (ERM). Yersinia ruckeri infections cause significant economic losses in salmonid aquaculture. ERM can be effectively controlled by antibiotic treatment and by application of whole cell based vaccines. Nevertheless, ERM outbreaks are still observed. More recently, vaccine resistant isolates caused disease throughout Europe. Research into the pathogenicity of Yersinia ruckeri is limited. Much more is known about the host-pathogen interaction of the three Yersinia species pathogenic to humans. The group of proteins, that mediate the host-pathogen interaction comprise adhesins anchored in the bacterial outer membrane. They are essential for infection as they possess the ability to bind a variety of host molecules such as collagen, fibronectin, laminin, β1 integrins.

Objectives

The aim of my research is to establish a fish infection model – zebrafish to understand the molecular mechanisms of adhesion and to identify adhesins of interest and their cellular receptors in order to assess their function in pathogenesis.

Methods

To examine whether adhesins of interest are involved in Yersinia ruckeri virulence, fish cell line and in vivo system (zebrafish embryos) will be used and optimise. Fluorescence microscopy will allow to visualize infections to assess the role of virulence genes. Infection of colour- fluorescence bacteria will be essential to follow the infection.

Conclusions
Understanding the molecular mechanisms of adhesion in fish, might prevent progression of infections. Moreover, the knowledge gained could help in developing new antimicrobials that could directly act as vaccines against bacterial fish diseases.
MOLECULAR CHARACTERISTICS OF BRUCELLA ABORTUS MUTANTS GENERATED WITH EZ-TN5TM PMODTM-3 TRANSPOSON

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Background
Brucella abortus is a well-known intracellular pathogen. Underlying mechanisms of the bacterial infection is very complicated to understand the pathogenesis even though most of clues have been solved. Therefore, prevention and control of the B. abortus infection is still problematic in animal and human. So, several methods including random mutation have been used to understand the mechanisms and find out the solution in control and prevention of the infection. Regardless the trials, the doors to be opened are still locked in understanding of the mechanism.

Objectives
To understand the underlying mechanism of B. abortus infection, mutants were generated with a transposon and function of the disrupted genes were revealed.

Methods
Mutants of B. abortus were generated by random insertion of a transposon, Ez-Tn5™ pMOD™-3 <R6Kyor/MCS> into chromosome. Molecular characteristics of the mutants were investigated using PFGE, Southern blot and sequencing of the genes.

Conclusions
B. abortus mutants were generated and insertion of the transposon was confirmed by Southern blot analysis with the transposon as a probe after PFGE of chromosome. Both sides of insert in the chromosome were sequenced and the location of interrupted genes were identified in chromosome. Using sequencing information, twenty-eight genes were identified from the mutants by comparison with wild type of the bacteria. Also, function of the genes were revealed. This work was supported by NRF (No. 2014R1A2A2A01007291).
FEMS-2498
Bacterial pathogenicity

CYTOKINE EXPRESSION PROFILE IN RAW 264.7 CELLS STIMULATED WITH BRUCELLA ABORTUS MUTANTS
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Background
Brucella abortus have been known as an important intracellular, zoonotic pathogen. Intestinal immune responses are induced in animals by the bacterial infection. Cytokine has key roles in activation of innate immunity and acquired immunity. Precious roles of cellular components of B. abortus infection have been remaining to be solved even though involvements of immune cells in activates of acquired immunity are already known. On the basis of current understanding, production profiles of inflammation cytokines such as NO, IL-1β, IL-4, IL-12 and IFN-γ, were compared with Brucella mutants. Moreover, it might be given a key to find immune system, pathological pathways, and mechanism.

Objectives
To understand the roles of cellular components of B. abortus in immune cells, production profiles of cytokines were investigated in a mouse macrophage cells, RAW264.7 cells after stimulation with B. abortus mutants.

Methods
RAW 264.7, murine macrophages, cells were stimulated with mutants of B. abortus. Culture supernatants were collected from RAW 264.7 cells after stimulation with B. abortus mutants at different time intervals. Amounts of cytokine such as NO, IL-1β, IL-4, IL-12 and IFN-γ, were quantified with ELISA.

Conclusions
Inflammatory cytokines were produced from Raw 264.7, murine macrophage, cells after stimulation with the B. abortus mutants. After then, production levels of inflammatory cytokines such as NO, IL-1β, IL-4, IL-12 and IFN-γ, were measured. The patterns of the cytokine productions will be presented. This work supported by NRF (No.2014R1A2A201007291) and IPET112012-03-1-HD020.
FUNCTIONAL INSIGHTS INTO THE ROLES OF AUTOTRANSPORTER ADHESINS IN BRUCELLA

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Background

\textit{Brucella} pathogens proliferate within several eukaryotic cells. Cumulative evidence indicates that adhesion of \textit{Brucella} to host cells is an important step to establish infection. We have shown that the unipolar monomeric autotransporter BmaC is involved in the adhesion of \textit{Brucella suis} to host cells through binding to cell-associated fibronectin.

Objectives

Genome analysis shows that the \textit{B. suis} genome encodes two trimeric autotransporters (BtaE and BtaF) and two additional monomeric autotransporters (BmaA and BmaB). The aim of this work was to explore the roles of these proteins in the interaction of \textit{Brucella} with the host.

Methods

Clean deletion mutants were analyzed by in vitro binding assays and in vivo using the mouse model. Adhesin localization was analyzed by immunofluorescence.

Conclusions

BtaE was involved in the binding of \textit{B. suis} to hyaluronic acid and fibronectin while BtaF may contribute to the binding to several ligands. Both the $\Delta$bta\textit{E} and $\Delta$bta\textit{F} mutants showed a reduction in the attachment to epithelial cells and were attenuated in mice. Similar to BmaC, BtaE and BtaF were detected in a low proportion of bacteria but in all cases at one particular pole. A remarkable variability in the number of motifs of autotransporter orthologues from different \textit{Brucella} species was observed.
Furthermore, while both trimeric autotransporters might be functional in all *Brucella* species, pseudogenization occurred in *bmaA* and *bmaB* orthologues. In particular, BmaA would not be functional in *Brucella abortus* while it is predicted to be functional in *B. suis* and *B. melitensis*. Thus, functional autotransporters may contribute to host preference.
Background

*Bifidobacterium breve* is a common inhabitant of the infant gut and its presence has been correlated with particular health-promoting effects, such as enforcement of the intestinal barrier, activation/modulation of the host’s immune response and protection against particular infections.

Objectives

Genome sequencing of *B. breve* JCM 7017 revealed the presence of the first verified bifidobacterial megaplasmid pMP7017 (190 kb of size). The objective of this study was to provide evidence of its horizontal transmission between strains of the same and/or different (bifido)bacterial species by means of a predicted conjugative machinery.

Methods

DNA sequencing was performed using Next Generation Sequencing platforms. The *in silico* predictions were performed using BLAST and PFAM databases as well as additional bioinformatics tools (e.g. prodigal, tRNAscan, CRISPR and GCUA). Proof of conjugal transfer was obtained by demonstrating the presence of the megaplasmid using Pulsed Field Gel Electrophoresis in combination with S1 nuclease treatment.

Conclusions

*In silico* characterization of the megaplasmid revealed several genomic features supporting a stable establishment in its host, illustrated by predicted CRISPR-Cas functions that are known to protect the host against intrusion of foreign DNA. Interestingly, pMP7017 is also predicted to encode a conjugative DNA transfer apparatus and consistent with this notion we demonstrate conjugal transfer of pMP7017 to representative strains of *B. breve* and *B. longum* subsp. *longum*. We furthermore demonstrate the presence of a megaplasmid with homology to pMP7017 in three *B. longum* subsp. *longum* strains, indicating that similar elements are also naturally present in other bifidobacterial species, thereby supporting our proof of cross-species transfer.
EXPLORING THE INFLUENCE OF CHEMOTACTIC PREFERENCES ON SUBSTRATE COMPETITION AND COMMUNITY FATE BY IN SILICO MODELING

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Background

Competition for substrate is a common mode of microbial interaction in natural environments. Both, growth and motility phenotype influence competition, with the former having been studied well, while the latter has received less attention.

Objectives

To elucidate the impact of microbial motility on competition, we focus on microbial communities in which two strains populating a homogeneous environment compete for a substrate. Strains share identical growth phenotypes but differ in their chemotactic preference, either responding more sensitively to substrate or to a chemoattractant that is excreted by the cells themselves. By systematically varying chemotactic preferences of the competing strains, we assess the impact of each chemotactic preference on competition and on overall community fate.

Methods

Microbial communities are simulated using an individual-based modelling approach. Competing strains are initially homogeneously populating a two-dimensional environment. Microbial growth and motility including random movement and chemotaxis towards substrate and microbial cells, and substrate diffusion are simulated, and the resulting spatio-temporal microbial distribution patterns are observed and analyzed.

Conclusions

Results show that, depending on the combination of chemotactic preferences, microbial distributions remained homogeneous or transient or permanent spatial patterns emerged. The presence of a competing strain is able either to suppress or to induce aggregate formation, indicating more complex interactions beyond simple substrate competition. Strains being more attracted to self-excreted compounds tend to form aggregates of high densities where starvation conditions prevail, leading communities to become dominated by the competing strain. The model indicates a mechanism by which aggregate formation can evolve, even if it is initially disadvantageous.
Background

One of the most important aims in ecology is to identify and comprehend the mechanisms that sustain biodiversity - often critically important for ecosystem viability. Many theoretical models have shown that species in competition can coexist - thus maintaining the ecosystem's biodiversity - if ecological processes such as competition and mobility occur on a local scale [1]. This is also true in the case of communities with non-transitive competition between species, the classical example being the rock-paper-scissors game.

Objectives

Initial community evenness has been shown to be a key factor in preserving ecosystem functional stability [2], but has not been accounted for in previous modelling studies. We formulate a model that allows initial community evenness to be varied in order to investigate the consequent impact on ecosystem biodiversity.

Methods

We consider an ecosystem of four interacting bacterial species, and present a stochastic, spatial individual-based model simulating the ecosystem dynamics. Interactions take place on a two-dimensional lattice. Three processes are incorporated: reproduction, competition and mobility. In addition to variable initial evenness, multiple competition schemes are implemented, modelling various possible communities, resulting in diverse coexistence and extinction scenarios.

Conclusions

Simulations show that long-term ecosystem behaviour is strongly dependent on initial evenness and competition structure. The ecosystem is generally unstable; higher initial evenness has a small stabilizing effect on ecosystem dynamics by extending the time until the first extinction.

Background
Antibiotic activity is assessed in vitro by quantifying the reduction of growth of a bacterial culture. Growth can be described as change in cell number on population level. On the single-cell level, physiological properties like cell size before division or translational capacity vary with growth rates: Cells adapt to their environment. Typical drug-effect models link population growth directly to the exposure of antibiotics. They rarely account for known mechanisms of action of the drug - which are particularly relevant for the analysis of synergistic or antagonistic effects of drug combinations.

Objectives
Development of a pharmacodynamic model which allows for mechanistic integration of antimicrobial effects on the cellular level to predict bacterial growth.

Methods
Control experiments without drug resulted in baseline values for population growth. An established single cell model predicted cell level parameters from this growth rate. Induction of cell killing and growth inhibition were linked to known drug characteristics (minimal bactericidal / inhibitory concentration). A transit compartment cell-cycle model integrated time dependent cellular adaptation processes. Two bacterial subpopulations were considered (resting and growing). We validated our model for predicting time-kill curves with E. coli exposed to tetracycline. Furthermore we predicted septation dynamics of B. subtilis during a shift-down from exponential to stationary phase.

Conclusions
Since both scenarios show good agreement between predicted and experimental data, these promising results are a first step to mechanistically model bacterial growth during exposure to multiple antibiotics.
Background

*E. coli* DacD is a DD-carboxypeptidase, having much weaker DD-CPase activity than PBP5. The reason for such a weak DD-CPase activity of DacD and its importance within an *E. coli* cell is unknown.

Objectives

Deciphering the role of active-site groove volume (AGV) on DD-CPase activity of *E. coli* DacD.

Methods

Here, we generated the 3D model of DacD using homology modeling and analysed the binding of specific substrates within AGV. Interestingly, the AGV of DacD is about one-third of PBP5 and such a low AGV might reduce the catalytic efficiency of DacD. To validate the result *in silico*, we mutated the secondary amino-acid residues around the DacD active-site by incorporating several transitional and transversional mutations and observed for the changes developed in the AGV. Such mutations brought a 70% increase in the mutant DacD AGV. PBP5 has higher AGV that can be correlated with DD-CPase activity. In addition, we superimposed PBP5 active-site onto mutated DacD and observed a similar geometric arrangement. Furthermore, the pentapeptide-binding affinity of the mutant DacD was increased as revealed by molecular docking analyses. Higher binding affinity of the mutant DacD towards peptidoglycan mimetic pentapeptide substrate possibly signifies a greater DD-CPase activity.

Conclusions

Active-site groove volume may act as a determinant for DD-CPase activity and the reduced DD-CPase activity of DacD is possibly due to its inadequate active-site groove volume.
Background

The most abundant periplasmic protein of *Vibrio cholerae* grown under inorganic phosphate (Pi) limitation is the high affinity Pi transporter, PstS, of the Pst2 system. It is encoded by *pstS* of the *pst2* operon, *pstSABC*, member of *V. cholerae’s* Pho regulon. Little is known about the regulation of *pst2* expression and the non-stoichiometric production of its genes products.

Objectives

To elucidate the regulatory mechanisms behind *pst2* expression in *V. cholerae*.

Methods

Bioinformatics analysis: Clustal Omega for protein sequences alignment, Mfold for the determination of RNA secondary structures and the corresponding free energy values and MEME/MAST to search for putative Pho boxes. Experimental procedures: SDS-PAGE and MS ESI-Q-Tof Micro for protein analysis and identification and promoter-*lacZ* fusion, RNA extraction, cDNA synthesis, endpoint-PCR and quantitative Real-Time PCR for the expression analysis.

Conclusions

Based on the bioinformatics analysis we proposed new start codons for *pstS* and *pstC*, a regulatory region upstream *pstS* containing potential Pho boxes and a *pstS-pstC* intergenic region, distinct from those predicted. Moreover, sequences able to form stem–loop structures followed by potential RNAse E-cleavage sites were detected in the intergenic regions *pstA-pstB* and *pstS-pstC*. The experimental analyses demonstrated that *pst2* is transcribed under Pi limitation in a PhoB-dependent manner as a full-length mRNA that is processed into minor transcripts of distinct stabilities. The most stable is the *pstS*-encoding mRNA, followed by *pstB*, *pstA* and *pstC* specific transcripts. The abundance of PstS relative to the other components of Pst2 system in *V. cholerae* seems to correlate to the higher stability of its transcript.
Background
The world’s microbial diversity is vast and the majority of microbes do not readily grow in culture, complicating the analysis and interpretation of this diversity. In addition, the lack of a unified and consistent annotation of the world’s microbial diversity has so far impeded a comprehensive comparative analysis.

Objectives
The “Microbe Atlas”, a global metaresource of all publicly available environmental marker gene sequence data, which will be available as a web resource. A dataset consisting of 212’000 samples with more than 3 billion identified small subunit ribosomal RNA sequences. This resource will enable a better understanding of what are the typical and atypical microbial taxa in samples, by establishing compositional baselines in terms of microbes that are often found in similar environments.

Methods
An analysis of all publicly available environmental marker gene sequence data.

Conclusions
This resource should become an essential tool for microbial biologists wishing to compare their samples to previously sequenced microbial communities, and identify both the typical as well as atypical microbial taxa in their samples.
LACCASE FROM PATHOGENIC GUT BACTERIA FACILITATE BIOTRANSFORMATION OF NON-STERoidal ANTI-INFLAMMATORY DRUGS (NSAIDs)

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Background
Non-steroidal anti-inflammatory drugs (NSAIDs) are used in the treatment of pain, inflammation, cardiovascular diseases, arthritis and cancer by inhibiting cell proliferation. Laccases (EC 1.10.3.2) are multicopper oxidases (MCO), recently reported from Escherichia coli, Salmonella enterica, Campylobacter jejuni, Pseudomonas aeruginosa with role in copper homeostasis, iron acquisition, manganese oxidation and in pathogenesis.

Objectives
To find out novel laccase from different serotypes of Yersinia enterocolitica and clone it into a suitable expression vector for the biotransformation of NSAIDs.

Methods
Amplified laccase gene (yack) from different serotypes of Y. enterocolitica was cloned in pTZ57R/T vector using TA cloning method. The positive clones were sequenced to confirm yack and construct a phylogenetic tree using Mega 5.0. Further, yack was subcloned into the desired restriction sites of pET28a expression vector and transformed into E. coli BL21 (DE3) expression host. The recombinant protein was refolded and purified using His-spin protein miniprep kit, for the biotransformation of aspirin and diclofenac (NSAIDs), which validates our in silico studies.

Conclusions
Laccase from different strains of Y. enterocolitica can be used as a taxonomic tool as it showed serotype specific clades. The HPLC analysis of laccase treated diclofenac and aspirin showed complete biotransformation after 24h of incubation. The FTIR results suggested a change in C-C and C-N bonds. Laccase transform NSAIDs and may have a significant role in the colonization of laccase positive gut microflora.
RELATIONSHIPS BETWEEN MICROBIAL GENE CONTENT, PHYLOGENETIC DIVERGENCE AND ENVIRONMENTAL PREFERENCES
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Background

Environmental characteristics impose some particular constraints to the species living in any given habitat. In many instances, prokaryotic taxa show environmental preferences that shape the diversity found in diverse different environments. It is reasonable to expect that these habitat preferences have a reflection on the metabolism of the prokaryotic organisms. All these metabolic choices require particular sets of genes, and, therefore, gene content is determined to some extent by environmental preferences. But it is unclear to what extent is this relationship determinant, or how extensive is the metabolic rearrangement leading to different environmental adaptations.

On the other hand, it seems obvious that genomic content is determined highly by phylogenetic proximity. The extent to which each of the two processes modulates the genomes has not been quantified.

Objectives

We present a comprehensive analysis of the relationships between phylogeny, metabolism and environmental similarity in the prokaryotic world. We aim to quantify the influences of phylogenetic descent and environmental preferences in genomic content, to answer the question of how much it is shaped by each of the contributions.

Methods

We have calculated measures of phylogenetic, genomic and environmental distances between pairs of prokaryotic taxa using their 16S distances, common gene content and environmental preferences.

Conclusions

The results indicate a strong influence of the phylogeny in the genomic content, although the influence of the environment is notable for some particular metabolic
classes. This allowed to discover sets of genes that can act as environmental markers and predictors of the possible habitats for new species.
DISCOVERY OF PUMILARIN, A NOVEL HEAD-TO-TAIL CYCLIZED PEPTIDE, BY BAGEL3

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Background
In the post-genomic era the amount of (public) genomic data is ever increasing. Bacteriocin mining tools can be used to extract valuable information from this data regarding the presence of post-translationally modified peptides.

Objectives
As a proof of principle we mined all publicly available bacterial genomes using Bagel3. For simplicity we limited the results to head-to-tail cyclized peptides only. We found a large variety of these (predicted) circular peptides in the Bacillus genus. Based on the mining results we screened several Bacillus pumilus strains for the production of novel (head-to-tail) bacteriocins.

Methods
This yielded the production of a novel bacteriocin that we call pumilarin. The overall homology with As-48 is about 50%. We purified pumilarin using protocols that are based on AS-48 purification methods.

Conclusions
Next we analysed its structure using mass-spectrometry and found that its mass was in accordance with its circular structure. After fragmentation masses could be observed that could be explained only by N- to C-terminal linkage of the peptide. The activity spectrum of pumilarin was benchmarked against the prototype head-to-tail cyclized bacteriocin AS-48 and found to show some interesting differences in spectrum.
MATHEMATICAL MODELLING OF BACTERIAL CROSS-FEEDING: APPLICATION IN THE HUMAN INTESTINAL TRACT

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Background
In nature bacteria are rarely found in isolation, rather they reside in close association and are commonly found within biofilms. Bacteria that develop alternative feeding mechanisms, such as cross-feeding, have a much greater chance of survival in a competitive environment. In the human intestinal tract, the many types of cross-feeding include the use of oligosaccharides and/or monosaccharides resulting from extracellular degradation of polysaccharides by a competing bacterium, the use of such molecules along with metabolites produced by other bacteria and solely using metabolites produced closely associated bacteria.

Objectives
We aimed to develop models to describe the three aforementioned types of microbial cross-feeding and use those models to understand the cross-feeding phenomenon.

Methods
We developed a system of mechanistic differential equations to describe the different types of bacterial cross-feeding. These models were fit to multi-response data found in the literature using nonlinear regression. The multi-response data included bacterial growth (Bacteroides, Bifidobacterium, Eubacterium, Roseburia), substrate degradation (inulin, oligofructose, fructose, and metabolites), and metabolite concentrations (succinate, acetate, lactate and butyrate). To ensure that the estimated model parameters were independent of scale, we minimized the diagonal of the matrix ZTZ, where Z is the residual matrix formed from the data and model fit.

Conclusions
Our results suggest that microbial cross-feeding follows second-order chemical kinetics even when the ratios of two substrates are not one-to-one.
COMPARATIVE GENOMIC ANALYSIS REVEALS SUBTELOMERIC GENE DUPLICATIONS AND METABOLIC POTENTIAL OF THE KLUYVEROMYCES GENUS

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Background

The yeasts *Kluyveromyces lactis* and *Kluyveromyces marxianus* are the best-studied species of the *Kluyveromyces* genus. *K. lactis* has been adopted as a model for non-*Saccharomyces* yeasts whereas *K. marxianus* is more widely used for a range of different industrial applications because of relevant phenotypic traits. A *K. lactis* genome sequence has been available for some time but only recently genome sequences for a number of different *K. marxianus* strains have been generated.

Objectives

Our aim to understand the genetic basis of the metabolic and physiological differences between *Kluyveromyces* and *Saccharomyces* and within the *Kluyveromyces* genus.

Methods

We compared the proteomes of *K. marxianus*, *K. lactis* and *S. cerevisiae* by protein clustering using OrthoMCL. We found 4,119 common clusters (in all 3 yeasts) and 596 exclusively shared by the two *Kluyveromyces* species. More detailed bioinformatics as well as phenotypic assays were carried out to verify the findings.

Conclusions

Some key differences arise in sub-telomeric regions, where the same genes can be found duplicated at each end of the chromosome or at the ends of a separate chromosomes. For example, the genes responsible for lactose assimilation are duplicated in the subtelomeric regions of *K. marxianus* but not in *K. lactis*. Indeed, detailed comparison this duplicated region in different *K. marxianus* strains identifies variation that may account for some of the phenotypic differences in lactose assimilation between strains. Other differences that give rise to phenotypic variation
between species have also been verified and new metabolic pathways identified in *K. marxianus*. 
COMPARATIVE GENOMIC ANALYSIS OF NINE SPHINGOBIUM STRAINS: INSIGHTS INTO THEIR EVOLUTION AND HEXACHLOROCYCLOHEXANE (HCH) DEGRADATION PATHWAYS

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Background

Species belonging to the genus Sphingobium are well known for their ability to degrade hexachlorocyclohexane (HCH) isomers due to the presence of lin pathway genes. However, the establishment of the lin pathway in sphingomonads under HCH pressure is still unknown.

Objectives

Here, the nine Sphingobium spp. (strains LL03, DS20, IP26, HDIPO4, P25 and RL3, B90A, UT26S and Sphingobium sp. SYK6) were compared in order to reveal the mode of acquisition of lin genes.

Methods

Genomes of Sphingobium spp. were assembled using ABysS1.3.3 followed by their validation with bwa0.5.9. Phylogeny was constructed with 16S rRNA, single copy genes, ANI and tetranucleotide frequency. Further, draft genomes were annotated using Glimmer-3 at RAST4.0 server. Orthologs were identified using CD-HIT and all versus all BLAST. Functional annotation was performed using KAAS and gene families were predicted using Minpath. dN/dS and dS for each gene pair was calculated using Hyphy2.1.2.

Conclusions

The phylogenetic analysis using genomic data clustered efficient HCH degraders in a closely related group comprising of UT26S, B90A, HDIPO4 and IP26, where HDIPO4 and IP26 were classified as subspecies with ANI value >98%. Total gene content shared among all nine strains was ~10% which increased to ~25% in eight HCH-associated strains. Additionally, an inter-genus plasmid pool between genus Sphingobium and Sphingomonas was observed. Further, the differences in lin gene sequences, copy number and arrangement with respect to IS6100 revealed possible
evolutionary acquisition mechanisms for this pathway. This study also reflected that HDIPO4, IP26 and B90A are better suited organisms for HCH bioremediation.
DIVERSITY OF SURFACE-ASSOCIATED BACTERIA AND ANTIBACTERIAL ACTIVITY OF THE MARINE SPONGES COLLECTED FROM MARINE AREAS OF TURKEY

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Background
Increasing resistance problem of human pathogenic bacteria against commercial antibiotic derivatives has become important problem of all over the world.

Objectives
Anti-bacterial activity of the methanolic extracts of the marine sponges against pathogenic bacteria and diversities of sponge-associated and free-living bacteria were investigated.

Methods
Taxonomic designations of the sponges were carried out using histological sample preparation technique. The samples were extracted using cold methanol extraction technique. The agar disc diffusion method was used to determine the anti-bacterial activity of the sponge extracts. The pure bacterial strains isolated from the sponges were identified using the automated micro identification system VITEK2 Compact30.

Conclusions
The methanolic extracts of all tested sponges displayed positive antibacterial activity at varying levels against pathogenic bacteria. Antibacterial effectiveness rate was found higher in the samples which were collected from the Aegean Sea than the Sea of Marmara. The community profiles of the sponge associated bacteria were found different from free-living bacteria. Though similar bacteria species were found in all sponge samples, the predominant bacterial community profiles were different. Of the 325 unit bacterial strains, isolated from the Aegean Sea sponges, 86% belonged to phylum Proteobacteria. 82% of 102 bacterial strains, isolated from the sponges of the Sea of Marmara, were Proteobacteria. The study resulted in increasing data to understand antibacterial activity of the marine sponges regarding diversities of the sponge/sponge-associated bacteria according to their exposure local environmental conditions in the in the different marine areas.
EFFECT OF TEMPERATURE ON BACTERIOPHAGE-MEDIATED BACTERIAL LYSIS EFFICIENCY

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Background
Bacteriophages (phages) are viruses that specifically target bacteria. This property makes them useful agents for the control of pathogenic bacteria. Despite their great potential there is great uncertainty about the environmental parameters affecting phage-mediated lysis efficiency.

Objectives
In this study, temperature was investigated as a key environmental factor.

Methods
Three distinct phages specific for *E.coli* were isolated from a single brook water sample. Lysis efficiency was monitored using the double agar overlay method and optical absorbance measurements. Plaque sizes and numbers varied dramatically depending on whether phage-host mixtures were incubated at 20°C, 30°C or 37°C with bigger and more plaques being visible at lower temperatures. Interestingly plaque formation was invisible at the optimum temperature of the host (37°C). Plaque numbers and sizes were found to additionally depend on the temperature history of the bacterial host prior to mixing with the phage. This applied both to the growth temperature of the *E. coli* (with higher plaque numbers and sizes when cells were grown at lower temperature) and short term temperature exposure of fully grown cells. Exposure of *E. coli* to sublethal heat stress resulted in a phage resistant phenotype.

Conclusions

These outcomes suggest that many phage-host interactions might be highly temperature-sensitive and phages seem to prefer host cells that were subjected to temperature conditions similar to the ones in the environment where the phages were isolated from. Phages-host interactions should be well characterized for their temperature preference and probably other environmental parameters prior to biocontrol and biosanitization applications to obtain optimal performance.
CHANGES IN MEMBRANE SUBPROTEOME IN VIBRIO HARVEYI DURING THE ADOPTION OF VIABLE BUT NONCULTURABLE STATE BY EFFECT OF STARVATION AND VISIBLE LIGHT

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Background

*Vibrio harveyi* is a pathogen bacterium for a wide range of marine animals that can be isolated as a planktonic form. The behaviour of *Vibrio* spp. in natural environments is conditioned, among others, by temperature, light, salinity or nutrient scarcity.

Objectives

The aim of this study was to evaluate the physiological and proteomic changes that take place during *V. harveyi* survival in seawater under exposure to visible light.

Methods

For this purpose, populations of *V. harveyi* were maintained in sterile natural seawater at 20°C and under illumination for at least 21 days. Along experiments, total, active and culturable populations were enumerated and the outer membrane subproteome was analyzed.

Conclusions

Experimental conditions had a negative effect upon culturability of *V. harveyi*, inducing its entry into the viable but nonculturable (VBN C) state after 6-7 days of exposure, and provoked changes in the membrane subproteome composition.

Along the period of study, some outer membrane proteins remained unchanged (e.g. OmpW), while a set of proteins increased (i.e. lipoproteins and agglutination proteins) or became detectable under tested conditions (in this last group the transport-related proteins are especially relevant). So, all these proteins could be likely essential for sustaining the *V. harveyi* viability under stress conditions. In contrast, several proteins and especially those related to chemotaxis processes decreased or were undetected along survival.

Therefore, in response to starvation and exposition to visible light *V. harveyi* experiments an adaptation process which includes the entry into the VBNC state and the membrane subproteome reorganization.
BIOTECHNOLOGICAL POTENTIAL OF BACTERIA ISOLATED FROM MARINE SPONGES

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Background

Marine sponge contains a great number of microorganisms present in their tissues. These ones perform symbiotic associations with their hosts, and are capable of producing secondary metabolites with various biotechnological properties.

Objectives

Aiming to know more about microbial diversity present in aquatic environments and get bacterial species with potential application in biotechnology

Methods

three species of marine sponges (Hymeniacidon heliophila, Amphimedon viridis and Aplysina fulva) were collected in two beaches of Rio de Janeiro, Brazil; 38 bacterial isolates from these sponges were tested for the production of amylase, cellulase and caseinase using standard methodology.

Conclusions

Of these, 19 showed enzymatic activity, 5 for amylase, 4 to cellulase and 16 for production of caseinase. Of all isolates tested only one colony was able to degrade the three substrates. The tests of biosurfactants’ production, showed that 17 colonies were able to emulsify diesel. Our results show that the species of marine sponges studied are a great substrate for prospecting microorganisms with potential biotechnological applications.
TOXICITY OF TERRESTRIAL CYANOBACTERIAL STRAINS ORIGINATED FROM FORESTS OF THE SERBIAN MOUNTAINS

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Background
As one of the oldest and highly diverse group of photosynthetic prokaryotic microorganisms, cyanobacteria are known as potent producers of various bioactive compounds with toxic effects.

Objectives
Toxicity of five forest terrestrial, filamentous cyanobacterial strains belonging to Oscillatoria and Nostoc genera was investigated.

Methods
The toxicity of intracellular extracts of cyanobacterial strains was tested in Artemia salina bioassay. The LC50 values were recorded in all tested strains after 24 h (Nostoc genera) and 48 h (Oscillatoria genera). In the case of three Oscillatoria strains, a dose-dependent response in mortality that increased over time was observed after 24 h (1% - 8%) and 48 h (72% - 98%) of exposure. However, for the intracellular extracts of two strains, Nostoc T7 and Nostoc M2, toxicity was higher after exposure time of 24 h (70% and 97%) and 48 h (93% and 98%). According to the LC50 values, cyanobacterial strain Nostoc M2 showed to be the most toxic strain with LC50 value 0.59 mg/ml, followed by strain Nostoc T7 (LC50=1.25 mg/ml). Concerning Oscillatoria genera, the recorded LC50 values were 3.49 mg/ml (Oscillatoria M1), 4.55 mg/ml (Oscillatoria M2) and 5.62 mg/ml (Oscillatoria T18).

Conclusions
The results presented in this paper indicate that toxin production is strain-specific property, whereby intracellular content of three strains belonging to Oscillatoria genera was found to be less toxic to A. salina compared to the strains belonging to Nostoc genera. An additional objective which should be done is analysis and identification of toxic compounds present in the extracts responsible for observed toxicity.
Studies on production and purification of bioflocculant by a bacterium species isolated from Adeti stream in Ilesa, Osun state
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Background

Water clarification methods like flocculation, coagulation and sedimentation are often inappropriate because of the high cost and scarcity of chemical coagulants. This study was designed to provide a safe, cheaper and biodegradable flocculating agent.

Objectives

The objectives include isolation of bacterial isolate with flocculating properties from wastewaters and soil samples; selection of a bacterium with the highest bioflocculant-producing potential; determination of the time course of bioflocculant production; and purification of the bioflocculant.

Methods

Wastewater and soil samples were collected from Ilesa and plated on yeast extract, peptone and glucose (YPG) agar. Plates were incubated at 25°C for 24 h. Colonies with mucoid appearance were selected. Pure isolates were stored on sterile YPG agar slants. Bioflocculant production was carried out by growing the organism in YPG medium at 25°C and centrifuged at 5000 rpm for 30 min. Flocculating activity was measured by using a suspension of kaolin clay. Time-course of bioflocculant production was determined by monitoring pH, cell growth and flocculating activity. Bioflocculant was purified by precipitation and gel-filtration chromatography.

Conclusions

The highest bioflocculant-producing isolate was identified as Pseudomonas extremoaustriis ABL19. Maximum bioflocculant production was observed at the 66th h. The optimal bioflocculant activity of 94.4% was achieved with glucose and peptone in the medium. The molecular weight was estimated to be 60.8kDa. The bioflocculant was effective at 80°C and alkaline pH.
CHARACTERIZATION OF BIOACTIVE COMPOUNDS PRODUCED BY SOIL STREPTOMYCES SPP.
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Background
In general special properties of different geographical area could be considered as best sources of bioactive compounds, therefore investigation on these fields might be increased treatment of patients and reduce the rate of occurrence of antibiotic-resistant bacteria.

Objectives
Major purpose of this study was isolation of antimicrobial metabolite producing Streptomyces spp. from soil samples and identification the structure of their antimicrobial metabolites.

Methods
To perform the present study, 70 strains of Streptomyces spp. were isolated from soil samples and evaluated for producing of their antimicrobial products against some pathogenic microorganisms. Then, the antimicrobial producing strains were identified using phenotypic and genotypic methods. The best growth phase of production was determined and finally structures of antimicrobial compounds were assessed by 13C NMR, 1H NMR and FT-IR methods. Regarding to determination of possible structures, all data obtained from 13C NMR, 1H NMR and FT-IR were analysis by NCBI PubChem Structure Search program.

Conclusions
Out of 70 strains of indigenous Streptomyces spp., three strains had potent activity for production of antimicrobial metabolites. The isolates recognized as Streptomyces griseus, Streptomyces phaeochromogenes and Streptomyces olivoviridis. Metabolites
showed antimicrobial activity against *Escherichia coli*, *Salmonella typhi*, *klebsiella pneumonia*, *shigella dysenteriae*, *Bacillus cereus* and *Staphylococcus aureus*. The best growth phase for antimicrobial metabolite production was stationary phase, therefore based on these data all the metabolites recognized as secondary metabolites. Regarding to determination of possible structures, all data obtained from 13C NMR, 1H NMR and FT-IR were analysis by NCBI PubChem Structure Search program and resulted suggestion three possible structures of bioactive compounds.
ISOLATION AND CONSERVATION OF FLUORESCENT PSEUDOMONADS STRAINS FROM RHIZOSPHERE OF WHEAT

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Background

Plant Growth Promoting Rhizobacteria (PGPR) meets those of inhomogeneous rhizobacteria groups which are patented to increase and promote plant growth. Pseudomonas bacteria is one of the most important PGPR bacteria which can increase yield, growth and health of the plant through direct and indirect mechanisms.

Objectives

In the present study, fluorescent pseudomonads isolated from Savadkooh’s soils have been investigated with respect to their biochemical tests and physiological characteristics.

Methods

In the first experiment, 30 soil samples were collected from the rooting zone of wheat at farmer’s field in Savadkooh, Mazandaran province, Iran. The soil samples were serially diluted up to 6–7, plated on King’s B (KB) agar medium and incubated at 28±2°C for 48 h. Distinct colonies showing fluorescence under UV light were picked and streaked on KB agar medium to check the purity. In the second experiment, for the identification of fluorescent pseudomonads, certain biochemical tests and physiological characteristics were conducted according to Bergey’s Manual for determinative bacteriology.

Conclusions

The results showed that 17 pseudomonas strains were detected based on yellow, green and blue pigments by viewing under UV light. Population of bacteria were $2.71 \times 10^5$ – $6.43 \times 10^7$ cell per gram of rhizosphere soil. Based on results of biochemical tests and physiological characteristics we were detected three species, $P. putida > P.$
fluorescens >> P. aeruginosa. Dominate species of fluorescent pseudomonads in rhizosphere of wheat was putida.
SEASONAL DYNAMICS OF SOIL MICROBIAL COMMUNITIES UNDER DOMINANT UNDERSTORY VEGETATION IN SPRUCE SWAMP FOREST

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Background

Spruce swamp forests (SSF) are exceptional ecosystems which are considered to be important sources of greenhouse gases (GHG). Understorey vegetation dominates Eriophorum vaginatum and Vaccinium myrtillus which create patchy environment in SSF may mitigate GHG emissions. Presence of Eriophorum or Vaccinium can affect GHG emissions through the quality and quantity of its litter and root exudates. Their input directly affects microbial community and its decomposition capabilities.

Objectives

Our two main objectives were (i) to describe the seasonal dynamics of bacterial and archaeal communities in spruce swamp forests under the two dominant understorey vegetations (Eriophorum and Vaccinium) and (ii) to evaluate the effect of dominant vegetation on activity of soil enzymes.

Methods

To characterize microbial community we sequenced variable region V4 of 16SrDNA using Illumina MiSeq platform. OTU-picking and taxonomic assignment was performed using the QIMIE 1.8.0 bioinformatics pipeline. Hydrolytic enzymes were analyzed by fluorometric method using 4-methylumbelifferone (MUB) labeled substrates.

Conclusions

Methanogenes (Methanomicrobia, Methanobacteria) dominated archaeal community followed by Thermoplasmata, MBGA and MCG (Fig. 1). Relative abundance of methanogens was significantly higher in Spagnum and Eriophorum than in Vaccinium sites. Eriophorum and Vaccinium sites were enriched by Proteobacteria in comparison to Shagnum sites (Fig. 2). On the other hand Sphagnum control site had higher relative amount of anaerobic bacteria (Firmicutes-Clostridia and Chloroflexi). Hydrolytic enzymes were dominated by phosphatase activity showing probably P
limitation which was highest on Sphagnum control site.

Community composition of Archaea

Fig. 1. Seasonal dynamics of archaeal community
Fig. 2. Seasonal dynamics of bacterial community
ANAEROBIC OXIDATION OF METHANE IN SURFACE SEDIMENTS OF MARINE LAKE GREVELINGEN, NORTH SEA

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Background

Sulphate reducing anaerobic oxidation of methane (AOM) is a well-established phenomenon occurring in deep marine environments (~ below 100 m). However, relatively shallow anaerobic sediments containing methane and sulphate may harbour anaerobic methanotrophic microorganisms (ANME). Marine Lake Grevelingen is a former estuary with a water depth of ~ 40 m and salinity of 31 g/kg, which is separated from the North Sea by a dam. High rates of degradation and deposition of organic matter have resulted in methane rich anoxic sediments which combined with sulphate from sea water renders the site a potential niche for AOM.

Objectives

The main aim of this research was to explore whether AOM occurs in the surface sediment of Marine Lake Grevelingen.

Methods

Pore water chemical analysis, serum bottle incubation for activity tests, and microbial analysis by fluorescence in situ hybridisation (FISH) were conducted.

Conclusions

AOM was evident at depths of 5 to 15 cm in the sediment with a steep decline in CH₄ concentrations from ~5 mM at 20 cm depth to negligible at the sediment surface; concurrently, sulphide concentrations increased to 5 mM. In vitro incubations with CH₄ and SO₄²⁻ showed sulphide production coupled to the consumption of sulphate at approximately equimolar ratios at 150 days. In contrast, sulphate and sulphide concentrations remained constant in biotic and abiotic control incubations. ANME and
archaeal cells were observed by FISH, which could be linked to AOM activity. Using three independent approaches, this study, for the first time provides evidence for the occurrence of AOM in sediments of Marine Lake Grevelingen.
EXOPOLYSACCHARIDE PRODUCTION BY RHIZOBIUM ETLI STRAIN ZD13, A HEAVY METAL-RESISTANT SOIL ISOLATE

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Background

Exopolysaccharide (EPS) production functions as a protective strategy which favors the survival of microbes under unfavorable environmental conditions, e.g., heavy metal stress. Rhizobia are Gram-negative, rod-shaped bacteria known for EPS production. Some EPS, due to the presence of negatively charged functional groups, may act as heavy metal-binding agents.

Objectives

Studies on the EPS produced by Rhizobium etli strain ZD13 (RheZD13), an isolate from a postindustrial soil of the Upper Silesia (southern Poland).

Methods

RheZD13 was grown in 4 liters of a culture medium under aerobic conditions (120 h, 30°C, 120 rpm). Bacteria were pelleted by centrifugation (60 min, 4°C, 11806 x g) and the supernatant was sterile-filtered (Ø 0.22 µm) and treated with cold (-20°C) ethanol to precipitate the crude EPS. The latter was dialyzed and analyzed by GC and GC-MS.

Conclusions

4.627 g of RheZD13 EPS were obtained. The EPS contained glucose and galactose (in a molar ratio 3:1) and pyruvic acid that was 4,6-linked to galactose. The acidic nature and the resulting multiplicity of potential binding sites for positively charged particles suggested that RheZD13 EPS may act as a biosorbent for heavy metals.
Water Treatment Plants (WTP) generate waste as residual decanters sludge due to the production of potable water. The objective of this study was to characterize microbiologically the sludge from three samples from the WTP of Londrina, Brazil, through microbiological indicators – as Enterococcus spp, sulphite-reducing Clostridium and pathogen Salmonella spp. The 'multiple tubes' technique was used to isolate bacteria. For physical-chemical analysis parameters it was used Biochemical Oxygen Demand (BOD), Chemical Oxygen Demand (COD), turbidity, pH and apparent color (according to APHA, AWWA and WEF-2005). The results showed that in sample 1, the most probable number (MPN) of microorganism per 100mL was 2,1x10^4 Enterococcus and 1,1x10^5 Clostridium; the sample 2 presented absence of Enterococcus and >2,4x10^5 Clostridium; and the sample 3, 1,0x10^5 Enterococcus and 2,4x10^5 Clostridium. Salmonella spp. was not detected in any sample analyzed. For sample 1, the BOD was 113,9 mg of oxygen/L, the COD of 6184,09 mg of oxygen/L, with turbidity of 24000 NTU, the pH 6,0 and apparent color of 100000 uH. For the sample 2, the BOD was 4334,59 mg of oxygen/L, the COD was not determined, with turbidity of 16800 NTU, the pH 6,38 and apparent color of 6000 uH. In sample 3, the BOD was 73,3 mg of oxygen/L, the COD was 2172,20 mg of oxygen/L, with turbidity of 4370 NTU, the pH 6,90 and apparent color of 16700 uH. The microbiological contamination was relevant, indicating high destructive potential for the environmental and healthy human population.
MICROBIOLOGICAL AND PHYSICAL-CHEMICAL CHARACTERIZATION OF FLUSHING WATER FILTER FROM WATER TREATMENT PLANTS IN BRAZIL

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Water treatment plants (WTP) yields waste as decanters sludge and flushing water filter (FWF). The objective of this study was characterizing FWF for presence of Enterococcus spp., Clostridium sulphite-reducing and Salmonella spp.. For physical-chemical parameters Biochemical Oxygen Demand (BOD), Chemical Oxygen Demand (COD), turbidity, pH, apparent color, AWWA and WEF 2005 were used. The results showed in WTP–A (collection one), most probable number (MPN) of microorganism per 100mL was 1,2x10² Ente-rococcus and >2,4x10³ Clostridium; Enterococcus and 1,1x10⁴ Clostridium. BOD was 4,8 oxygen/L, COD 218,30mg oxygen/L, turbidity 754 NTU, pH 6,07 and apparent color 300 uH; In WTP-B, for collection 1, absence Enterococcus, 7,0 x10¹ Clostridium; BOD was 3,2 mg oxygen/L, COD 294,5 mg oxygen/L, turbidity 3990 NTU, pH 6,07 and apparent color 3000 uH. For collection 2, absence Enterococcus and 4,3x10² Clostridium; BOD 2,43 mg oxygen/L, COD 1,3 mg oxygen/L, turbidity 881 NTU, pH 6,57 and apparent color 5500 uH; In WTP-B, for collection 2, absence Enterococcus and Clostridium; BDO 5,1 mg oxygen/L, COD 106,8 mg oxygen/L, turbidity 889 NTU, pH 6,78 and apparent color 4500 uH. For collection 3, 3,9x10²; BOD 11 mg oxygen/L, COD 628,83 mg oxygen/L, turbidity 1600 NTU, pH 6,98 and apparent color 54 uH; 4,0x10¹ Enterococcus and 4,0x10¹ Clostridium; BDO 3,7 mg oxygen/L, COD 170,9 mg oxygen/L, turbidity 190 NTU, pH 7,11 and apparent color 1100 uH. Salmonella spp. was not isolated. The microbiological contamination was relevant and high levels of physical-chemical parameters were detected in the flushing water filter, indicating destructive potential for the environment and population.
ENTERIC VIRUSES INACTIVATION BY GAMMA IRRADIATION

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Background
Enteric viruses, like norovirus and adenovirus are frequently detected in environmental systems and have been implicated in many outbreaks. Normally, the viruses exist in the environment and acquire certain capability to endure conventional treatment processes. Alternative disinfection methods must be developed.

Objectives
The main goal of this study was to investigate the inactivation of murine norovirus type 1 (MNV-1), as a Norovirus surrogate, and human adenovirus type 5 (AdV-5) by gamma irradiation.

Methods
Six different substrates: Phosphate buffer saline (PBS), demineralized water, tap water, fetal bovine serum (FBS) and aqueous solutions of 10% and 50% FBS, were inoculated with MNV-1 and AdV-5 and irradiated in a Co-60 source at several doses (1 up to 10 kGy). The inactivation of viral particles was tested by plaque assay using Raw 264.7 and A549 cells. The $D_{10}$ values were estimated for each virus and substrate.

Conclusions
A reduction on MNV and AdV titers of $4 \log_{10}$ PFU/ml was achieved after irradiation at 3 kGy on PBS and water suspensions. However, MNV and AdV were approximately 3 times more resistant to gamma irradiation when irradiated in FBS, and at a dose of 10 kGy it was detected the presence of infective viral particles. The observed $D_{10}$ values ranged between 0.76 kGy (water) and 3.22 kGy (10% FBS) for MNV; and between 0.87 kGy (PBS) and 2.94 kGy (FBS) for AdV. These results show that inactivation of MNV-1 and AdV-5 by gamma radiation strongly depends on the substrate where the viruses are inoculated.
RECOVERY OF AN ACETATE-CONSUMING SULFATE REDUCING CONSORTIUM AT ACIDIC CONDITIONS

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Background
Sulfidogenic reactors have been used to eliminate dissolved metals from industrial wastewaters, due to their possibility to obtain metal sulfides. Sulfate reducing bacteria (SRB) are responsible for the production of sulfide which reacts with dissolved metals, forming precipitates (metal sulfides). However, the major drawback of sulfidogenic reactors is that acetate cannot be degraded completely, reducing the reactor’s efficiency, another problem is that metal containing wastewaters are usually acidic (pH 2-4).

Objectives
The objective of this work was to obtain a consortium from a natural acidic environment (an abandoned sulfur mine) able to work at low pH (<4) and produce sulfide from the complete oxidation of the substrate.

Methods
The enrichments were performed from sediments cultured in anaerobic bottles (120 mL) and fed with acetate, glycerol, or lactate, and sulfate. The pH was initially 3 or 4, and we evaluated their capability to completely consume the substrate and produce sulfide. When the enrichment consumed the substrate and produced sulfide, we took 10 or 20 % (v/v) as an inoculum to develop a new enrichment of sulfate-reducing consortium.

Conclusions
Supernatant (only the liquid part) used as inoculum (10-20%) was not favorable for the growth of SRB. The slurry (combination of sediment and liquid) used as inoculum (20%) promoted the sulfate-reducing activity at acidic pH. Acetate, used as the sole electron donor, was not suitable for the SRB consortium. Complex electron donor, such as glycerol or lactate, was more suitable to obtain a sulfate-reducing consortium at low pH able to consume acetate and produce sulfide.
BACTERIAL DIVERSITY AND DYNAMICS FROM START-UP TO STEADY CONDITIONS IN A FULL-SCALE MUNICIPAL SOLID WASTE (MSW) ANAEROBIC REACTOR BY 454 PYROSEQUENCING TECHNOLOGY.

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Background

Anaerobic digestion is an environmentally acceptable means of reducing and stabilizing organic fraction from MSW. Functioning and stability of an anaerobic reactor is directly related to the microbial community within it.

Objectives

The bacterial diversity of a full scale anaerobic reactor treating MSW from Madrid (Spain) was investigated using high-throughput 454 pyrosequencing technology.

Methods

The digested MSW samples were taken in two moments of the treatment: start-up phase and steady-state conditions. The bacterial 16S rRNA genes were amplified and sequenced using 454 pyrosequencing FLX machine.

Conclusions

118,392 bacterial 16S rDNA reads were yielded and after trimmed 35,981 (start-up phase) and 18,290 reads (steady conditions) were clustered into OTUs. Fifteen phyla were identified: Bacteroidetes, Firmicutes, Actinobacteria, Proteobacteria, Chloroflexi, SR1, Synergistetes, Planctomycetes, Tenericutes, Deinococcus-Thermus, Acidobacteria, Chlorobi, Cyanobacteria and Gemmatimonadetes. Shifts in the bacterial communities were observed and proteolytic bacteria dominated during all the process, being the major diversity identified in the steady-state condition of the reactor. Fermentative bacteria belonged to Bacteroidetes and Firmicutes phyla predominated, and a great number of reads were identified to genus level. Proteiniphylum, Gallicola and Fastidiosipila genera carried out most biodegradation processes during the MSW treatment. A great number of genera are involved in the
sugars and carbohydrates metabolisms, although its coverage is low. The high concentration of ammonium appears to be the reason for the prevalence of the proteolytic bacteria. The use of the next generation sequencer technology revealed a great diversity of rare organisms and increased our knowledge about the dynamics of bacterial communities in anaerobic MSW reactors.
BACTERIAL DIVERSITY IN RHIZOSPHERE AND SOIL ON MINING WASTE IN THE ARID ENVIRONMENT OF THE SAHARA

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Background

Relatively few studies have been devoted to the microbial diversity of the Sahara. In this study, we have investigated an arid area located in south-west Algeria. In this area, spoil heaps of coal mine waste are colonized by a Saharo-Mediterranean vegetation dominated by Caroxylon sp.

Objectives

As part of an on-going study aiming to understand the role of micro-organisms in the adaptation of this plant to the site, we have studied its associated microflora.

Methods

The bacterial community was investigated by fingerprinting (RISA). The diversity was much higher in the Caroxylon rhizosphere than the surrounding soil, both on the waste heaps and in the desert soil. Libraries of 16S rRNA gene were obtained from rhizospheric and non rhizospheric soil–extracted DNA. Sequences from Firmicutes and Actinobacteria were abundant in all libraries. As expected from the fingerprints, the diversity was higher in the rhizosphere libraries, with 1.6 more phyla detected than in non rhizospheric soil.

Conclusions

Depending on the library, 25 to 65 % of the sequences had percentages of identity with known species below the 97% threshold. Potential new species are located in very diverse phyla. The most distant ones (less than 90% identity) belong to Acidobacteria, Chloroflexi, Deltaproteobacteria, Cyanobacteria and Nitrospirae. The results show that we have only begun to explore the diversity of soil bacteria in the arid environment of the Sahara.
POTENTIAL OF CADMIUM RESISTANT AND BIPHENYL UTILIZING MARINE BACTERIA IN BIOREMEDIATION

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Background
Utilization of aromatic compounds by marine microbial communities in nutrition and pollution stress has evolved several catabolic pathways for bioremediation. Of which, biphenyl and cadmium are highly toxic compounds having harmful effects on flora, fauna, and mankind.

Objectives
The main objective of this study was to find out the catabolic potential and pathway of marine bacteria in aerobic utilisation of biphenyl as sole carbon and energy source under toxic metal (cadmium) stress.

Methods
Two bacterial strains *Pseudomonas aeruginosa* (JP-11) and *Achromobacter xylosoxidans* (JP-22) were isolated from the coastal regions of Odisha, India, which could resist up to 1000 ppm and 630 ppm of cadmium respectively. They utilized 97.88 % and 89.6 % of biphenyl respectively. Combination of chromatographic and spectrophotometric techniques identified 2-Hydroxy-6-oxo-6-phenylhexa-2, 4-dienoate and benzopropanol as the intermediate product of biphenyl metabolism in JP-11 and JP-22 respectively. Biosurfactant production was observed which increased the bioavailability of this hydrophobic compound into the cells resulting in the effective removal from the environment. Of both the isolates, the partial structure of the biosurfactant produced by JP-11 was similar to that of rhamnolipid which showed an upregulation of rhamnolipid synthesis gene (*rhlAB*) with increased biphenyl concentration (upto 200 ppm).

Conclusions
Thus, these bacterial isolates can be efficiently utilized for bioremediation of the contaminated sites with toxic metal and organic pollutants.
DIVERSITY OF OIL DEGRADING BACTERIA FROM GEOGRAPHICALLY DISTANT REGIONS

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Background
An investigation of oil degrading bacteria is significant for practical and fundamental reasons. The diversity of metabolic systems allows these microorganisms to utilize a wide spectrum of substrates and produce different biological active substances.

Objectives
The main aim of this research was to characterize oil degrading bacterial strains isolated from different geographical regions (Belarus, Libya, Iraq, and Antarctic).

Methods
Physiological, biochemical and molecular genetic methods were used.

Conclusions
Taxonomy of 22 oil degrading bacterial strains was established. They were determined as *Rhodococcus pyridinivorans*, *R. opacus*, *R. erythropolis*, *Bacillus licheniformis*, *B. beijingensis*, *B. flexus*, *Arthrobacter* sp., *Micrococcus* sp., *Enterobacter* sp., *Planococcus maitriensis*, *Acinetobacter radioresistens*, *Pseudomonas* sp., *P. stutzeri*, *Dietzia* sp., *Deinococcus* sp.

*R. pyridinivorans* 5Ap was shown to have the widest spectrum of utilizing substrates (it utilized 19 investigated organic substrates). Different strains were able to grow under extreme conditions. For example, *B. licheniformis* FD9 grew at 54 ºC; *P. stutzeri*, *R. erythropolis*, *Arthrobacter* sp. - at 4 ºC; *Enterobacter* sp. FD1 - pH=4, *B. beijingensis* FD4 - pH=11, *P. maitriensis* FD3 – with sodium chloride concentration up to 7 %; *A. radioresistens* and *Deinococcus* sp. were UV-resistant.

It was shown the presence of naphthalene catabolic plasmids in the strains of *R. pyridinivorans* and *R. opacus*.

The ability of oil degrading bacteria to produce the surfactants was shown. The most effective production of surfactants was determined for *R. erythropolis* A29-k1 when it was cultivated on mineral medium with hexadecane. Lichenisyn biosynthesis genes were found in *B. licheniformis* FD9.
MICROBIOLOGICAL INVESTIGATION OF MUD VOLCANOES FROM EASTERN ROMANIA FLUIDS

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Background

The largest areas of terrestrial mud volcanoes in Europe are located in the Carpathian Mountains in eastern Romania. Till now the few microbiological investigations were focused on the study of Archean communities implicated in methane- and hydrocarbon cycling.

Objectives

The purpose of this study was to isolate, identify and characterize the pathogenic and resistance features of bacterial strains isolated from the bubbling fluids expelled from the mud volcanoes from Buzau in order to establish the contribution of these protected environments to the natural reservoirs of resistance and pathogenicity.

Methods

All water samples (500 ml) were collected according to current ISO guidelines using sterile glass bottles, at a depth of approximately 30 cm and stored in cold bags at 4°C until analysis. The samples were diluted and analysed through the standardized membrane filtration method. The identification of aerobic strains obtained in pure culture was based on Gram staining, oxidase reaction and API galleries. Simultaneously, the samples were processed for DNA extraction and molecular assays based on ribosomal DNA amplifications with general bacterial 16S rRNA primers. The isolated strains have been investigated for antibiotic susceptibility and for the production of cell-associated (adherence to inert and cellular substratum, biofilm development) and soluble, enzymatic (hemolysins, lecithinase, lipase, caseinase, gelatinase, amylase, esculin hydrolysis, DN-ase) virulence factors, using phenotypic (disk diffusion method, selective media for enzymatic factors production) and PCR-based methods.
Conclusions

The virulence and resistance profiles of cultivable strains were very poor, suggesting the absence of any selective pressure agents in the investigated areas.
IDENTIFICATION AND CHARACTERIZATION OF GENES INVOLVED IN SIDEROPHORE PRODUCTION OF PANTOEA ANANATIS, THE CAUSATIVE BACTERIUM OF RICE SHEATH ROT

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Background

Pantoea ananatis is known to cause disease in maize, eucalyptus, onions, and rice. This bacterium is also known as an opportunistic human pathogen. P. ananatis PA13 is a rice grain and sheath rot bacterium, which produces an antibacterial compound.

Objectives

In the present study, we have tried to identify and characterize genes involved in antibacterial activity in P. ananatis PA13.

Methods

We performed a large-scale, mariner transposon-based genetic screening and gene replacement strategy based on homologous recombination. Antibacterial activity was monitored against P. ananatis HY02, onion center rot bacterium, and Yersinia enterocolitica.

Conclusions

A number of proteins that involved in transcription, catalytic enzymes, siderophore transporting systems, and TonB-dependent transporter system were found to affect the antibacterial activity of strain PA13. These data indicate the iron uptake is a major factor of antibacterial activity in P. ananatis PA13. We focused then siderophore biosynthetic genes. P. ananatis has genes homologous to siderophore biosynthesis, aerobactin (iucABCD), alcaligin (alcA) and rhizobactin (rhbF). Single mutant of ΔiucABC exhibited slight reduction of antibacterial activity, whereas ΔalcA and ΔrhbF showed normal antibacterial activity. This result led to generation of double and triple mutants of the genes. Mutants ΔiucABCΔrhbF and ΔiucABCΔalcA exhibited strong reduction in antibacterial activity, whereas mutant ΔalcAΔrhbF showed normal antibacterial activity. These data indicate that siderophores play an important role in antibacterial activity of P. ananatis PA13. This work will give novel insights into iron
uptake regulation and contribute to a better understanding of how these genes function to regulate iron uptakes.
HYDROGEN PRODUCTION AND VFA ACCUMULATION IN A CONTINUOUS STIRRED TANK REACTOR USING MOLASSES WASTEWATER AS FEEDSTOCK

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Background
Molasses wastewater as a sugar-rich waste is a liquid by-product generated in large amount from food industries. Molasses wastewater is one of promising feedstock for bioenergy production because it has high strength of organics and relatively high bioavailability.

Objectives
For bioenergy production and wastewater treatment simultaneously, two-stage fermentation processes employing hydrogenesis and methanogenesis have been developed. Hydrogenesis has advantage of volatile fatty acids (VFAs) as main byproducts as well as hydrogen, and VFA production is necessary to stabilize the waste stream in the secondary stage for methane production. Hydrogen production and VFA production was characterized in a continuous stirred tank reactor (CSTR) fed molasses wastewater as feedstock.

Methods
The CSTR reactor achieved a stable performance at an organic loading rate from 20.3 to 36.1 g-COD·L⁻¹·d⁻¹. Average gas production rate was 1.2 L/d with a hydrogen concentration of 33%. The maximum hydrogen production rate was 1.02 L·H₂·L⁻¹·d⁻¹ at 31.0 g-COD·L⁻¹·d⁻¹. The abundant VFAs were butyric acid (50%) and acetic acid (38%). Total VFA production kept at around 7,135 mg/L level during operation period. Active microbial community was analyzed using rRNA-based massively parallel sequencing technique. Clostridiales, Lactobacillus and Clostridium were relatively abundant in the CSTR reactor. It has been reported that appearance of Clostridium species was in accordance with increase of butyrate and acetate concentration. Lactobacillus species has been known to be involved in anaerobic lactate degradation to acetate.

Conclusions
VFAs produced in the CSTR reactor can be utilized as substrates for methane production in downstream processes.
Background
Biohydrogen and biomethane can be produced in Upflow Anaerobic Sludge Blanket (UASB) reactors treating high strength wastewaters such as molasses wastewater and distillery wastewater.

Objectives
Even though many researches have been reported on the performance of UASB reactors and their microbial properties, there is little information about the effect of starvation on the UASB performance. In this study, the effect of starvation on methane production was characterized in an UASB reactor.

Methods
Anaerobic digested sludge was used as an inoculum source, and the effluent (10,000 mg-COD/L) from an hydrogen-producing UASB reactor treating molasses wastewater was used as influent wastewater for the methane-producing UASB reactor. The UASB reactor was operated for 18 days at hydraulic retention time of 3 d. After confirming the UASB performance reached at steady-state, the UASB was maintained at starvation condition for 14 d or 30 d. After starvation period, the UASB was re-operated by feeding the influent without re-inoculation, and COD removal and methane production were compared with their performance at steady-state (COD removal and methane production rate were 98% and 0.91 L-CH_{4}L^{-1}d^{-1}, respectively).

Conclusions
After 3-days of 14 d-starvation, COD removal and methane production rate were 87% and 1.51 L-CH_{4}L^{-1}d^{-1}, respectively. The UASB performance improved speedy, and the UASB performance could be recovered by 15-days operation after restarting. However just after 30 d of starvation, no COD removal and methane product was observed, and over 27-days operation was necessary to recovery the UASB performance. These findings can be used to plan the operation strategy of methane-producing UASB reactors.
APPLICATION OF BIOLOG PHENOTYPING MICROARRAYS FOR ASSESSING THE FUNCTIONAL DIVERSITY OF ENVIRONMENTAL STRAINS OF BACILLUS

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Background
The ability to link genotypes to corresponding phenotypes is of interest in biotechnological manipulation of metabolic pathways. In order to assess a rapid functional and phenotypic profiling under different metabolic conditions, a high-throughput approach has been developed by BIOLOG Inc. (Hayward, CA): the Phenotype Microarray (PM) technology.

Objectives
The objective of the study was to determine phenotypic changes in environmental strains of Bacillus by Phenotyping MicroArrays.

Methods
The four Bacillus strains were tested on PM 96-well plates (PM01-02 carbon sources, PM03 nitrogen sources, PM04 phosphorus and sulfur sources, PM09 with osmolytes, PM10 with another pH, PM11-13 with various antibiotics). Bacillus subtilis ATCC as reference strain was used to determine the differences between strains. Data were analyzed with Omnilog-PM software. The software suite called 'DuctApe' which analyzes and links together both the genomic and the phenomic data and suggests genetic explanations of metabolic phenotypes was also used.

Conclusions
The environmental isolates as compared to the reference strain were found to be the most metabolically versatile strains, either by looking at the number of 'more active' metabolic features or by the proportion of active compounds for each category. The strains showed higher proportion of active compounds for carbon and nitrogen sources and in the resistance to chemical agents. The DuctApe software is good tool for the visualization of PM data, the exploration of both the genomic information and the phenotypic expression of Bacillus spp. tested. Phenotypic profiling is an essential step for understanding genotype differences for environmental microorganisms of interest in contaminant remediation, biofuels production, and climate change.
THE STRUCTURAL AND FUNCTIONAL BIODIVERSITY OF MICROORGANISMS IN SOIL UNDER EX SITU AIDED PHYTOSTABILIZATION

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Background
The high toxicity of heavy metals causes the need to remove them from the contaminated soil using minimally invasive remediation solutions. One of them seems to be aided phytostabilisation.

Objectives
The aim of this project was to investigate the impact of organic by-products as heavy metals stabilizers in soil on the biodiversity of microorganisms under ex situ aided phytostabilisation.

Methods
Green waste compost and pulp from the processing of grain were introduced into highly contaminated with Zn, Pb and Cd soil. As a phytostabilizer grass Festuca arundinacea was used. After 18 weeks the activity of enzymes: dehydrogenases, phosphatases and urease as well as the structural and functional biodiversity of microorganisms were tested. The biomass of grass after crop was also analysed.

Conclusions
The activity of enzymes in soil supplemented with grain was significantly higher in comparison with their activity in soil amended with compost. Simultaneously, the addition of grain into soil caused the highest increase the metabolic activity and biodiversity of microorganisms in comparison with soil amended with green compost. Additionally, after plant cutting the grass achieved the highest biomass in soil with grain.
REMOVAL OF HEXAVALENT CHROMIUM BY OCHROBACTRUM SP. CUST210-1 AND CHROMATE REDUCTASE
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Background
Chromium is extensively used in resistant alloys, electroplating, dye productions and leather tanneries. Hexavalent chromium (Cr⁶⁺) is a toxic, mutagenic and carcinogenic chemical. Physicochemical processes have been applied in Cr⁶⁺ treatment, however, their setup and operating costs are high for large-scale treatment. Based on this consideration, biotechnology is a potential alternative.

Objectives
In this study, we would like to reveal the characteristics of Ochrobactrum sp. CUST210-1 and its crude chromate reductase in removing Cr(VI).

Methods
We isolated chromium-resistant bacterium from leather industry wastewater by chemostat. Its chromate reductase was purified and further identified by SDS-PAGE. Characteristics of bacterial cell and chromate reductase for Cr(VI) removal were evaluated.

Conclusions
A facultative Ochrobactrum sp. CUST210-1 was isolated and it achieved high removal efficiency at relatively low Cr(VI) concentration (<350 mg/L) under the aerobic condition. Cu²⁺, Zn²⁺, Ni²⁺, SO₄²⁻ and Cl⁻ with 10-150 mg/L did not affect Cr(VI) removal by CUST210-1. Thus, the cell technique could be applied in treating chromium-containing wastewater (e.g. leather industry wastewater). By SDS-PAGE analysis, we confirmed that crude chromate reductase was successfully isolated. NADH rather than carbohydrate was optimal e⁻ donors for the activity of chromate reductase. Thus, immobilized enzyme technique could be applied in treating chromium-containing wastewater with low concentrations of organic compounds (e.g. electroplating wastewater). To our knowledge, this is the first report to demonstrate the application range by bacterial cell or immobilized enzyme in removing Cr(VI).
INVESTIGATION OF METABOLICALLY ACTIVE BACTERIA LEVELS AROUND GÖKÇEADA ISLAND (THE NORTHERN PART OF AEGEAN SEA), TURKEY

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Background

Determination of the metabolically active bacteria level in the studied area is essential in terms of describe of bacteria that contribute to the ecological in aquatic ecosystems. Metabolically active bacteria have a well-developed polysaccharide capsule whereas inactive bacteria rapidly release the capsule (the term "capsulated bacteria was used in reference to "capsule bearing bacteria").

Objectives

The aim of this study was to investigate total and metabolically active bacteria levels at the surface sea water samples taken the coastal and off shore areas which have chosen around Gökçeada Island, northern part of Aegean Sea (Fig.1).
Fig. 1. Sampling area

**Methods**

Sea water samples were taken 19 stations around Gökçeada Island (northern part of Aegean Sea) seasonally for the autumn, winter, spring as monthly for the summer in 12 times in total between March 2012 - November 2013. The frequency of the metabolically active bacteria was determined with modified staining technique comparing the number of the intact cells to the total (live and dead bacterial cells) number of bacteria (Plante and Shriwer, 1998; Stoderegger and Herndl., 2001).
**Conclusions**

The highest metabolically active bacteria percentage was determined to be 33.12% in the seawater samples which were taken from Aydincik Bay in 2012 summer and the metabolically active bacteria levels in coastal station was found higher than the off-shore stations.

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**References**


DEGRADATION OF NAPROXEN BY A SELECTED FUNGAL STRAIN

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Background

Pharmaceuticals are widely used for treating human and animal diseases. These active compounds and their metabolites can enter aquatic environments via urine, feces, discharge from production facilities and hospital wastes (Kosjek et al. 2005). Naproxen and its sodium salt are members of the α-arylpropionic acid group of non-steroidal anti-inflammatory drugs. Due to excessive usage of naproxen, this drug has been detected in aquatic environments and even in drinking water (Marco-Urrea et al. 2010).

Objectives

The principal aim of this research was assessment of naproxen degradation capabilities of four fungal strains, include two white-rot fungi (Phanerochaete chrysosporium, Funalia trogii), yeast Yarrowia lipolytica, filamentous fungus Aspergillus niger and identification of byproducts.

Methods

The LC/MS spectra were taken on a Waters Micromass ZQ connected with Waters Alliance HPLC, using ESI (-) method, with C-18 column. For the structural elucidation of isolated metabolites, low resolution LC-MS and NMR spectroscopy have been used. ¹H (400 MHz) was recorded employing a VARIAN MERCURY 400 MHz FT spectrometer.

Conclusions

Aspergillus niger found to the most efficient strain with %98 degradation rate. Two main byproducts of degradation, O-desmethylnaproxen and 7-hydroxynaproxen were identify using LC/MS and ¹H-NMR.

References

THE INFLUENCE OF ENDOPHYTIC BACTERIA IN WHEAT PLANTS FERTILIZED WITH NITROGEN

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Background
The plant growth-promotion bacteria improve nutrient uptake, reduce fertilization cost and also minimize the environmental pollution by decreasing nitrogen leaching. However, there is not enough data about plant growth-promotion bacteria and grasses interaction, especially with endophyte bacteria.

Objectives
The aim of this work was evaluate the possible benefits of three diazotrophic endophytic bacteria about the role in the nitrogen metabolism, ability to synthesize plant hormones and the influence of different N level on wheat plant.

Methods
Wheat plants were fertilized with three nitrogen levels (no-fertilizer, half recommended N-fertilizer and full recommended N-fertilizer) and treated with different nif H+, endophytic bacteria: IAC-AT-8 (Azospirillum brasilense), IAC-HT-11 (Achromobacter insolitus) and IAC-HT-12 (Zoogloea ramigera). The enzymes nitrate reductase, glutamine synthetase and chlorophyll content analyses were analyzed as well as the IAA production by the strains. Before the harvest the nitrogen and the growth promotion were also evaluated.

Conclusions
All the three bacteria were able to modify the nitrate reductase and the glutamine synthetase activities and improve the chlorophyll content, and promote the root and shoot weight. The strains also produced IAA in pure cells culture. The N content and the N-use efficiency index were dependent on the endophytic bacteria strain and on the plant tissue assessed. In general, the performance of the above parameters was strongly affected by the nitrogen rate. The different responses suggest that the successful colonization and the growth promotion are achieved by distinct mechanisms.
ANTIBIOTICS PROMOTE AGGREGATION WITHIN AQUATIC BACTERIAL COMMUNITIES
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Background

The release of antibiotics (AB) into the environment poses several threats for human. Knowledge on the impact of AB on natural bacterial communities is missing both in terms of spread and evolution of resistance mechanisms, and of modifications of natural community composition and productivity.

Objectives

We performed a chemostat-based experiment with 4 coexisting bacterial strains mimicking a freshwater bacterial community to study their response to antibiotics in low and high doses.

Methods

Bacterial abundance rapidly decreased by 75% in the presence of AB, independently of their concentration, and remained constant until the end of the experiment. The bacterial community was mainly dominated by \textit{Aeromonas hydrophila} and \textit{Brevundimonas intermedia} while the other two strains, \textit{Micrococcus luteus} and \textit{Rhodococcus} sp. never exceed 10%. Interestingly, the bacterial strains, which were isolated at the end of the experiment, were not AB-resistant; in addition reassembled communities composed of the 4 strains, isolated from treatments under AB stress, significantly raised their performance (growth rate, abundance) in the presence of AB compared to the communities reassembled with strains isolated from the treatment without AB. By investigating the phenotypic adaptations of the communities subjected to the different treatments, we found that the presence of AB significantly increased co-aggregation by 5-6 fold.

Conclusions

These results represent the first observation of co-aggregation as a successful strategy of AB resistance based on phenotype in aquatic bacterial communities, and can represent a fundamental step in the understanding of the effects of AB in aquatic ecosystems.
ISOLATION AND CHARACTERIZATION OF THERMOTOLERANT PETROLEUM-OXIDIZING BACTERIA FROM GEOGRAPHICALLY REMOTE REGIONS

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Background
Bioremediation of oil-polluted ecosystems in the hot climate regions can be performed by microbial consortia of thermotolerant bacteria.

Objectives
The objective of our work was to isolate thermotolerant bacteria from soil and water samples, physiological and biochemical analysis and taxonomic characterization of the bacteria.

Methods
In the work we used such methods as bacterial cultivation, DNA isolation, PCR, electrophoresis in agarose gel, direct sequencing, phylogenetic analysis, IR-spectroscopy.

Conclusions
Among 85 petroleum-oxidizing bacteria isolated from soil and water samples of Russia, Kazakhstan and Antarctica 15 thermotolerant gram-positive strains have been found which can utilize petroleum and particular hydrocarbons at 45°C and higher. The isolates were identified as the members of genera Gordonia, Paenibacillus and Rhodococcus and were closely related to known species of those genera. The alkB gene (encodes for alkane 1-monoxygenase) has been determined in all Gordonia isolates. Most thermotolerant isolates of genera Gordonia, Paenibacillus and Rhodococcus utilize petroleum in salt-enriched media.

The strains Rhodococcus sp. 5Ap, Rhodococcus sp. Par7 and Gordonia sp. 1D were the most effective thermotolerant petroleum-oxidizing bacteria at 45°C – 27%, 14% and 20% of petroleum, respectively. At the temperature 24°C the most effective petroleum-oxidizing strain was Gordonia sp. 1D – up to 59% of petroleum.
ADAPTATION OF ARTHROSPIRA SP. PCC 8005 TO NITROGEN STARVATION: UNEXPECTED EXPRESSION OF HETEROCYST-LIKE PROTEINS

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Background
Among the most spread microorganisms on Earth, cyanobacteria play a dominant role in nitrogen and carbon cycles. The main limiting factor in cyanobacterial growth results from essential nutrients deprivation (e.g., nitrogen starvation). To deal with that, cyanobacteria develop survival strategies, including morphological and metabolic changes.

Objectives
This study aims to investigate the morphological and metabolic changes within Arthrospira sp. PCC 8005 under N starvation.

Methods
In order to characterize the biological response of this strain, a multidisciplinary investigation was carried out (i.e., proteomic, electronic microscopy, intracellular compounds assay,...).

Conclusions
Arthrospira sp. PCC 8005 underwent to metabolic reprogrammings of C and N metabolisms. Proteomic investigations suggested a degradation of the phycobilisomes, cyanophycine and potential endogenous N sources (e.g., nitrile, urea, cyanates & formamide) to counteract the lack of N in the medium. As N metabolism is regulated thanks to N-to-C ratio, N starvation results in C over-excess, which was stored as glycogen as indicated by proteomic and transmission electronic microscopy investigations. TEM confirmed proteomic results as shown in the decreased of thylakoids, cyanophycin, and in glycogen granules accumulation. Arthrospira sp. PCC 8005 exhibited a survival capacity during long-N starvation, likely by maintaining a basic activity in cells named HFN⁺, probably to sustain colonies viability. Interestingly, proteomic results showed an increased of heterocyst-like proteins (HetR & HglK), while the colonies are non-heterocystous. In order to characterize the role of these heterocyst-like proteins during N starvation in our strain, we carried out immunolocalization of HetR, and studied its potential protein partners by co-immunoprecipitation.
ANALYSIS OF EXTRACYTOPLASMIC FUNCTION SIGMA FACTORS IN THE PLANT GROWTH-PROMOTING RHIZOBACTERIA BURKHOLDERIA PHYTOFIRMANS PSJN

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Background
Nutrient availability makes plant internal tissues and rhizosphere attractive spaces for microbial colonization. However, bacteria interacting with plants need to adapt to constant environmental changes, reacting to fluctuations in their immediate surroundings through global gene regulation. Sigma factors are dissociable subunits of RNA polymerase that regulate gene transcription initiation by recognition of specific promoter sequences. Their total number in the genome of a specific bacterial strain is usually determined both by its lifestyle and its genomic size. The extracytoplasmic function sigma factors (ECFs) are the largest and most diverse group of these factors (classified into 43 subgroups), but little is known about their functions, although they are predominant in environmental and plant associated bacteria, which makes them interesting candidates for global gene regulation during plant colonization.

Objectives
The aim of this study was to inactivate and to characterize ECFsf present in the plant growth-promoting rhizobacteria Burkholderia phytofirmans PsJN.

Methods
This strain contains 18 ECFsf, which were inactivated by insertional mutagenesis.

Conclusions
Results suggested that at least one ECFsf could be related with growth and general metabolic processes in strain PsJN, meanwhile others ECFsf may be specifically involved in oxidative stress tolerance and biofilm formation, processes related to plant colonization by bacteria. In general, the results support a possible role of ECFsf in plant bacteria interactions at the level of the rhizosphere.

MELANIN SYNTHESIS BY BACILLUS WEIHENSTEPHANENSIS STRAINS ISOLATED FROM SOIL IN NORTHEASTERN POLAND
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Background
Melanin is known for protecting living organisms from harmful physical and chemical factors, however, its synthesis is rarely observed among Bacillus cereus sensu lato. So far, only two melanin-positive wild strains, B. thuringiensis subsp. dendrolimus [1] and B. thuringiensis subsp. kurstaki [2], have been described, but melanin production by B. weihenstephanensis has not been reported yet.

Objectives

B. weihenstephanensis isolates originated from soil in Northeastern Poland were investigated in order to assess physicochemical properties of a blackish-brown pigment they produced and the mechanism of its synthesis in relation to their genotypic and phenotypic characteristics.

Methods

Phenotypic and genotypic investigation of the melanin producers were done using API system, Multi-locus Sequence Typing, 16S rRNA sequencing, and Pulsed-Field Gel Electrophoresis. Fourier transform infrared spectroscopy we used to compare produced pigment with commercial melanin. The mechanism of the pigment production was assessed by biochemical tests and the whole genome sequencing of one B. weihenstephanensis strain.

Conclusions
For the first time, we report that psychrotolerant B. weihenstephanensis can produce melanin pigment, which is probably associated with laccase activity. These environmental isolates are closely related to the B. weihenstephanensis DSMZ 11821 reference strains. The ability of melanin synthesis by soil B. weihenstephanensis strains seems to be a local adaptation to a specific niche.

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References
MICROBIOLOGICAL ANALYSIS OF THE SURFACE OF THE UNDERGROUND REINFORCED CONCRETE STRUCTURES

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Background
In public buildings and facilities, in areas with high humidity and specific climatic conditions, microbiological corrosion is an important factor affecting the reliability and durability of structures made of metal, concrete and reinforced concrete. Therefore, the problem of potential protection of building materials and structures is very complex and actual.

Objectives
Estimation of the degree of contamination of reinforced concrete structures of Almaty subway by corrosive dangerous microflora.

Methods
Isolation of corrosive dangerous microorganisms was performed by seeding selected water samples and swabs of concrete materials on selective media.

Conclusions
7 stations of Almaty subway were examined in spring 2014. 31 samples were selected, including 5 water samples and 26 swabs from different visually damaged surfaces of reinforced concrete structures. Studies have shown that pH of the water samples and scrapings were neutral or alkaline. Heterotrophic bacteria, filamentous fungi and denitrifying microorganisms were found in almost all samples. Their number was high enough. This is an alarming fact, because of their intensive development can contribute to a change in environmental conditions that are favorable for the development of thione and sulfate-reducing bacteria, which are corrosive dangerous microorganisms. Further, these group of microorganisms may also participate in the destruction of the concrete.

Thione and sulfate-reducing bacteria were detected in small amounts. However, their presence shows that they can grow intensively in creating the favorable conditions and thereby contribute to the development of corrosion processes in reinforced concrete structures.
THE DIVERSITY OF MICROBIAL COMMUNITIES AT ACIDIC MINE WATER OUTLETS IS DRIVEN BY STOCHASTIC PROCESSES

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Background

Explanation how the vast microbial diversity with multiple concurrent organisms may persist at a single site is still elusive for microbial ecology. One of plausible hypotheses is that a large number of spatially separated microcommunities exist within each complex habitat. Thus, each real assembly consists of a small and largely random selection of organisms from the total observed diversity.

Objectives

One of plausible ways to test these hypotheses is to compare microbial communities from homogeneous microhabitats that exist in multiple physically separated copies at a single site. We selected ferruginous stalactites growing in large groups at acidic mine drainage outlets as a model of such microhabitats.

Methods

Two groups of microbial stalactites growing at a single acidic mine drainage outlet and six stalactites from other sites were sampled. Prokaryotic 16S rDNA was amplified using universal primers and microbial diversity was characterized by tRFLP and amplicon pyrosequencing.

Conclusions

Both methods showed that simple and homogeneous microbial communities inhabited all stalactites, but species-level OTU composition substantially differed even between adjacent stalactites. Although the sampling sites influenced composition of microbial communities, all OTUs including the most abundant ones unpredictably alternated in all samples. As a result, a large portion of communities clustered independently of locality or other variables by both phylogenetic beta diversity and OTU abundance metrics. Thus, each community probably contains a random selection from the group of plausible species. The environmental traits influence the probability of the presence of individual species but do not determine composition of simple microbial assemblies.
THE MICROBIOME OF THE ESTERO SALADO MANGROVE: A 16S TAG APPROACH TO DETERMINE DIFFERENCES IN BACTERIAL COMMUNITY, ON SEDIMENTS CONTAMINATED BY HEAVY METALS.

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Background
Mangrove ecosystems are coastal estuarine systems confined to the tropical and subtropical regions. The Estero Salado mangrove located in Guayaquil, Ecuador, has suffered constant disturbances during the past 20 years, due to industrial wastewater release. Previous work determined that this estuary is one of the most disrupted on earth due to its relationship with anthropogenic disturbance.

Objectives
To determine the concentration of heavy metals and nutrients in surface sediments of the Estero Salado in equatorial summer season (February).

To determine the influence of high concentrations of heavy metals in terms of composition of the bacterial microbiome.

Methods
A proper understanding of the spatial variations of microbial communities will provide clues about the underline mechanisms that structure microbial groups. In the present study, the analysis of concentration of heavy metals, nutrients and tag sequencing of 16S rRNA V4 was conducted in surface sediments in two mangroves areas in Guayaquil. The first one under the influence of heavy metals and other pollution free

Conclusions
Using the high throughput sequencing method, revealed a detailed picture of the spatial variations of the bacterial community structure in the superficial sediment on Estero Salado mangrove. Heavy metals directly influence on bacterial community due to in the contaminated area had a relatively lower alpha-diversity than the area without its presence. We were able to identify bacterial groups significantly enriched in specific locations. This is the first approach in a mangrove area using tag sequencing and correlating with heavy metals levels.
BEHAVIOUR OF SOME BACTERIA ISOLATED FROM CITRUS RHIZOSPHERE INFESTED BY TYLENCHULUS SEMIPENETRANS, IN TUNISIA

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Background
The cape Bon area (Tunisia) is the world leading producer and exporter of Maltese half blood oranges. However the sour orange (Citrus aurantium) rootstock is naturally infested by a phytopathogen nematode, Tylenchulus semipenetrans, a major nematode pest. Some bacteria and nematophagous fungi demonstrated a nematicidal effect.

Objectives
Only few studies have been conducted on the interaction between bacteria and nematodes in the rhizosphere of citrus trees. The aim of this study is to better understand the role of some bacteria in the complex microbial ecosystem of the Tunisian Citrus rhizosphere.

Methods
Sixty bacterial strains were isolated from Citrus root trees and egg masses of nematodes. The isolates were screened for i) their nematicidal activity on eggs and on second stage juvenile of Tylenchulus semipenetrans, ii) their antifungal activity and iii) their effect on the growth of sour orange roots. To identify the isolates rDNA and/or rpoB gene sequencing was carried out.

Conclusions
Among the 60 bacterial isolates, the majority belonged to either Bacillus subtilis or Bacillus cereus. However, the strains showed a great diversity in their effects i) on the plant host Citrus, ii) on the nematode Tylenchulus semipenetrans, iii) on the rhizosphere-associated fungi (i.e. the phytopathogen Fusarium solani and predatory nematodes Monacrosporium cianopagum and Arthrobotrys conoides). Almost 30%
showed antifungal activities against F. solani. These bacterial strains could be used to improve the quality of soil when used in the field by mixing various bacteria, thereby combining antifungal activity and plant growth-promoting bacteria. Large-scale trials should be performed to confirm these promising results.
ICE NUCLEATION ACTIVITY IN THE WIDESPREAD SOIL FUNGUS
MORTIERELLA ALPINA

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Background
Biological residues in soil dust are a potentially strong source of atmospheric ice nucleators (IN). So far, however, the sources and characteristics of biological - in particular, fungal - IN in soil dust have not been characterized.

Objectives
Thus, the objective of this study is a regional investigation of the identity and relative abundances of culturable INA fungi in topsoils, an essential base for improving our understanding of the effects of microorganisms on climate and the hydrological cycle.

Methods
Soil fungi, from a range of different land use and ecosystem types in south-east Wyoming, were analyzed for ice nucleation activity. Ice nucleation active (INA, i.e., inducing ice formation in the probed range of temperature and concentration) fungi were found to be both widespread and abundant, particularly in soils with recent inputs of decomposable organic matter. Across all investigated soils, 8% of fungal isolates were INA. All INA isolates initiated freezing at -5°C to -6°C, and belonged to a single zygomycotic species, Mortierella alpina (Mortierellales, Mortierellomycotina). The IN produced by M. alpina seem to be proteinaceous, <300 kDa in size, and can be easily washed off the mycelium.

Conclusions
Ice nucleating fungal mycelium will ramify topsoils and probably also release cell-free IN into it. If these IN survive decomposition or are adsorbed onto mineral surfaces, these small cell-free IN might contribute to the as yet uncharacterized pool of atmospheric IN released by soils as dusts.
DEVELOPMENT OF GREEN ALTERNATIVE COATINGS TO PROTECT THE METALLIC STRUCTURES AGAINST THE MICROBIAL INDUCED CORROSION PROCESS

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Background
Microbial Influenced Corrosion (MIC) is a very aggressive form of corrosion with many proposed mechanisms for its prevention, but existing antifouling solutions include not environmentally friendly biocides. This fact has caused an urgent demand for greener, non-toxic or low-toxicity and longer lasting antifouling compounds and technologies.

Objectives
BIÓCORIN project, within FP7-ENVIRONMENT program, aims to develop a green alternative to the coatings and solutions used up to date for MIC corrosion protection and prevention in infrastructures.

Methods
The project is based on a biological phenomenon commonly occurring in nature for microbial population regulation. Microorganisms secrete compounds that inhibit the growth of other microorganisms due to existing antagonism between microbial populations. First step was the isolation and identification of those microorganisms that play a major role in fouling and MIC for different environmental conditions. Several samples were collected from different climate locations for the isolation and identification of microbial populations from metal surfaces. As a result, several microorganisms belonging to bacteria, fungi and yeasts were isolated from the four
different samples and were identified by means of 16S and 18S rRNA gene sequencing.

Conclusions
The search of microorganisms with anti-MIC properties resulted in the identification of two candidates with the ability to synthesize compounds with antifouling properties. These candidates were integrated in a coating based on sol–gel technology to prevent the MIC corrosion phenomena in a green way. Currently, this technology is under evaluation by means of three case studies with different environmental conditions.
METABOLOMЕ ANALYSIS OF THE ANTIMICROBIAL COMPOUNDS SECRETED BY ENVIRONMENTALLY ISOLATED ANTI-MIC MICROORGANISMS

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Background
Corrosion is a worldwide problem and the direct economic loss due to metal corrosion of infrastructures and equipments is estimated to exceed €1.32 trillion per year, which accounts for 3 to 4% of the Gross Domestic Product of industrialized countries. Among corrosion types, Microbial Induced Corrosion (MIC) is involved in at least 10% of the corrosion of structures, and up to 50% in the case of subterranean pipes.

Objectives
BIOCORN (New Biocoating for Corrosion Inhibition in Metal Surfaces) is a project funded by the European Union under the activities of the Seventh Framework Programme (FP 7-ENVIRONMENT) that aims to develop a green alternative to the coatings and solutions used up to date for MIC corrosion control. Some of the results of this project are presented here, with the identification of several relevant antifouling compounds secreted by environmentally isolated anti-MIC strains via a metabolomic approach.

Methods
Metabolites were extracted from three anti-MIC strains. 100 ml cultures incubated in Terrific Broth (TB) at 30ºC for 24 h in the presence and absence of a pool of MIC extracts (inducers) originated from several climatic locations around Europe. Each culture broth was recovered and analyzed by means of LC-MS and GC-MS (Acquity UPLC - SYNAPT G2 HDMS system QToF with ESI injection and a Thermo Scientific Focus GC - DSQ II respectively).

Conclusions
The metabolomics analysis pointed out the differential expression of 2 compounds by GC-MS and more than 10 compounds for LC-MS (positive mode) between the analyzed anti-MIC strains.
DETERMINATION OF FUNGUS PRESENT IN THE AIR TAKEN IN PM10 FILTERS TWO NETWORK STATIONS MONITORING AIR QUALITY IN THE VALLE DE ABURRÁ (MEDELLÍN – COLOMBIA)

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Background

Air quality is an issue that has come to be of great concern in recent years due to increased production of particulate material. The presence of fungi in the air is partly responsible for some allergies, infections and diseases. Because of this there is a need to study fungal species associated with inhalable particulate matter and its effects on health.

Objectives

The aim of the research is to isolate the fungi present in PM10 filters in sampling two Network Monitoring Air Quality in Aburrá Valley, which presented the highest levels of pollution in the city. After isolation fungi are identified and characterized by the types of pathogens.

Methods

The microorganisms are absorbed into quartz filters that capture particles diameters smaller than 10 microns (PM10). The filters are exposed to atmospheric air for 24 hours in the HI-VOL PM10 equipment. Subsequently, the filter was incubated in enrichment medium, serial dilutions were made and plated on selective media for fungi (PDA and Sabouraud in this case). After growth, morphological identification is performed.

Conclusions

The results of the tests reveal the presence of \textit{Cladosprium} sp., \textit{Fusarium} sp., \textit{Alternaria} sp. y \textit{Rhodotorula} sp. These microorganisms are considered allergens and are associated with the generation of respiratory diseases like asthma, rhinitis, tracheitis, bronchitis, pneumonia, bronchopulmonary mycosis, among others. These findings demonstrate the importance of knowing the microorganisms associated with particulate material breathe as increased air pollutants in the atmosphere increases morbidity and mortality.
DIFFUSION SANDWICH SYSTEM, A TOOL TO ACCESS THE MICROBIAL DARK MATTER
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Background

Prokaryotes are the most abundant and diverse life form on Earth. However, to date, we know that only between 0.001% and 1% of total environmental bacteria have been cultivated through standard culture techniques and media, remaining then a rich diversity of bacterial taxa to be isolated and cultured, the so-called microbial dark matter.

Objectives

The aim of this study was to carry out a proof of concept of a system of miniature diffusion chambers which we have named Diffusion Sandwich System (DSS), as a cheap and user-friendly tool to gain access to previously uncultured bacterial taxa.

Methods

We did a comparative study of the cultivability and bacterial diversity which could be isolated by using the DSS and with the classical approach based on picking up bacterial colonies from Petri dishes. Further both culture-dependent approaches were compared with the bacterial diversity present in the same ecological niche but analyzed in-depth by pyrosequencing the metagenomic 16S rRNA gene.

Conclusions

The culturability obtained by using the miniaturized chambers was 2-log10 units higher than with the classical approach. Unlike the classical approach, the relative abundances of taxa isolated with the DSS was in line with those actually present in the ecological niche, allowing in fact to isolate representatives of several underrepresented phyla such as Acidobacteria, Verrucomicrobia and Gemmatimonadetes, which constitute new taxa into these phyla.
Background
Little is known about the factors determining the eligibility of marine phages as specific markers of hydrological flow and reactive transport of colloidal particles in the Earth’s Critical Zone (CZ). Marine phages and their bacterial hosts are naturally absent in soil, can be applied as tracers at high concentration and be detected as little as one phage/mL in water.

Objectives
Within the DFG Collaborative Research Center – AquaDiva we evaluated the effects of phage characteristics (morphology, surface property), water flow regimes as well as the role of mycelial networks on transport of marine phages in laboratory experiments.

Methods
Phages were characterized by electron microscopy, dynamic light scattering and water contact angle analysis (CA) for their size/morphology, surface charge (ζ) and hydrophobicity. Sand-filled percolation columns and a modified high-throughput plaque assay combined with fluorescence microscopy counting were used to quantify phage deposition during percolation.

Conclusions
Our data show that marine phages exhibit differences in their deposition efficiency depending on their size, surface charge and hydrophobicity. For instance, the bigger (60 nm), more hydrophobic (CA = 78°) and poorly charged (ζ_{TM, pH=7.8} = -13 mV) PSA-HM1 phage exhibited a higher deposition efficiency than smaller (30 nm) and less hydrophobic (CA = 52°) H6/1 phages. Interestingly however, both marine phages showed lower retention than terrestrial T4 phages exhibiting similar characteristics than HM1. We conclude that marine phages have a high potential for the use as sensitive tracers in terrestrial habitats with their surface properties playing a crucial role for their transport.
Background
Pentachlorophenol (PCP) is a hazardous agent in water and soil that poses a significant risk to human health. The toxicity associated with PCP occurs in low doses and mainly affect liver, kidney and central nervous system.

Objectives
The purpose of this study was to investigate the biosorption of PCP on Aspergillus niger biomass as a method for removal of PCP from aqueous solutions.

Methods
The modified A. niger biomass with NaOH was used to adsorb the PCP. The impacts of various experimental parameters such as initial PCP concentration, pH of solution, contact time, and biomass dosage were investigated and optimal conditions were identified. The experimental data were evaluated by various isotherm and kinetic models.

Conclusions
The correlation of contact time, pH and initial concentration with the biosorption of PCP by A. niger biomass was statistically significant (P<0.001). The PCP removal increased with decreasing of solution pH and maximum adsorption was obtained at the pH 3. The equilibrium sorption capacity was increased by increasing initial PCP concentration from 10 to 40 mg/l of 4.23 to 11.65 mg/g, while the PCP removal efficiency decreased from 87 to 55%. Both Langmuir and Freundlich isotherms well described adsorption equilibrium of PCP on A. niger biomass. For all cases, the correlation coefficients for the second order kinetic model were close to one.

A. niger biomass can be used to reduce the toxicity of aqueous solutions containing PCP in acidic pH conditions. PCP concentration in water and soil poses a great hazard to human health, so its removal by fungi is applicable and commercial.
CULTURE-INDEPENDENT APPROACH FOR THE CHARACTERIZATION OF THE BACTERIAL COMMUNITY IN HAEMODIALYSIS WATER

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Background

The microbiological purity of haemodialysis water and dialysis fluid is crucial in the treatment of patients with chronic renal insufficiency. The presence of bacteria and bacterial-derived products (i.e., endotoxins, sphingolipids, oligonucleotides) can contribute to silent chronic inflammation, secondary amyloidosis, pyrogenic reactions and anemia.

Objectives

The main objective was to characterize the microbial community present in the haemodialysis water used to reconstitute the dialysis fluid at the Hospital Universitario Son Espases (Majorca, Spain). Bacterial community was assessed in samples obtained in November 2012 and 2013 to evaluate the evolution of the species colonizing these pure waters.

Methods

Dialysis water was filtered through 0.22 µm pore size filters. DNA was extracted from the filters, purified and was amplified with two different set of primers: i) 16S rDNA universal primers and ii) selective \textit{Pseudomonas} primers designed for the \textit{rpoD} gene. Amplicons were pyrosequenced with a 454 System+ GS FLX platform from Roche. Reads obtained were analyzed in order to assess the phylogenetic identification of bacteria present in the community.

Conclusions

The bacterial community present in the dialysis water is very complex, with microorganisms highly diverse and adapted to the oligotrophic habitats, not usually detected routinely in clinical microbiology laboratories. This community is maintained in the two years period of this study. Based on the 16S rRNA gene pyrosequencing analysis the \textit{α-Proteobacteria} class is the most abundant. The pyrosequencing of the \textit{rpoD} amplicon showed a high number of putative novel species of \textit{Pseudomonas} showing a high diversity still not described.
Background
Because of their massive utilization, hydrocarbons are major pollutants of soils and aquifers. Biodegradation is a key aspect of the fate of pollutants in the environment.

Objectives
It is based on the Gas Chromatographic and Thin Layer Chromatography (TLC) coupled with a Flame Ionization Detector (FID) analysis of hydrocarbons, after incubation in optimal conditions of major compounds in crude oil (saturates and aromatics fractions).

Methods
Using a method of gene sequencing 16S rDNA, the identification of bacteria adapted to soil was determined. Based on nearly full length 16S rRNA gene sequencing analysis, a phylogenetic trees was constructed. Capacity of degradation of selected bacteria and the presence or absence of functional genes coding for the initial oxidation of hydrocarbons (alkB, nahAc, nidA) were studied.

Conclusions
The results indicate that the Gamma-Proteobacteria group was the main actor of this degradation. Pseudomonas sp, specific bacterial group suitable for petroleum hydrocarbon pollution has the ability to metabolize (single culture) a high mass residues, Pristane (n-C19) at 35.11% and Benzo[a]Pyrene (n-C20) at 33.93% and co-metabolize (in consortium) fractions in petroleum-hydrocarbon, 50.44% of saturates and 30.42% of aromatics compounds. Biodegradation efficiency of soil microflora depended on the specific metabolic pathways of some microorganisms and on the cooperation within microbial population. This study provides a better understanding of the adaptation of bacteria inhabiting polluted environments and for developing and implanting adequate bio-strategies in the future to enhance oil in contaminated soil and the biotreatment of oily waste water in refineries.
EXTRACTIVE MEMBRANE BIOFILM REACTORS (EMBFR) FOR BIOREMEDIATION OF METHYL TERT-BUTYL ETHER CONTAMINATED GROUNDWATER: A LAB-SCALE STUDY.

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Background
Among the strategies developed for contaminated groundwater bioremediation, those based on the use of bacteria adhering to inert supports and established as biofilms, have gained great importance in this field. The EMBFR technology offers productive solutions for the removal of volatile compounds like methyl tert-butyl ether (MTBE) and other contaminants from water bodies. The EMBFR is based on extractive semipermeable (silicon) membranes through which the contaminants migrate to the biological compartment in which microorganisms with MTBE biotransformation and/or mineralization capacities grow in mineral salts medium, forming a catalytic biofilm on the membrane surface.

Objectives
The objective of this study was to assess the use of three bacterial strains (Paenibacillus sp. SH7 CECT 8558, Agrobacterium sp. MS2 CECT 8557, and Rhodococcus ruber EE6 CECT 8612) previously isolated from gasoline contaminated environments, and used as inoculum in a lab-scale EMBFR running for 28 days under aerobic conditions.

Methods
Three different retention times (1h, 6 h, and 12 h) were employed. Degradation rates were determined daily by gas chromatography - mass spectrometry, and the biofilm established by the bacterial strains on the semipermeable membrane was observed by FESEM at the end of each experiment. Acute toxicity of the effluents was determinate by Microtox® assay (EC₅₀).

Conclusions
The results achieved from MTBE degradation, biofilm formation and toxicity analysis showed that these bacterial strains could be excellent candidates for use as selective inocula in EMBFR technology for MTBE bioremediation.
DIARRHEAL BACTERIAL PATHOGENS ARE FREQUENT IN THE CHOQUEYAPU RIVER IN BOLIVIA, AND POSE A RISK OF TRANSMISSION TO FRESH PRODUCE BY CONTAMINATED IRRIGATION WATER

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Background
Water borne diarrheal pathogens might accumulate in river water and present a potential contamination risk of drinking and irrigation water.

Objectives
In this study, we show how pathogens can accumulate and be transported in the river Choqueyapu in La Paz and contaminate fresh produce irrigated by river water.

Methods
Using qPCR absolute quantification methods, we found a seasonal and spatial variation in the number of pathogen gene copies per ml of river water with a high peak of Campylobacter ssp. that contaminated otherwise clean water in a watershed at the Altiplano plateau, after the peak of the rainy season in February 2014. High levels of Campylobacter, Salmonella and enterotoxigenic E. coli (ETEC) were subsequently isolated in sampling points downstream along the river and finally identified in vegetables rinse water.

Conclusions
These results indicate that distant water contaminations may be transported all the way to the agricultural area. In addition, we found that ETEC carrying the heat labile toxin (LT) alone was much more prevalent in water than other ETEC carrying the heat stable toxin (ST). Hence ETEC with different toxin profiles might have different reproduction and survival ability in water. The results show that the absolute levels of several diarrheal pathogens are high in contaminated river water in the central part of La Paz and although this water is not accessible by humans at this spot several of the pathogens end up in the downstream river water used for irrigation of crops where they might contaminate vegetables.
STUDY ON O, M, P-XYLENE DEGRADATION AND ITS RELATED GENES IN RHODOCOCCUS SPP. J7
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Background

Xylene, an aromatic hydrocarbon, can be easily found in gasoline contaminated soil and composed of three isomers [ortho (o-), meta (m-) and para (p-) xylene]. Many bacterial strains have been reported to grow on omp-xylene isomers, but few studies have been performed to degrade them simultaneously. Xylene is known for its wide usage in tissue processing, staining and cover slipping and can find easily in groundwater. Due to volatile property of xylene isomers, they can spread easily from groundwater to the air.

Objectives

Once human is exposures to xylene, central nervous system could be damaged. So we tried to decrease amount of xylene emitted from factory using soil bacteria.

Methods

Strain J7, belong to the genus Rhodococcus was isolated from total petroleum hydrocarbon contaminated site in Korea. Xylene degradation ability was evaluated using 90ppm of xylene mixture (30 mg/L individually) and analyzed by Agilent GC 6890N with Flame ionization detector (FID). The metabolites are analyzed using GC-MS and the pathway of degradation was construed. Target was amplified by PCR using genomic DNA and PCR products were sequenced at Marcrogen (Korea) and compared with other reference genes.

Conclusions

From this study, we confirmed that strain J7 can degrade o, m, and p-xylene at the same time and based on the data, we have constructed that it follows TOL pathway with high efficiency. The catechol-1,2-dioxygenase gene was amplified and conforming that it followed ortho-cleavage pathway.
Keratin azure and feather degradation in soil by some fungi using singly and in combination

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Keratin azure and feather degradation in soil by some fungi using singly and in combination

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Abstract

The large amount of keratinous waste produced and their localized accumulation create a serious disposal problem leading to environmental pollution. Keratin in forms of animal waste such as hairs, nails, horns, hooves and feathers ultimately comes to soil and if it remains there make pathogenic reservoirs because it is degraded by fungi which have pathogenic potential. On the other hand keratin when degraded by fungi nitrogen is slowly released in the soil and may be utilized by plants as manure. Keratinophilic fungi frequently occurring in soil were selected for hen feather and keratin azure degradation individually and in combination in different sets in soil. Five selected fungi Acremonium implicatum, Chrysosporium queenslandicum, Chrysosporium pannicola, Malbranchea pulchella and Verticillium lecanii, when used in combination with Chrysosporium keratinophilum showed feather and keratin azure degradation. Further feather and keratin azure degradation was monitored in soil when these were used individually and in combination with C. keratinophilum and almost above 90% degradation was recorded. The determination of amount of keratin degradation in soil by using individual fungus and fungi in combination is of immense importance in order to find out how much keratin is degraded in soil. The measurement of feather and keratin azure degradation in soil by using five fungi singly and in combination with C. keratinophilum presented here for the first time.
LOW BIOMASS DNA AND RNA EXTRACTION IMPROVES BY USING G2 BLOCKING REAGENT WITHOUT ANY BIAS IN METAGENOMIC ANALYSIS

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Background
Low biomass samples can be extremely challenging when extracting DNA and RNA. Many scientists have been adding various sources of RNA or DNA to the samples to aid a high recovery, however this makes the samples only useable for specifically directed analysis, while metagenomic analysis is hampered.

Objectives
A new product G2 that are shown to improve DNA and RNA yield from low biomass samples are tested for use in metagenomic analysis as well as if several labs can repeat the results in a ring test.

Methods
We have included analysis of influence of G2 on both full genome sequencing and various amplicon sequencing techniques. Further a Q-PCR analysis was performed after 12 laboratories in a ring test were applying G2 to one sample type.

Conclusions
G2 was shown to be free of contaminating DNA both using 16SrDNA based Q-PCR and amplicon sequencing, further a metagenomic analysis clearly indicated that no traces of the original DNA could be found in a metagenome analysis. The ringtest showed that the participating laboratories all had a higher yield using G2 but also that the participating laboratories did perform very different.
ANTIMICROBIAL ACTIVITIES AND BIOLOGICAL DIVERSITY OF CULTURABLE ACTINOBACTERIA ISOLATED FROM FRESHWATER FISH GUT MICROBIOTA

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Background
Terrestrial microorganisms, especially those from the order Actinomycetales, formed the backbone for the golden age of antibiotic discovery, and are the source of the vast majority of FDA-approved antimicrobial agents.

Objectives
In this study, the antimicrobial activity and diversity of genes associated with secondary-metabolite biosynthesis of Actinobacteria isolated from fresh fish gut microbiota namely *Schizothorax zarudnyi* and *Schizocypris altidorsalis* were investigated.

Methods
Strains of six groups of fish and human pathogenic bacteria were selected and all the isolated actinomycetes were tested for their antibacterial activity against the pathogenic bacteria The antagonistic activity of actinomycetes was tested by the perpendicular cross streak method.

The PCR-based approach was applied to detect the presence of three important biosynthetic genes including polyketide synthases (PKS-I and PKS-II), non-ribosomal peptide synthetase (NRPS). The PCR amplicons and their origins were further confirmed by sequencing.

Conclusions
According to results all isolates possess at least two types of the investigated biosynthetic genes. This study demonstrates the significant diversity of genes associated with secondary-metabolite biosynthesis of Actinobacteria in the fish gut microbiota and it’s potential to produce biologically active compounds.
EFFECTS OF NITROGEN FERTILIZER ON NITRITE-DEPENDENT ANAEROBIC METHANE OXIDATION BACTERIA IN A FLOODED RICE PADDY

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Background
Nitrite-dependent anaerobic methane oxidation (n-damo) may be an important process in the biogeochemical cycles of carbon and nitrogen. However, the inquiry of n-damo bacteria in rice paddy is rarely explored.

Objectives
The aim of this work was to investigate the effects of nitrogen fertilizer on the communities of n-damo bacteria in a flooded rice paddy.

Methods
Different amounts of urea (0, 45, 90, and 180 kg ha⁻¹) were applied to a flooded rice paddy transplanted with a rice cultivar (Oryza sativa) and soil cores were sampled after 15 days of third urea application. The soil cores were divided into top (0–5 cm), middle (10–15 cm), and bottom (20–25 cm) layers and the diversity and abundance of n-damo were investigated using T-RFLP (Terminal-Restriction Fragment Length Polymorphism) and quantitative PCR approaches, respectively. A clone library was constructed for the T-RFLP analyses of 16S rRNA gene fragments of n-damo. DNA samples were retrieved from the soil core samples and the abundances and diversities of n-damo were investigated using T-RFLP.

Conclusions
Abundance of n-damo in a flooded rice paddy was not significantly different although different amounts of urea were applied. However, the phylogenetic analysis of Methylomirabilis oxyfera-like 16S rRNA gene sequences revealed that n-damo was a little diverse depending on urea application and soil depths. Sequences of Top 90 soil sample fell into just one group, while sequences of Bottom 180 were classified into three groups. In addition, the diversity analysis of n-damo in response to different amounts of urea will be performed and discussed.
MICROFLUIDIC CHIP-ASSISTED DNA PREPARATION FROM ENVIRONMENTAL SAMPLES SPIKED WITH FRANCISELLA TULARENSIS AND BACILLUS THURINGIENSIS

Background

The detection of bacteria from environmental samples using nucleic acid amplification depends on efficient DNA extraction and purification to remove PCR Inhibitors.

Objectives

DNA extraction and purification from spiked air, natural water, swab and soil samples was performed using a microfluidic chip device. A protocol was established that was suitable for the detection of two representative bacteria, Francisella (F.) tularensis (Gram-negative) and Bacillus (B.) thuringiensis (Gram-positive).

Methods

The procedure is composed of thermal, chemical and enzymatic lysis methods combined with on-chip DNA purification using paramagnetic particles for reversible DNA attachment. Polymeric microfluidic chips with rhombic cavities were mounted on a device with a regulated heating unit and a movable magnet that enables particle mixing for careful washing and elution. Serial 10-fold dilutions of the bacteria suspensions were tested. DNA preparations were analysed by real-time PCR assays to determine the detection limits that can be achieved.

Conclusions

Detection of both species of bacteria was possible from all tested matrices. The detection limits in air samples corresponded to 7 GE/10 l for F. tularensis and 0.7 GE/10 l for B. thuringiensis. In river water the limits were $10^2$ GE/ml B. thuringiensis and $10^3$ GE/ml F. tularensis. The achieved detection limits for soil samples corresponded to $2.7 \cdot 10^3$ GE/mg F. tularensis and $2.7 \cdot 10^2$ GE/mg B. thuringiensis and
with swab samples $4 \cdot 10^3 \text{ GE/cm}^2$ \textit{F. tularensis} and $4 \cdot 10^2 \text{ GE/cm}^2$ \textit{B. thuringiensis} were detected. Future developments will focus on the development of integrated lab-on-a-chip system combining sample preparation with on-chip PCR technologies.
PLANT GROWTH PROMOTING RHIZOBACTERIAL EFFICACY IN COWPEA (VIGNA UNGUICULATA (L.) WALP.)

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Background

Plant growth promoting rhizobacteria (PGPR) are known to influence the growth of the plant by various plant growth promoting activities such as Indole Acetic Acid (IAA) production, Phosphate solubilization activity, Ammonia production (NH3), Nitrogen fixation, Siderophore production and Hydrogen cyanide production (HCN). In search for the potential PGPR from the Rhizosphere soil of cowpea, a total of 16 rhizobacteria (B1-B6 & D1-D9) were isolated. By comparing the results two potential rhizobacteria (B6 & D4) have been selected for the further evaluation of Plant Growth Promoting (PGP) activity. These isolates biochemically characterized, B6 and D4 confirmed as Pseudomonas and Bacillus by 16SrRNA gene sequencing. Invitro and Invivo Nursery field studies revealed that 100% seed germination rate using Bacillus safensis and 92% for Pseudomonas aeruginosa showing promising results. The study showed the characters of benefits to use these organisms as biofertilizers for sustainable agriculture to improve crop yield.

Objectives

This study was aimed to focus on the PGP activities of the two potential rhizobacteria and growth promotion

Methods

IAA production (Lwin et al. 2012)

Siderophore production (Arnow 1987)

Phosphate Solubilization Activity (Qureshi et al. 2012)

Hydrogen Cyanide Production (Ahmad et al., 2008)

Ammonia Production (Cappucino and Sherman, 1992)

Nitrogen fixation activity (Lwin et al. 2012)

16S rRNA Sequencing

Seed germination rate and seedling vigor index (Ng LC et al. 2012)
Conclusions

Rhizobacteria isolated from the cowpea rhizosphere *Pseudomonas aeruginosa* VRKK1 and *Bacillus safensis* VRKK2 acts as the potential PGPR. *Pseudomonas aeruginosa* showed 100% germination rate, *Bacillus safensis* showed 92% germination.
PROTEOMICS REVEALS OSMOTIC STRESS ADAPTATIONS OF SHEWANELLA BALTICA, PARACOCCUS SP. AND FLAVOBACTERIUM SP. IN A UNIQUE ENVIRONMENT OF THE BALTIC SEA

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Background

Baltic Sea constitutes a unique environment of brackish water conditions, with dynamic, regional shifts in salinity, mainly due to varying freshwater inflows.

Objectives

The goal of the study was to identify physiological and molecular strategies that allow Baltic Sea bacteria to confront this challenging habitat.

Methods

Three bacterial Baltic Sea representatives, identified as Shewanella baltica, Flavobacterium sp. and Paracoccus sp. on the basis of 16S rRNA sequencing and MALDI-TOF analyses, were cultured under controlled growth conditions in a chemostat system and analyzed with a combined approach of 2-DE, multiplex fluorescent staining of proteins in gels and LC-ESI-MS/MS and/or MALDI-TOF/TOF-driven protein identifications.

Conclusions

Tracking down of proteome dynamics in Shewanella baltica, Flavobacterium sp. and Paracoccus sp. in response to the salinity shifts characteristic of the Baltic Sea (2, 7 and 20‰) allowed the identification of a subset of proteins, both of the general stress response group and of other functional categories, differentially expressed under osmotic challenges tested. Wide variations in adaptations to osmotic stress were revealed in the three strains studied, with Paracoccus sp. requiring a significantly less
number of proteins to survive in the osmotic stress conditions in comparison to *Shewanella baltica* and *Flavobacterium sp.*

Taken together, our study uncovered well-coordinated and highly regulated protein inventories in the marine bacteria studied, with a large number of novel candidates for stress response players, thus laying a ground work for understanding of the complexity of microbial adaptive processes in the intriguing ecosystem of the Baltic Sea.
BIOSORPTION OF CHROMIUM BY PSEUDOMONAS SP ISOLATED FROM SOIL CONTAMINATED WITH PETROLEUM OF KHOUZESTAN, IRAN

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Background
In recent decades, the biological methods have been used for remove environmental contaminants such as heavy metals. Resistance to metals is important for this aim.

Objectives
The objective of this research is the isolation of chromium resistant strains.

Methods
Five samples of soil were collected under sterile conditions and transferred to laboratory immediately. Soil samples were homogenized and diluted up to 10⁻¹⁰ with sterile normal saline. These samples were cultured in LB agar containing 5ppm of chromium in order to isolation of resistant strains. These strains were isolated and cultured in Macconkey agar for isolation of appropriate strains. Isolated bacteria were identified by biochemical tests. Then, the MIC test was used for screening of resistant strains. The best conditions of bacterial growth were found in the presence of chromium in various temperatures, rate of shaking and values of pH by spectrophotometry at 600nm in the overnight of cultivation. Metal adsorption test measured under optimal conditions by atomic absorption spectroscopy.

Conclusions: From total of 24 strains of isolated Pseudomonads, 14 strains were resistant to chromium. The top strain (Mac2) has eliminated 35.60% of chromium from aqueous culture mediums in optimal conditions (pH: 8, Temp: 40°C and Shaking rate: 200rpm). The results were verified using statistical method named the general linear model. The contaminated zones should be clean, because heavy metals in various physicochemical forms have been considered as environmental pollutants. This bacterium can be used for studies of bioremediation of contaminated sites to chromium.
MICROORGANISMS IN CAVES: IMPORTANCE, DIVERSITY AND CONSERVATION

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Background

Harsh conditions and isolation lead to formation of unique microbial communities, making caves a promising yet poorly understood field for study of biodiversity. Studies of microbial communities in caves may provide insights of microbial relationships, biogeochemical processes, evolution and even functioning of extraterrestrial life. Furthermore, they can lead to the discovery of novel bioactive compounds and draw attention to the importance of cave conservation.

Objectives

The goal of the project was to analyze the microbial communities of different caves and evaluate the impact of human activity within pristine and disturbed cave areas.

Methods

Barcode pyrosequencing of 16S rRNA genes was used to investigate the microbial diversity of two oligotrophic Karabi Jaila (Crimea) caves and the world’s deepest cave Krubera-Voronja (Caucasus).

Conclusions

It was found that 12 % of 16S rRNA gene sequences from the more frequently visited areas of Crimean caves belonged to previously unclassified bacteria. In the samples of rarely visited branches the sequences belonging to unclassified bacteria comprised 46 %. On the other hand, only 5 % of sequences found in more pristine areas of Krubera-Voronja cave belonged to unclassified bacteria. 6 % of unclassified sequences were found in the main branch of the cave where human activity is particularly intense. Disturbed areas of all caves demonstrated broader diversity of known phyla as well as significantly higher abundance of sequences belonging to Bacteroidetes, which are known to be indicators of fecal pollution. The authors are thankful to the Lithuanian Science Council for the financial support (project No. MIP-005/2014).
IDENTIFICATIONS OF PROPER METHODS TO DETERMINE LEGIONELLA PNEUMOPHILA BACTERIA EXPOSED TO DIFFERENT ENVIRONMENTAL CONDITIONS
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Background

Legionella which is the agent of Legionnaires’ disease in natural and man-made water systems. Legionella bacteria pass to the viable but non-culturable (VBNC) phase due to the environmental conditions such as temperature, pH and disinfectants. After examining under laboratory conditions the water samples containing Legionella bacteria, have been exposed to environmental factors, it was shown that those bacteria couldn’t colony form medium, but this result does not mean that the bacteria are not present in a sample or an environment.

Objectives

The aim of the present study is to detect the most suitable methods for the recovery of L. pneumophila bacteria exposed to different environmental conditions.

Methods

In the current study, first of all, the detection limits of culture, fluorescent in situ hybridization (FISH) and semi-nested polymerase chain reaction (PCR) methods for different L. pneumophila concentrations (10²-10¹⁰ cell/L) was determined. Later, the recovery of L. pneumophila bacteria from the water samples including L. pneumophila bacteria that expose to different environmental conditions (temperature, pH and biocide) has been assessed by culture, FISH and semi-nested PCR methods.

Conclusions

In the present study, the detection limit of L. pneumophila bacteria was determined as 10³ cell /L for the culture method, 10⁵ cell/L for FISH method and 10⁶ cell /L for semi-nested PCR method. It was determined that FISH and semi-nested PCR methods are suitable methods for the recovery of L. pneumophila bacteria that are subject to different environmental conditions from the water samples.
ANTIBIOTIC RESISTANCE PATTERN OF PSEUDOMONAS SPP. ISOLATED FROM THE TOTAL COURSE OF THE RIVER DANUBE


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Background
This study was part of the Joint Danube Survey 3 (JDS3), the world's biggest river research expedition of its kind.

Objectives
The aim of the study was to determine the presence of acquired antibiotic resistance of different bacterial species with clinical impact.

Methods
During JDS3 180 water samples were taken over the total course of the river Danube. Samples were mixed with glycerol and stored at -70°C. For isolation five ml of the thawed samples were plated in 0.5 ml portions on Pseudomonas selective agar and incubated at 37°C. The isolates were tested for species identification with mass spectrometry MALDI-TOF MS (Shimadzu, Japan). Resistance testing was performed as recommended by the European Committee on Antimicrobial Susceptibility Testing (EUCAST). To estimate the presence of carbapenemase modified hodge tests were performed with all isolates resistant to at least one carbapenem.

Conclusions
Up to now 496 Pseudomonas were isolated, 343 (67%) Pseudomonas putida, 120 (25%) Pseudomonas fluorescens, all other species were represented by less than 10 isolates but no Pseudomonas aeruginosa was observed. Most common resistance was against meropenem (20.4%/76 isolates) piperacillin/tazobactam (4.3%/16 isolates) and ciprofloxacin (4.3%/16 isolates). None of the antibiotics were susceptible in all tested strains. Resistances were present all over the tested sampling points with no significant changes downstream. One isolate (Pseudomonas fluorescens) was tested positive for the presence of a carbapenemase in modified hodge test. Although the isolated Pseudomonas species are clinically less frequent, they are a
possible reservoir for resistance acquisition for other species (especially *Pseudomonas aeruginosa*).
SELECTION OF A SYNTHETIC SOIL MICROBIAL COMMUNITY FOR QUANTIFYING EFFECTS OF HYDRATION DYNAMICS ON COMMUNITY STRUCTURE AND DIVERSITY

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Background

Soil bacteria play a central role in many ecosystem services such as formation of rhizosphere communities, nutrient transformation and global biogeochemical cycles. Although molecular techniques have uncovered the tremendous bacterial diversity in soil, the mechanisms that control the assembly, functioning and maintenance of complex microbial communities remain largely unknown.

Objectives

The study aims at linking dynamics of soil hydration conditions and related biophysical factors with variations in microbial composition and ecological functioning. Understanding factors that shape the dynamics of species abundance is essential to get an insight into the assembly of a microbial community, its functioning and response to induced changes in water availability, a key parameter of soil environments.

Methods

To assist with definitive community level observations we designed a synthetic microbial community comprised of 10 well-characterized bacterial species spanning a wide range of soil phyla to be inoculated onto model porous surfaces mimicking soil habitats. The experimental system consists of sand layer placed on porous ceramic surface connected to nutrient reservoir. Fluctuations in hydration conditions are induced by changing the reference nutrient reservoir level to simulate wetting-drying cycles. The activity and composition of the bacterial community is studied using 16S fingerprinting and quantitative PCR.

Conclusions

Spatial self-organization of microbial community is highly dependent on the hydration level of their habitat. Subjecting the synthetic community to a range of controlled physico-chemical conditions allows us systematic evaluation of the role of environmental fluctuations on changes in community composition.
TRANSCRIPTOME ANALYSIS OF THE BALTIC SEA ISOLATES: SHEWANELLA BALTICA, PARACOCCUS SP. AND FLAVOBACTERIUM SP. SHOWS THEIR SPECIFIC RESPONSE TO OSMOTIC STRESS.

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Background

The Baltic Sea constitutes very specific environment. One of its characteristics are brackish water conditions that are caused by high water input from the surrounding lands and rivers. However, because of its characteristic geographical location, salinity levels change significantly in different parts of the Sea and can extend from 2, through 7 and even up to 20‰.

Objectives

The aim of our work is to assess which genes are essential for marine bacteria to adapt to the changes in salinity of their environment.

Methods

Three bacteria isolated from Baltic Sea were sequenced using Illumina Hiseq 2,000 platform and characterized as Shewanella baltica, Paracoccus sp. and Flavobacterium sp.. Then, they were cultured in a chemostat system in the minimal marine medium at various salinity conditions. During steady-state growth cells were harvested for RNA extraction and Illumina-based RNA-Seq reads were analyzed with DESeq2 software.

Conclusions

The transcriptome analysis of the three strains Shewanella baltica, Paracoccus sp. and Flavobacterium sp. at various salinity levels (bacteria cultured in 2, 7 and 20 %) resulted in identification of groups of genes responsible for bacterial survival in the
challenging environment of the Baltic Sea. Each strain is characterized by different response to osmolarity stress, underlying the importance of diverse categories of functional genes in this marine habitat.
FUNCTIONAL RESILIENCE OF MICROBIAL ECOSYSTEMS IN SOIL: HOW IMPORTANT IS A SPATIAL ANALYSIS?

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Background
Microbial life in soil is exposed to fluctuating environmental conditions influencing the performance of ecosystem services such as biodegradation.

Objectives
However, as this environment is typically very heterogeneous, spatial aspects can be expected to play a major role for the ability to recover from a stress event.

Methods
To determine key processes for functional resilience, scenarios with varying stress intensities were simulated within a microbial simulation model and the biodegradation rate in the recovery phase measured. Besides temporal monitoring, results were analyzed regarding spatial and mechanistic aspects. Parameters including microbial growth and dispersal rates were varied over a typical range to consider microorganisms with varying properties.

Conclusions
Results of the mechanistic and spatial view show that key factors for functional recovery depend on stress intensity and the location of the observed habitats. The limiting factors near unstressed areas are spatial processes (motility and substrate diffusion), with increasing distance microbial growth becomes more important. To confirm this, we repeated the simulations including a dispersal network representing fungi in soil. The system benefits from an increased spatial performance due to higher bacterial mobility.

With these simulations we show the importance of spatial aspects even at the mm-scale for recovering after a severe stress event in a highly heterogeneous environment such as soil. In consequence a spatial-mechanistic view is necessary for examining the functional resilience as the temporal view alone could not have led to these conclusions.

Further research should explore the importance of a spatial view for quantifying functional resilience also after complex stress regimes.
UNCOVERING STABILITY MECHANISMS IN MICROBIAL ECOSYSTEMS - COMBINING MICROCOSM EXPERIMENTS, COMPUTATIONAL MODELING AND ECOLOGICAL THEORY IN A MULTIDISCIPLINARY APPROACH

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Background
Although bacterial degraders in soil are commonly exposed to fluctuating environmental conditions, the functional performance of the biodegradation processes can often be maintained by resistance and resilience mechanisms.

Objectives
However, there is still a gap in the mechanistic understanding of key factors contributing to the stability of such an ecosystem service.

Methods
Therefore we developed an integrated approach combining microcosm experiments, simulation models and ecological theory to directly make use of the strengths of these disciplines. In a continuous interplay process, data, hypotheses, and central questions are exchanged between disciplines to initiate new experiments and models to ultimately identify buffer mechanisms and factors providing functional stability. We focus on drying and rewetting-cycles in soil ecosystems, which are a major abiotic driver for bacterial activity.

Conclusions
Functional recovery of the system was found to depend on different spatial processes in the computational model. In particular, bacterial motility is a prerequisite for biodegradation if either bacteria or substrate are heterogeneously distributed. Hence, laboratory experiments focussing on bacterial dispersal processes were conducted and confirmed this finding also for functional resistance. Obtained results will be incorporated into the model in the next step. Overall, the combination of computational modelling and laboratory experiments identified spatial processes as the main driving force for functional stability in the considered system, and has proved a powerful methodological approach.
BIOEGO: DEVELOPMENT OF A FUNGAL-BACTERIAL CONSORTIUM-BASED SOIL INOCULANT FOR PLANT GROWTH PROMOTION AND BIOCONTROL

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Background

There is an emerging need for environment-friendly soil inoculants with favorable effects on crop plants.

Objectives

We aimed to develop BioeGO, a fungal-bacterial consortium-based soil inoculant ensuring increased nitrogen fixation, phosphorous mobilization, stem degradation and humus production on the treated fields and providing protective effects against phytopathogenic fungi.

Methods

Nitrogen-fixing component was selected from bacteria based on growth capabilities in nitrogen-free medium. Phosphorous-mobilizing and stem-degrading component was selected from Trichoderma strains based on cellulase- and phosphatase-producing abilities. Humus-producing component was selected from bacteria based on peroxidase-producing abilities, while biocontrol component was selected from Trichoderma strains based on in vitro antagonism towards phytopathogenic fungi.

Conclusions

An Azotobacter vinelandii strain is the nitrogen-fixing component of the soil inoculant with the potential to provide excess nitrogen for crops. Phosphorous mobilization and stem degradation are ensured by a Trichoderma harzianum strain producing cellulose-degrading enzymes in the absence of stem residues, while this ability is increased 10-15 fold in the presence of grinded maize stem. It also produces large amounts of enzymes liberating organically bound phosphorous. A Streptomyces albus strain with excellent peroxidase-producing abilities was selected as the humus-producing component, while a Trichoderma asperellum isolate with outstanding antagonistic abilities towards Fusarium, Phoma, Alternaria, Botrytis and Rhizoctonia.
strains is the biocontrol component of BioeGO. Microbial components of the consortium do not have negative effects on each other, thereby all beneficial effects can occur simultaneously, complementing each other. This research was supported by grant TÁMOP-4.1.1.C-12/1/KONV-2012-0012. Csaba Vágvölgyi thanks the visiting professor program, Deanship of Scientific Research, King Saud University, Riyadh.
CHARACTERIZATION OF POTENTIAL AEROSOLIC IN ACTIVITY OF FUNGAL SPECIES

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Background

Biological aerosol particles are ubiquitous in the atmosphere. Recent findings identified a pool of these particles as potential ice nucleators (IN), which are capable of catalyzing ice formation at relatively warm subfreezing temperatures and thus impact cloud formation and precipitation. Some fungal species have been identified as ice nucleation active (INA) e.g., the ice nucleation capacity of Fusarium acuminatum was described already some decades ago but recently more species were discovered to posses IN activity (e.g., Mortierella alpina). For fungal IN activity, a protein or at least a proteinaceous compound is hypothesized to be responsible.

Objectives

The aim of the study is to analyze the ability to release cell-free IN into the atmosphere. Therefore, the IN machinery of fungal species is characterized by analytical and molecular biological methods as well as chamber experiments with the ultimate goal to clarify their potential impact on atmospheric cloud formation and water precipitation.

Methods

In this study, liquid chromatography and mass spectrometry is performed to identify proteinaceous compounds of fungal IN. A specific IN trap chamber (IN-TC) is designed to analyze the aerosolic transport capacity of fungal IN. For testing functional IN activity a customized droplet freezing assay is used.

Conclusions

Functional IN active fractions of fungal protein surface extractions can be identified by liquid chromatography and SDS-PAGE. Preliminary IN-TC results show a successful proof of principle suggesting the chamber is capable of collecting aerosolic IN generated of fungal washing water.
DEVELOPMENT OF HEAVY METAL ION BIOSENSORS USING SYNTHETIC GENETIC CIRCUITS

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Background

We previously determined the genome of *Bacillus oceanisediminis* 2691 that was isolated from marine sediment of the South Korean coast and found that many genes encoded heavy metal resistance and efflux systems in the genome. Genes encoding putative cadmium efflux pumps, arsenic efflux pumps, a chromate transporter, and lead-, cadmium-, zinc-, and mercury-transporting ATPases were found in the genome, which are apparently regulated by CadC homologous repressor-type transcription factors.

Objectives

We attempted to develop heavy metal ion biosensors using synthetic genetic circuits with *cadC* genes.

Methods

It was confirmed the transcriptional modulations of CadC-controlled genes in response to various heavy metals in *B. oceanisediminis* 2691 using real-time PCR. Six *cadC* promoter-operator-structural genes were transcriptionally fused with *egfp* gene to make recombinant *E. coli* cells. A variety of heavy metals were treated and specific fluorescence intensities were measured.

Conclusions

Taken together, the results showed that CadC proteins specifically respond to heavy metals and may play separate roles in heavy metal resistances, which have been evolved in the heavy metal abundant marine sediment milieu. In biotechnological aspects, CadC-controlled transcriptional modules could be used in the development of harmful heavy metal bio-sensor.
CHARACTERIZATION OF COD REMOVAL, ELECTRICITY GENERATION AND MICROBIAL COMMUNITY IN MICROBIAL FUEL CELLS TREATING MOLASSES WASTEWATER

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Background

Wastewater microbial fuel cells (MFCs) have advantages in treatment wastewater and electricity production simultaneously. Molasses wastewater (MW) is one of suitable feedstock for wastewater MFCs.

Objectives

In this study, electricity generation, COD removal, and microbial communities were investigated using 3-types of MFCs which were single-chamber MFCs without proton exchange membrane (PEM) and with PEM (AC-MFCs and ACM-MFCs, respectively) and two-chamber MFCs (H-MFCs).

Methods

Diluted MW (10,000 mg·L⁻¹ COD) was continuously fed into the MFCs. Reactor performances in the single-chamber MFCs were similar, indicating that PEM couldn’t enhance the reactor performances. COD removal in the single-chamber MFCs (90%) was higher than in the H-MFCs (50%). However, current density in the H-MFCs (80 mA·m⁻²) was 1.4 times of that in the single-chamber MFCs (57 mA·m⁻²). Power density in the H-MFCs (17 mW·m⁻²) was 2.2 times that in the single-chamber MFCs (7.7 mW·m⁻²). Microbial communities of the MFCs were examined using 16S rRNA-iron torrent sequencing. For bacterial community, Clostridium (7.6%) and Geobacter (4.2) were abundant in the AC-MFCs, and Clostridium (7.6%) and Desulfovibrio (5.1%) in the ACM-MFCs. Bacterial community in the H-MFCs was different with the single-chamber MFCs. Pseudomonas (12.9%) was dominated in the H-MFCs, followed by Clostridium (7.0%) and Geobacter (2.9%). For archaeal community, Methanothrix (96.4%) was remarkably dominated in the single-chamber MFCs. However, Methanobacterium (35.1%) and Methanosarcina (28.3%) were abundant in the H-MFCs.

Conclusions

These results suggest that chamber design is a key factor to influence on COD removal and electricity generation in the MFCs treating MW.
GENOMIC INSIGHTS INTO THE CATABOLIC VERSATILITY OF RHODOCOCCUS SP. STRAIN P52 TOWARDS ENVIRONMENTAL POLLUTANTS

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Background
Rhodococcus sp. strain p52, an isolated dioxin-degrader, can aerobically utilize a variety of aromatic compounds, such as dibenzofuran, dibenzo-p-dioxin, 2-chlorodibenzo-furan, 2,8-dichlorodibenzo-furan, dibenzothiophene, biphenyl, naphthalene, fluorene, phenanthrene, and anthracene. Degradation of dioxin by strain p52 involves two dioxygenases, DbfA and DfdA, encoded by genes on two plasmids. Additionally, strain p52 can use linear alkanes (tetradecane, tetracosane, and dotriacontane) and branched alkane (pristane) as sole carbon and energy sources.

Objectives
The objective of the present study is to gain insight into the underlying genetic information for the catabolic versatility, and biodegradation potential of strain p52.

Methods
Complete nucleotide sequence of the strain p52 genome was sequenced with the PacBio RS II system. The genome was annotated by BLAST against database, including KEGG, COG, SwissProt, TrEMBL and NR. The predicted genes involved in degradation of aromatic compounds and alkanes were tested for their transcriptional activity. And the function of the genes was confirmed by heterologous expression in recombinant hosts.

Conclusions
Co-presence of catabolic plasmids and multiple catabolic genes, as well as genes involved in chemotaxis, solvent tolerance, and transportation in the strain favors its catabolic ability and environmental adaptation. The present study provides insights into the bioremediation potential of strain p52.
BIODEGRADATION OF P-NITROPHENOL BY DIFFERENT MIXED CULTURES

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Background
One of the most important environmental pollutant among nitroaromatic compounds is p-nitrophenol (4-nitrophenol, PNP). It is used in the manufacture of a wide range of products and is the intermediate of some organophosphorus pesticide degradation (1). PNP has been classified as a priority pollutant by United States Environmental Protection Agency (EPA), and its concentration in natural waters is restricted to less than 10 ng/L (2).

Objectives
The aim of this study was to investigate the biodegradation of PNP by three different mixed cultures (TUS, AM and KOV) isolated from contaminated soil and sediment.

Methods
The samples were inoculated into minimal medium with increasing PNP concentrations. The biodegradation process was monitored by UV spectroscopy. The toxicity of PNP solution and PNP degradation intermediers to luminescent bacteria Aliivibrio fischeri was evaluated according the ISO 11348-3.

Conclusions
The mixed culture TUS showed the greatest biodegradation potential. Mixed microbial population was capable of degrading up to 100 mg/L of PNP as a sole carbon and nitrogen source. Toxicity was significantly reduced after the degradation process.

References:
CHARACTERIZATION OF A THERMO-STABLE BIOFLOCCULANT PRODUCED BY METHYLOBACTERIUM SP

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Background

Chemical and organic synthetic flocculants are, currently, predominantly employed in different industrial processes such as wastewater treatment and drinking water purification. The use of bioflocculants is desirable as most chemical and organic synthetic flocculants have been shown to possess deleterious effects to human health and, also, are recalcitrant to biodegradation thus, polluting the environment.

Objectives

This study assessed the bioflocculant production potential of Methylobacterium sp. Obi isolated from Tyume River in the Eastern Cape Province, South Africa.

Methods

Methylobacterium sp. Obi (accession number HQ537130) identified through 16S rDNA and sequence BLAST analysis was evaluated for bioflocculant production and the physicochemical properties optimal for the production of bioflocculant was determined using standard technique. The bioflocculant was characterized for novelty using FTIR and thermo-gravimetric analysis.

Conclusions

Culture conditions for optimal production of the bioflocculant included the following: glucose as carbon source; complex nitrogen source of urea, yeast extract and (NH₄)₂SO₄; inoculum size of 1% (v/v); initial pH 10; and 1% Ca²⁺. Maximum flocculating activity of 90% was attained after 36 hr of cultivation with the bioflocculant exhibiting thermal stability up to 100°C. Optimal culture conditions resulted in purified bioflocculant yield of 4.61 g/l. Optimal dose for kaolin suspension clarification 0.3 mg/ml in the presence of 1% Ca²⁺. Fourier transform infrared (FTIR) spectrum of the purified bioflocculant showed the presence of carboxyl, amino and hydroxyl groups known to be critical in the flocculation process. Thermo gravimetric analysis revealed that the produced bioflocculant was thermo stable.
Background

Over the years, diversity of beneficial rhizobacteria associated with agronomical/horticultural crops, especially the influence of geographical distribution (altitudinal variation) has been well documented. However, the influence of physical geography (eastern and western aspect of a hill farm), on the community structure of beneficial rhizobacterial population, irrespective of the altitudinal variation in unclear.

Objectives

The nutrient and health status of crops growing on two opposite geographical aspects of a hill farm could be influenced by presence or absence of beneficial rhizobacterial communities, with exceptional abilities of nutrient mobilization and pathogen antagonism.

Methods

In the present study we investigated rhizobacterial population associated with apple trees (four different apple cultivars) grown on hill farms at similar altitudes on two opposite geographical aspects. In the present study, a total of 259 rhizobacteria were isolated from the rhizosphere of apple trees, of which 50 were selected and further examined on the basis of their abilities of nutrient mobilization and pathogen antagonism. Beneficial rhizobacterial diversity analyzed using molecular analysis (16S rDNA-ARDRA) and supported by statistical tools revealed the presence of dominant siderophore producing bacterial populations (nutrient mobilizers and plant pathogen antagonists) on the eastern aspect of hill farm.

Conclusions

Results from microbial diversity study correlated with the survey analysis, for the presence of 40-60% of diseased apple trees and high (80-90%) non-survivability of re-planted apple plants on the western aspect of hill farm, as compared to 10-15%
diseased apple trees and low problem (15-20%) of survivability of re-planted apple plants on eastern aspect.
ENTEROCOCCUS FAECIUM REDUCED MALODOROUS COMPOUNDS IN SWINE MANURE SLURRY IN VITRO AND IN CONFINEMENT HOUSES

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Background
Reducing malodorous compounds are still swine producers’ problem.

Objectives
This study evaluated different bacteria for their ability to reduce malodorous compounds.

Methods
Previously isolated and identified bacteria with 99% similarities were used in this study. These were Enterococcus faecium(M1), E. faecalis(M2), Acetobacter tropicalis(M3) and Bacillus subtilis subsp. subtilis(M4). The bacteria were tested using in vitro technique (serum bottles and plastic boxes) and the best bacterium was selected and used as spray in the confinement pig houses.

Conclusions
In serum bottles technique, pH values were reduced (P ≤ 0.05) and hydrogen sulfide (H₂S) gas decreased (P ≤ 0.05 and 0.01) from 12 to 24h while opposite was observed in NH₄⁺-N concentration in the liquid phase. Total volatile fatty acid, histamine, methyamine, ethylamine, putrescine and cadaverine as well as total biogenic amine concentration was lowest (P ≤ 0.05, H₂S, total mercaptans) and liquid NH₄⁺-N were comparatively lower (P ≤ 0.05 and 0.01). M1 was selected and used for spray to confirm the odor reduction at different periods in the confinement pig houses. Odor compounds were lessened by 28.21%, 42.15% and 15.02% for NH₃, H₂S, and total amines, respectively up to 3 days after spraying. E. faecium can reduce malodorous compounds in swine manure slurry in vitro and in confinement houses.
COMPARISON OF DIFFERENT BACTERIA ON REDUCING MALODOROUS COMPOUNDS USING IN VITRO SWINE FERMENTATION

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Background
Bacteria play a significant role in odor production.

Objectives
This experiment was conducted to determine the appropriate bacteria for reducing malodorous compounds during fermentation in pig.

Methods
The bacteria were isolated and identified from rectum of pigs and were used in this study. They have 99-100% similarities to: Enterococcus faecium CWBI B1430(M1), E. faecium DSM 20477(M2), E. faecalis(M3), Lactobacillus acidophilus(M4), L. plantarum(M5), Bacillus subtilis ZHA10(M6), Bacillus subtilis XJASZB15(M7), Acetobacter malorum(M8), and E. faecium NRIC0114(M9). One hundred ml of fecal slurry was transferred to serum bottles containing 1g of soluble starch, inoculated with or without 1ml of different microbial cultures (1.0×10⁷CFU/ml) and incubated anaerobically for 12 and 24h.

Conclusions
Ammonia-nitrogen (NH₃-N) contents (gas phase) were decreased rapidly from 12 to 24h but was not found in M7 and M9 at 24h while NH₃-N contents (liquid phase) of M6 was significantly lowest after 24h. Hydrogen sulfide (H₂S) contents (gas phase) were detected in M5, M6, M7, M8 and M9. Total biogenic amine was found lowest (P Enterococcus faecium NRIC 0114(M9) is the best among the all of the treatments on the basis of toxic compound production.
GRAZING BY NEMATODES AFFECTS MAIZE RHIZOSPHERIC BACTERIA

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Background

Both mutualistic and pathogenic bacteria colonize the plant rhizosphere. Mutualists promote plant growth by providing nutrients or defense against pathogens. Nematodes and protozoa are also enriched in the rhizosphere, but little is known about the interactions of the grazing microorganisms with the bacterial community. The diversity of protozoans and nematodes in the corn rhizosphere has’t been studied and will provide a basis for the further analysis of their grazing preferences and their impact on bacteria.

Objectives

Identify and isolate protozoans and nematodes from the corn rhizosphere with metagenomic and culture-dependent approaches.

Methods

The diversity of the eukaryotic microorganisms was assessed analyzing 18S rRNA gene sequences from DNA extracted from corn rhizosphere samples. Nematodes were cultured in Nematode Growth Medium (NGM) and identified as described above. Feeding preferences were assayed in Petri dishes with NGM medium in the presence of different bacteria. The effect of nematodes was tested by adding the nematode to roots of maize plants growing in flasks with vermiculite.

Conclusions

The metagenomic analysis was performed with DNA samples from Huitzilac in the state of Morelos and in Amecameca in the state of Puebla. In cultures, the single nematode present was identified as Oscheius tipulae. This nematode has a preference for the bacterium Rhizobium phaseoli Ch24-10, and decreases the growth of corn plants in laboratory assays. However, in the presence of a group of selected root bacteria such as Rhizobium phaseoli, Methylobacterium extorquens and Chromobacterium violaceum, the growth of the corn plants was enhanced compared to control plants without the nematode.
VIBRIO VULNIFICUS EMPLOYS DIFFERENT DEFENCE MECHANISMS AGAINST PROTOZOAN GRAZING

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Background

Vibrio vulnificus is an opportunistic pathogen responsible for wound infections and septicaemia following ingestion of contaminated seafood, and has the highest reported mortality rate for seafood-related diseases. It is an autochthonous inhabitant of coastal marine environments where it is exposed to predation by heterotrophic protozoa.

Objectives

Protozoan predation on bacteria acts as a selective force, leading to evolution of antiprotozoal mechanisms that may also function as virulence factors. This study investigated the correlation of the place of isolation, genotype and virulence potential with grazing resistance of V. vulnificus.

Methods

Twenty two strains of V. vulnificus were exposed to grazing by Tetrahymena pyriformis, a filter feeding ciliate, and Acanthamoeba castellanii, a surface browsing amoeba in microtitre plates. V. vulnificus numbers in planktonic phase were determined by plate counts and photometry while biofilm biomass was assessed by crystal violet staining. The health and number of protozoa were determined by microscopy.

Conclusions

Although there was no significant difference in grazing resistance between C-genotype (clinical origin) and E-genotype (environmental origin), an environmental and a clinical isolates exhibited unique grazing defence mechanisms against the protozoa: the oyster isolate showed toxicity against T. pyriformis while the blood
isolate employed biofilm and pellicle formation to escape predation. Furthermore, the role of capsular polysaccharide (CPS) in grazing resistance of *V. vulnificus* was examined; no statistically significant differences were found between the encapsulated, "opaque" strains and the non-pathogenic non-encapsulated, "translucent" strains. These results show that the defence mechanisms of different strains of *V. vulnificus* have evolved along different evolutionary pathways.
MICROBABIAL COMMUNITY DYNAMISM DURING START-UP OF ANAEROBIC DIGESTER

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Background

The start-up of anaerobic digester has not been studied in detail from the microbiological point of view. Knowledge of microbiology process could help to determine whether an anaerobic digester has reached steady state, in the event that physicochemical data are not successful.

Objectives

The objective is to determine the dynamism of the microbial community during the start-up of an anaerobic digester.

Methods

The start-up of two-mesophilic anaerobic digester (fed with mix sludge) was studied through methane productivity, volatile suspended solids (biomass), along with other chemical variables. The inoculum was obtained from 'La Farfana”, wastewater treatment plant located in Santiago, Chile.

DNA was extracted from samples of sludge operation days 0, 13, 25 and 50. The genes studied were: 16S (Bacteria and Archaea) through DGGE and 4 functional genes (Hydrolytic, Homoacetogenic, Methanogenic and Acidogenic) through real time PCR.

Conclusions

The digester reached steady state after 30 days of operation, with productivity of 140 mL methane/ (Liter day) and 40% increase in biomass.
When the digesters reached the steady state the number of haplotypes (DGGE) increased about 40%, while the copies number of functional genes increased twice in
relation to the beginning of the process.

Figure 2 DGGE 16S of digester 1: (A) *Archaea* and (B) *Bacteria*
Fig 3. Copy number of hydrolytic gene during start up of two anaerobic digester
The digester 1 is represented for ● and digester 2 is represented for ○
MICROBIOLOGICAL SOIL QUALITY IN CITRUS ORCHARD UNDER DIFFERENT MANagements

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Background
The inter rows of citrus orchards in Brazil are generally managed with mowing, while in the lines mainly use the herbicide glyphosate. There is a great predominance of Urochloa decumbens in the inter rows of orchards. As there are reports of allelopathy of U. decumbens to citrus and damages that result from the continued use of glyphosate in other cultures.

Objectives
This project aims to study through microbiological parameters influences on soil quality using two vegetation in inter rows, managed with different mowing, with and without herbicide in the lines of Tahiti acid lime orchard.

Methods
Since 2010, trial was conducted in Mogi Mirim, Sao Paulo State, Brazil, in split plot design, where in the plots were composed of two types of vegetation: U. decumbens and U. ruziziensis; the sub plots for two types of mowing: conventional and ecological; and the sub sub plots: by applying and absence of the herbicide glyphosate on planting line. Samples of soil from the rhizosphere of plants were collected 0-10 cm depth, in three seasons and the following parameters evaluated microbial biomass carbon (BMC), basal respiration (RB) soil and metabolic quotient.

Conclusions
The results showed that there were significant differences for the microbiological parameters and soil treatment with U. decumbens, ecological mowing and herbicide use were better when compared with others. There was no damage to the vegetative and productivity development of Tahiti acid lime, so the changes in management can be recommended.
EPIBACTERIAL AND PLANKTONIC BACTERIAL COMMUNITIES RESPOND TO INCREASED PCO2 AND RISING TEMPERATURE LEVELS: A THREE-MONTH MESOCOSM STUDY

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Background

In the framework of the BMBF-funded consortium project BIOACID (Biological Impacts of Ocean Acidification) we are interested in the response of marine epibacterial communities attached to biological surfaces to increased pCO₂ and temperature. In spring 2014, we performed a three-month experiment in the benthic mesocosm facility at the Alfred Wegener Institute in List (Germany) which allows low and high tide simulation to mimic Wadden Sea conditions. We manipulated the seawater pCO₂ and temperature as predicted by the end of this century to simulate ocean acidification and warming events of the future North Sea.

Objectives

Both stress factors were tested for single and combined effects on epibacterial communities on the surface of the brown macroalgae Fucus vesiculosus forma mytili and in comparison on the planktonic microbiota of the surrounding seawater.

Methods

Changes in the bacterial phylogenetic diversity were analyzed by bacterial 16S rDNA (V1-V2) amplicon sequencing using the Illumina MiSeq technology.

Conclusions

The generated data indicated significant shifts in the bacterial community structure over a period of three months (April to June), driven by rising temperature and lowered seawater pH levels. Overall, our findings suggest significant changes in the microbial diversity under anthropogenic pressures such as global warming and ocean acidification with potentially severe consequences for the future Wadden Sea ecosystem.
REMOVAL OF LEAD BY PSEUDOMONAS SP ISOLATED FROM KHUZESTAN'S PETROLEUM SOILS, IRAN

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Background
Biosorption of heavy metals is effective technology for removal of toxic metals. Algae, bacteria and fungi are useful biosorbents for metal elimination from polluted zones.

Objectives
The objective of this study was isolation of lead resistant Pseudomonas strains.

Methods
Soil samples were homogenized and diluted with sterile normal saline. These samples were cultured in LB agar containing 5ppm of lead in order to isolation of resistant strains. These strains were isolated and cultured in Macconkey agar for isolation of appropriate strains and identified by metabolically tests. Maximum lead resistance of strains was evaluated by MIC test. The bacterial growth were optimized in the presence of lead in various temperatures, rate of shaking and values of pH by spectrophotometry in 600nm at 24h of cultivation. The bacterial strains were tested for ability to grow in the presence of different antibiotic by disc diffusion method. Metal adsorption test measured under optimal conditions by atomic absorption spectroscopy.

Conclusions
In this study 24 colonies were resistant to lead and the bacterium, Msap, had more resistant than others. The best conditions of growth in temperature, pH and rate of shaking were 37, 9 and 200. The results were verified using the general linear model. The highest of biosorption capacity was 91.79%. The strain, Msap, showed resistance to ampicillin, tetracycline, chloramphenicol, erythromycin, and kanamycin. Msap, have a great potential for bioremoval of lead from aquatic polluted environments. In conclusion, this study reveals the significance of using the Pseudomonas sp in bioremediation of lead contaminated zones.
CHARACTERIZATION OF BIOAEROSOLS - VALIDATION OF A REDUCED SAMPLING METHOD -
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\textbf{Background}

Bioaerosol monitoring is a rapidly increasing field of environmental risk assessment, which includes measurement of different microbiological and chemical parameters in both indoor and ambient air. Generally, bioaerosols are particles of biological origin with an aerodynamic diameter of up to 100 nm, containing bacteria, fungi, viruses, pollen, plant debris as well as their metabolites like endo-/mycotoxins, allergens and (M)VOC’s. In the last years, numerous studies showed associations between biological particles and a wide range of adverse health effects such as infections, intoxication and sensitization.

\textbf{Objective}

The aim of the study was the validation of a reduced and optimized sampling procedure necessary for a comprehensive bioaerosol risk assessment.

\textbf{Methods}

We investigated bioaerosols from laying hen-, swine- and cattle livestocks on different measuring sites (within the barn, 50 m downwind and background) using impaction and impingment. For the microbiological testing the total viable count of bacteria and fungi as well as a selective enrichment for MRSA, ESBL and VRE was carried out. Additional species identification (bacteria/ fungi) was performed by MALDI-TOF-MS. The chemical characterization included the detection of particulate matter (PM\textsubscript{10}, PM\textsubscript{2.5}, PM\textsubscript{1}, UFP), endotoxin, metal and VOC content.

\textbf{Conclusion}

The interspecies comparison show clear variation in the microbiological (bacterial/ fungal) and chemical components (PM, endotoxin and (M)VOC level) profile between the livestocks (indoor). Up to 50 m downwind no differences compared to the background samples of the corresponding livestock were detected. Due to the harmonized combination of selected chemical and microbiological indicator parameter, the reduced sampling program constitutes a constructive alternative for a comprehensive risk assessment.
ISOLATION AND IDENTIFICATION OF CYANIDE TOLERANT BACTERIA FROM WASTEWATER DISCHARGED FROM AN ELECTROPLATING PLANT IN CAPE TOWN, SOUTH AFRICA

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Background

The pollution of water sources is one of the major challenges municipalities face worldwide. The industrial wastewater discharged into the municipal sewers often contain toxic substances. Electroplating industries contribute to the pollution of water sources with cyanide, cyanide metal complexes and heavy metals used in their metal plating processes.

Objectives

The aim of this study was therefore to isolate and identify cyanide tolerant bacteria from wastewater discharged from an electroplating plant in the Western Cape, South Africa which was suspected of disposing of its potentially hazardous wastewater into the municipal sewers.

Methods

Bacteria were isolated from the biofilm and wastewater discharged from the electroplating plant. The pure colonies were streaked on nutrient agar supplemented with 400 mg/L cyanide to test for tolerance. To characterise and identify the cyanide tolerant isolates the following tests were performed; gram and endospore staining, biochemical tests, selective media, VITEK 2 Compact, Genomic DNA extraction, Polymerase chain reaction (16s rRNA gene amplification) and sequence analysis.

Conclusions

Two of the isolates were identified as *Pseudomonas* species and the other two as *Bacillus* species. The four identification methods used (biochemical testing, selective media, VITEK 2 and sequence analysis (16s rRNA gene) could confirm 100% identity up to the genus level even though a few discrepancies in identifying the species existed. Further studies are however needed to study other genes to differentiate the species. The isolates are further to be tested for cyanide degradation and as possible candidates for bioremediation of cyanide and metal cyanide complexes.
MYXOBACTERIA IN PEAT BOGS AND FEN – AN ASTONISHING DIVERSITY

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Background

Increasingly appearing resistances to antibiotics and the subsequent need for new bioactive substances is one of the fundamental challenges of infection research. Myxobacteria have a rich secondary metabolism, which places them among the best known natural product producers. To date more than 100 new metabolites and more than 500 derivatives were described from these soil dwelling, predatory microorganisms. In the past it turns out that in particular new families, genera and species of myxobacteria are reliable sources for new compounds.

Objectives

Promising sources for the isolation of new myxobacteria are uncommon habitats like, for example, peat bogs or fen, which are characterized by low pH values. This habitat has been neglected with regard to the isolation of myxobacteria in the past. In our study, the diversity of myxobacteria in moor-samples from the Harz-region was evaluated by standard cultivation methods as well as by cultivation independent 16S rRNA-clone bank analyses.

Methods

A total of 297 clones of myxobacterial origin were sequenced and compared to sequences of cultures, isolated from these samples, by phylogenetic analyses. It turned out that the majority of myxobacteria is only represented by clone sequences and could not be cultivated. Comparing these sequences to sequences of a public database (NCBI) revealed that most of these uncultivated myxobacteria are exclusively related to other uncultivated myxobacterial sequences from acidic soils.

Conclusions

This suggests that peat bogs and fen harbor a big diversity of new myxobacterial species, genera and even families which themselves harbor a great potential of urgently needed, new secondary metabolites.
EVOLUTIONARY HISTORY INFLUENCES THE SALINITY PREFERENCE OF BACTERIAL TAXA IN WETLAND SOILS

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Background
Salinity is a major driver of phylogenetic bacterial community composition across the globe and, given the importance of microbial community composition for ecosystem function, it is necessary to understand how salinity structures bacterial communities. If phylogenetic groups exhibit coherence in their salinity preferences, characterizing these preferences will enhance our ability to predict how changes in salinity will affect microbial communities and their associated functions.

Objectives
Consequently, we aimed to examine the response of wetland bacterial community composition and function to changes in salinity and to assess whether salinity preferences were phylogenetically conserved in bacteria.

Methods
To address these aims, we used a reciprocal transplant experiment of fresh- and saltwater wetlands soils.

Conclusions
The salinity of both the origin and host environments affected bacterial community structure (16S rRNA gene sequences, Illumina) and activity (e.g., CO₂ and CH₄ production). Phylotypes were categorized as exhibiting a preference for freshwater, saltwater, or having no salinity preference by comparing presence/absence patterns across treatments. When these preferences were related to phylogeny, a significant influence of evolutionary history was seen in all preference categories. This phylogenetic signature was corroborated by differences in the salinity preferences of high-level taxonomic groups. The majority of a-proteobacteria and g-proteobacteria phylotypes preferred saltwater, while the phylotypes of b-proteobacteria were more likely to prefer freshwater, and phylotypes within Verrucomicrobia tended to have no salinity preference. Overall, our results indicate that salinity structures bacterial communities by selecting for organisms with phylogenetically clustered salinity preferences, and these effects on community composition may have consequences for ecosystem function.
METABOLIC PATHWAY INVOLVED IN H2 PRODUCTION FROM FORMATE BY DESULFOVIBRIO VULGARIS

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Background
Formate is an important metabolite in anaerobic environments, where it is one of the metabolic products of fermentative organisms and an energy source for many bacteria such as sulfate-reducing bacteria (SRB). SRB are anaerobic organisms that possess a high number of formate-dehydrogenases (FDHs) and hydrogenases (Hases), enzymes responsible for H2 production from formate [1]. In sulfate-limited environments SRB grow fermentatively in syntrophy with other organisms producing H2. Moreover, it was recently demonstrated that in the absence of sulfate, Desulfovibrio vulgaris displays a very high H2 production from formate [2].

Objectives
In the present work, the electron transfer mechanism involved in formate-driven H2 production by SRB was studied.

Methods
H2 production by D. vulgaris deletion mutants of the two main FDHs (ΔfdhABC3 and ΔfdhAB) and of the four periplasmic Hases (Fe-only Hase (ΔhydAB), two [NiFe] Hase (ΔhynAB1 and ΔhynAB2), and a [NiFeSe] Hase (ΔhysAB)) was compared with the wild-type strain in order to disclose the role of each enzyme in the metabolic pathway involved in H2 production from formate [3].

Conclusions
This work contributes to our knowledge about the electron transfer pathways involved in H2 and formate metabolism in D. vulgaris, showing which of the FDHs and Hases play an important role in this process.


BACTERIAL PROFILING OF SAHARAN DUST DEPOSITION IN THE ATLANTIC OCEAN USING SEDIMENT TRAP MOORINGS – YEAR ONE RESULTS

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Background

Large quantities of dust are transported from the Sahara Desert across the Atlantic Ocean towards the Caribbean each year, with a large portion of it deposited in the ocean. This dust brings an array of minerals, nutrients and organic matter, both living and dead. This input potentially fertilizes phytoplankton growth, with resulting knock-on effects throughout the food chain. The input of terrestrial microbial life may also have an impact on the marine microbial community.

Objectives

The aim is to understand the links between dust input and the bacterial community and how this relates to ocean productivity and the carbon cycle.

Methods

The current multi-year project consists of a transect of floating dust collectors and sub-surface sediment traps placed at 12ºN across the Atlantic Ocean. Sediment traps are located 1200m and 3500m below the sea surface and all are synchronized to collect samples for a period of two weeks.

Conclusions

The first set of sediment trap samples were recovered using the RV Pelagia in November 2013 with promising results. Results from 7 sediment traps (three at 1200m and four at 3500m) were obtained. In general, the total mass flux decreased as distance from the source increased and the upper traps generally held more material than those at 3500m. Denaturing Gradient Gel Electrophoresis (DGGE) was used as a screening technique, revealing highly varied profiles, with the upper (1200m) traps generally showing more variation throughout the year. Preliminary analysis of 454 sequencing from a subset of the samples supports this observation, and detailed analysis will be presented.
Background

By the recent years, it has increased the air pollution in urban environments as well as rural, due to anthropogenic activities. This has been reflected on respiratory illness or infections related to air pollutants, mainly to fine inhalable particulate matter PM2.5; nonetheless, it is rarely discussed in the aburra valley about the microorganisms associated to this sort of particles, which can cause human illness ranging from mucosa irritation and allergies to asthma and pneumonia.

Objectives

The purpose of this research is the isolation and identification of microorganisms present in PM2.5 filters of the stations MED-UNFM and MED-PJIC of the Air Quality Monitoring Network from the Aburrá Valley.

Methods

Air sampling equipment BGI PQ200 semiautomatic was used for collection of biological material in PM2.5 Teflon filters 47mm in diameter. A sample of the latter was inoculated and incubated in an enrichment broth; Serial dilutions were made and were inoculated onto selective media. Each of the strains isolated, were characterized morphologically and biochemically (VITEK® 2 Compact).

Conclusions

In this research, several microbial species from PM2.5 filters, were isolated and identified like *Bacillus cereus*, *Acinetobacter baumannii*, *Enterococcus faecium* and *Bacillus anthracis*, all those consider as pathological microorganisms. This allows inferring that air quality, besides generating problems associated to respiratory illness, can increase the possibility of contracting bacterial infections.
Background
The Iron Quadrangle (Minas Gerais, Brazil) is one of the world’s largest mining regions, being historically explored for over 300 years. Since then, many toxic metals and metalloids were released into the environment leading to contamination of water bodies. It is well known that microbial community influences the bioavailability of these toxic elements and its importance in the ecological balance of various environments.

Objectives
This study investigated the taxonomic and functional diversity of the microbiome from Mina Stream sediment (MSS), historically metal-contaminated, especially arsenic.

Methods
The environmental DNA extracted was sequenced using SOLiD and Miseq platforms for the functional and taxonomic analysis, respectively.

Conclusions
Taxonomic profile obtained by Greengenes database revealed a complex microbial community, with dominance of Proteobacteria and Parvarcheota. Contigs recruited bacterial and archaeal genomes, especially Candidatus Nitrospira defluvii and Nitrosopumilus maritimus, and their presence implicated them in C and N cycling in the MSS. Functional reconstruction revealed a large diverse set of genes for ammonium assimilation and ammonification. These processes have been implicated in the maintenance of N cycle and healthy of the sediment. SEED subsystems functional annotation unveiled a high diversity of metal resistance genes, suggesting that the prokaryotic community is adapted to metal contamination. Furthermore, a high metabolic diversity was detected in MSS, suggesting that the historical arsenic contamination is no longer affecting it. Finally, the results reported herein may contribute to expand the current knowledge of the microbial taxonomic and functional composition of tropical metal-contaminated freshwater sediments.

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BIODEGRADATION OF AN ORGANOPHOSPHATE, CHLORPYRIFOS BY
MICROBES ISOLATED FROM AGRICULTURE SOIL
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Background
Organophosphorous Pesticides (OP) are being used for pest control in many
countries. Extensive use of these compounds results in their accumulation in
agricultural soil, water source, affecting insects and humans. Chlorpyriphos (CP) a
broad-spectrum insecticide has been used for controlling termites, beetles on paddy
fields and vegetable crops. CP acts as a neurotoxin by inhibiting acetyl-
cholinesterase, which degrades the neurotransmission agent, acetylcholine, in the
nervous system. Biodegradation of CP results in the formation of a compound TCP
(3,5,6-trichloro-2-pyridinol) classified as mobile, toxic also has antimicrobial property.

Objectives
Screening, isolation and identification of the microbial isolates for Chlorpyrifos and
TCP degradation

Methods
Resting cell assay for Chlorpyrifos degradation in liquid media and
HPLC analysis

Conclusions
Present investigation showed microbial strains isolated and screened from
agricultural soil amended with CP. Six bacterial strains (CPD-03, CPD-5C, CPD-7,
CPD-18, CPD-20 & CPD-33) showed significant degradation of CP in minimal salt
media supplemented with CP @ 10 µg/ml. A resting cell assay was carried out at pH
7.2, 37 °C with CP @ 10 µg/ml and monitored over a period of 48 h, and analyzed
using HPLC. The degradation efficiency of these isolates varied between 65% to 67%
within 48 h. The isolates CPD-18, CPD-20 and CPD-33 were able to degrade CP and
metabolize TCP up to 64% - 65 % within 48 hr compared to CPD-03, CPD-5C, CPD-
7 which degraded up to 65% - 67%. These six microbial strains can extend our
understanding of pesticide degradation and would be beneficial in using as a
consortium for cleaning up pesticide contaminated environment.
PREVALENCE AND ANTIBIogram PROFILING OF ESCHERIChIA COli PATHOTYPES RECOVERED FROM TYUME RIVER IN THE EASTERN CAPE PROVINCE OF SOUTH AFRICA
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Background

Waterborne diseases are among the leading causes of morbidity and mortality in developing countries and diarrheagenic *Escherichia coli* are one of the most important etiologic agents of acute diarrhea and represent a major public health problem in developing countries like South Africa.

Objectives

In this study, we assessed the prevalence and antibiogram profiles of *E. coli* pathotypes in Tyume River South Africa.

Methods

Twelve water samples were collected from three sites along the rivers between August 2010 and July 2011 and *E. coli* was isolated by the membrane filtration method on mFC and Chromocult agar. Identification and pathotyping of the isolates were done by polymerase chain reaction technique using species/pathotype specific primers and antibiogram was conducted using the disk agar diffusion method.

Conclusions

Of the 202 presumptive *E. coli* isolates recovered from the river 35% were confirmed as *E. coli*. The pathotypes prevalence was as follows: ETEC (30%); EPEC (35%); EAEC (35%); and EIEC (16%). All five *E. coli* pathotypes showed high-level resistance to ampicillin, tetracycline, cotrimoxazole, and chloramphenicol but were highly susceptible to quinolones, aminoglycosides, and novobiocin. All the EAEC, ETEC, and EIEC pathotypes were resistant against ampicillin, while 87.5% of the EAEC pathotype were resistant to carbenicillin. Also, all the pathotypes were susceptibility to quinolones. *TEM* genes and integron conserved segments were observed in 80% of the isolates. The Tyume river appears to be a reservoir of pathogenic *E. coli* strains and antibiotic resistance determinants, and hence a subject of public health importance.
PHYLOGENETIC CHARACTERIZATION AND COMMUNITY DIVERSITY OF HYDROCARBON-DEGRADING BACTERIAL POPULATIONS IN SOIL MICROCOSMS ENRICHED WITH VARIOUS AROMATIC HYDROCARBONS

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Background
High quantities of various aromatic hydrocarbons such as toluene, benzene, ethylbenzene and xylene (BTEX) are ubiquitously found in nature as a result of both natural and anthropogenic influences. Despite previous research efforts devoted to understanding the degradation of hydrocarbon pollutants in various environments, yet, there is still paucity of knowledge concerning the diversity of indigenous bacterial populations that are mostly responsible for the biodegradation of various hydrocarbons.

Objectives
The primary focus of this study was to isolate and characterize indigenous bacterial populations in soil microcosms previously enriched with various aromatic (BEX) hydrocarbons. Also, to compare the diversity of hydrocarbon degraders between the three microcosm communities examined.

Methods
Combinations of standard microbiological (phenotypic) and molecular (16S ribosomal RNA gene sequencing) were used to isolate, characterize and identify 45 bacterial isolates with putative hydrocarbon-degrading capability in the three soil microcosms. Additionally, various bioinformatics and analytical tools were employed for sequence alignments and diversity measurements.

Conclusions
A total of 45 bacterial isolates belonging to 5 distinct phyla were phylogenetically characterized among indigenous bacterial populations with putative hydrocarbon-degrading in the soil microcosms. In general, bacterial members of the γ-proteobacteria (mostly Pseudomonas and Acinetobacter) dominated (between 60 to 94%) among the isolates from the three microcosms. Bacterial members belonging to the β-proteobacteria (Comamonas and Delftia sp) and Firmicutes (Bacillus sp) were also represented. Overall, the differences observed in bacterial phylotypes among the microcosms are probably attributable to the direct effects of the chemical properties of each hydrocarbon pollutant on the indigenous microbial community.
NOSOCOMIAL INFECTIONS IN NIGERIA: A CASE STUDY FROM DELTA STATE, NIGERIA
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Background
Nosocomial or hospital acquired infections (HAIs) is a serious global public health issue, causing the suffering of about 1.4 million people across the world at any given time.

Objectives
This study was undertaken to ascertain the prevalence of HAI in Delta State of Nigeria.

Methods
This is a descriptive case study design in which structured questionnaire was administered to 96 doctors, 170 nurses, 24 pharmacists and 40 medical laboratory scientists from randomly selected government hospitals and 90 doctors, 180 nurses, 30 pharmacists and 30 medical laboratory scientists from randomly selected private hospitals, all located in the 25 Local Government Areas of Delta State. The study was carried out between May and July 2013 after Ethical Approval was granted by the Ministry of Health, Delta State. The commonest HAIs encountered are Urinary Tract Infection (61.4%) followed by Hospital acquired Pneumonia (55.6%) and Staphylococcus aureus (54.4%) for the government hospitals. In the private sector, Hospital acquired Pneumonia (66.1%) is the commonest. The study also showed that the frequency of occurrence of HAI is higher in government than private hospitals. This can be attributed to the higher population of patients leading to overcrowding. The modes of transmission of hospital acquired pneumonia was observed to follow this order airborne > contact with blood and body fluids > contaminated instruments > contaminated hands > needle sticks.

Conclusions
The prevalence of nosocomial infection in this state is very high. Efforts are needed to reduce it, improve the quality of life of patients and healthcare workers and improve healthcare delivery.
BACTERIAL BIODIVERSITY IN THE RICE RHIZOSPHERE AND HIGH-SALT SOIL AND SEDIMENT ENVIRONMENTS FROM THE CAMARGUE REGION (FRANCE)

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Background

The Camargue territory in the south of France is a delta triangle of 1500 km² of the Rhône River as it flows into the Mediterranean. This zone consists mostly of salty land inhabited by a diverse community of birds, wild white horses, black bulls and waterfowl, the most famous of which are pink flamingos. Rice production covers approximately 10,000 hectares which accounts for 30 percent of France’s rice consumption.

Objectives

We are examining the microbial composition of high salinity salt ponds, soils and the rhizosphere of rice plants in the Camargue region for two consecutive years (2013-2014) in order to establish a variable-temporal comparison regarding the physical and chemical properties of the environment versus the bacterial communities present in similar (saline) environments.

Methods

Bacterial biodiversity of five rhizosphere sites, growing two varieties of rice (Arelate and Gageron), plus sixteen sites of sediments and soils from nearly saline wetlands, were studied using pyrosequencing of PCR-amplified 16S V3-V4 rDNA amplicons from total extracted genomic DNA.

Conclusions

Our preliminary results revealed bacterial populations dominated by members of the Bacteroidetes and Proteobacteria phyla. However, in sandy soils with a lower degree of salinity than the 25 ppm, we observed, members from the Acidobacteria phylum representing an important fraction. Ongoing analyses of diversity (Chao1, Shannon and Simpson index) and statistics such as principal component analyses (PCA) will permit a detailed description of the bacterial populations among the different sampling sites, and better comparisons with similar types of sites worldwide.
INVESTIGATION OF AEROBIC HETEROOTROPHIC BACTERIA IN OIL RESERVOIRS IN THE SOUTHEAST ANATOLIA REGION

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Background
Petroliferous soil has unique environment due to the presence of a variety of aliphatic and aromatic hydrocarbons, creating a distinctive habitat for certain microbial communities. Although hydrocarbons are constituted only of two elements, relative to their electron acceptors; they are exploited as growth substrates by numerous microorganisms. Some aerobic bacteria, especially *Pseudomonads* are one of the most frequently isolated bacteria from hydrocarbon-impacted environments due to their reported metabolic pathways.

Objectives
The aim of this study was to determine the presence, abundance and the predominant culturable aerobic heterotrophic bacteria as well as *Pseudomonads* in 4 different production wells located in the Southeast Anatolia Region.

Methods

Petroleum samples were warmed to 40-60°C for a few hours until water-phase could be observed under oil phase. Diluted water-phase samples were inoculated onto Nutrient Agar, Tryptone Soy Agar, R2A, Cetrimid Agar and *Pseudomonas* Agar Base with and without salt. Petri plates were incubated at 30°C for 2 days (R2A for 5 days). Genomic DNA was extracted from the pure cultures using UltraClean Microbial DNA Isolation Kit. 16S rDNA gene fragments were amplified using U968F and L1401R primers. The amplified gene fragments were subsequently analyzed using sequencing.

Conclusions
It was determined that the number of aerobic heterotrophic bacteria and *Pseudomonads* was generally low/high (min. 9 cells/ml and max. 1358 cells/ml) in the oil reservoirs. 15 pure cultures were obtained from the samples. Among the cultured bacteria, *Acinetobacter guillouiae* (99.3%), *Halomonas daqingensis* (99.7%), *Suttonella indologenes* (93.1%) and *Pseudomonas stutzeri* (99.8%) were present.
EVALUATING THE RELATIVE INFLUENCE OF BIOTIC AND ABIOTIC FACTORS ON EFFICIENCY OF BIOAUGMENTATION MEDIATED EX-SITU BIOREMEDIATION

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Background

Soil and ground water pollution with xenobiotic chemicals is a major environmental concern. Bioremediation has been regarded as an efficient and environment friendly approach for restoration of contaminated niches. Amongst different bioremediation procedures, bioaugmentation is considered as one of the most potent approach for bioremediation. However, till date very few reports have shown successful application of this approach. It is often suggested that variety of biotic and abiotic factors e.g. indigenous microbial community, physico-chemical nature of contaminating pollutant and target soil may influence the efficiency and outcome of bioaugmentation process.

Objectives

To systematically address this issue, we performed a study for determination of relative influence of biotic and abiotic factors on efficiency of p-nitrophenol (PNP) degradation during bioaugmentation mediated ex situ bioremediation.

Methods

PNP degradation was monitored in microcosms prepared with soil samples collected from seven geographically distinct Indian provinces (viz., Assam, Andhra Pradesh, Gujarat, Karnataka, Maharashtra, Rajasthan and Tamil Nadu) having diverse physico-chemical character and native microbial communities. Soil samples were spiked with 70 ppm of PNP and subjected to bioaugmentation using pre-grown cells of degradative strain.

Conclusions

Quantification of residual PNP showed apparent enhancement of pollutant degradation; however, kinetics of pollutant removal varied significantly as a function of soils properties. Results from statistical analysis of PNP degradation patterns and matrices generated with microbial community and physico-chemical characteristic of soil highlighted relatively strong influence of indigenous bacterial community on biodegradation efficiency. It was observed that soil microbial community renders such impact by differentially modulating the survival of bioaugmented strain.
AN O-NITROBENZOATE (ONB) SENSITIVE TRANSPONsoon MUTANT OF ARTHROBACTER PROTOPHORMIAE STRAIN RKJ100 INDICATES FOR INVOLVEMENT OF N-ACETYGLUCOSAMINIDASE IN BACTERIAL RESISTANCE TOWARDS ITS ELEVATED CONCENTRATIONS

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Background

Microorganisms capable of thriving under extreme environments are of great significance for basic as well as applied studies. Consequently, microorganisms showing survival under extreme environments have attracted considerable interest. Several studies have shown isolation and characterization of extremophilic microorganism. However, relatively fewer studies have reported isolation and characterization of microorganisms from niches characterized by elevated concentration of anthropogenic, xenobiotic environment pollutants. Previously we had characterized Arthrobacter protophormiae RKJ100 for its ability to tolerate extremely high concentrations of o-nitrobenzoate (ONB), a toxic xenobiotic environmental pollutant.

Objectives

Main objective of present study was to characterization of gene(s)/ gene cluster(s) involved in resistance of strain RKJ100 towards elevated concentrations of ONB.

Methods

Random transposon mutants of strain RKJ100 were generated and screened for ONB sensitive phenotype. A sensitive mutant was defined as one which exhibits sensitivity towards ONB at ≥30 mM. Molecular and biochemical characterization of this mutant was carried out to identify the gene disrupted due to the transposition.

Conclusions

Results from the characterization of ONB sensitive mutants showed disruption of endo-β-N-acetylglucosaminidase (ENGase) gene. ENGase is a non-essential enzyme involved in oligosaccharide processing and cell wall recycling many life forms. It has been shown to have role in cellular homeostasis. Results obtained
during this study present first evidence for its role in bacterial resistance towards elevated concentrations of toxic xenobiotic compound.
BIODEGRADATION OF BISPHENOL A-LIKE ENDOCRINE DISRUPTORS BY DIVERSE MICROORGANISMS FROM RIVER SEDIMENT

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Background
Bisphenol A (BPA) is a widely used chemical in the production of plastics and resins. Tetrabromo-BPA (TBBPA), the BPA derivatives with bromide atoms, is one of the most commonly used brominated flame retardants. The broad distribution of BPA-like pollutants in the environment raises great concern because of their endocrine-disrupting characteristics and toxicity.

Objectives
Developing cheap and low environmental impact remediation methods for BPA-like pollutants is emergent. Therefore, the biodegradation abilities of diverse microorganisms toward BPA and TBBPA were investigated.

Methods
The microcosm from acclimated sediment culture was used to evaluate the biodegradation ability of diverse microorganisms toward BPA and TBBPA were investigated.

Conclusions
The microcosm completely removed BPA (10 mg L⁻¹) within 28 h and transformed the contaminant into several metabolic intermediates. During the degradation process, the microbial composition fluctuated and finally restricted to Pseudomonas knackmussii and Methylomonas clara. The four isolated BPA degraders were all genetically similar to P. knackmussii. Although the degradation ability of mixed strains was higher than that of single strain, it was far less than that of the original microcosm. These results demonstrated the novel role of P. knackmussii in BPA degradation, as well as the crucial role for microbial diversity in pollutant decomposition. From the acclimated sediment culture, a novel TBBPA degrader was isolated. It degraded 70 % of the pollutant (4 mg L⁻¹) within 5 days and transformed the pollutant to one degrading by-product. Our results are helpful to understand the natural attenuation processes of BPA-like endocrine disruptors, as well as to remediate those structural similar pollutants.
PRESENCE OF P. SHIGELLOIDES IN TEMPERATE CLIMATE ZONE LAKES

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Background

Plesiomonas shigelloides is a Gram-negative rod-shaped bacterium found in aquatic environments. Most of the reports on its isolation are from tropical or subtropical areas. Furthermore, the bacterium is considered as an emerging opportunistic pathogen.

Objectives

Aim of this study was to detect presence of P. shigelloides in lakes situated in temperate climate zone (Central Europe).

Methods

Total of 42 water samples taken from Vojvodina, i.e. northern Serbia and southern Hungary lakes were examined. Some samples were taken from same locations but at different time intervals, so all isolates were obtained from geographically and/or chronologically different sources. The samples were spread directly onto Inositol Brilliant Green Bile Agar and incubated at 37 °C for 24 hours. The colonies of inositol fermenting bacteria were randomly selected, subcultivated and several preliminary tests were performed (Gram staining, catalase and oxidase test, indol production, gelatin hydrolysis and citrate utilization test). P. shigelloides was additionally characterized and identification was confirmed by PCR using specific primers (Ref, god). Additionally, RAPD analysis was performed to estimate genetic relationships among the isolates using 5 different primers.

Conclusions

The results indicate the frequent presence of bacteria in the waters in moderate climate, since P. shigelloides was isolated from 81% of the samples. RAPD analysis revealed distinct DNA fingerprint patterns for each P. shigelloides strain. The results confirm high presence of P. shigelloides in temperate climate zone lakes in Central Europe, which is probably a result of global climate changes. In addition, P. shigelloides can represent a potential health concern in this region.
Background

Factory sites, wastewater treatment plants or hospitals are representative environments containing high concentrations of xenobiotics. β-lactam antibiotics (BLA) in these environments could directly exert a selective pressure on bacteria, resulting in spontaneous mutations or horizontal transfer of genetic elements among them. These genetic changes could pose a threat to human health, if these bacteria become multiresistant pathogens and migrate to the clinical settings. Our knowledge about the taxonomic composition of bacterial consortia from BLA contaminated environments is scarce.

Objectives

To investigate and characterize bacterial community subjected to long-term selection pressure of BLA, soil samples from the factory site of a pharmaceutical company, which has been producing β-lactam antibiotics for more than 50 years, were taken, analyzed, and taxonomic compositions were compared to those obtained from non-polluted soils of the same area.

Methods

In this study we assessed the microbial diversity on three different sites among the factory campus and two control sites (approx. 5km from the factory) using tag-encoded 454pyrosequencing of the 16s rRNA genes.

Conclusions

The selection pressure of BLA did not reduce the diversities of bacterial consortia compared to those in the reference samples, although composition of the dominant species was different. The data from culture dependent study showed that the overall resistome of the polluted soil communities was boosted not only in terms of
resistance to BLAs but also within the tetracycline, macrolide, chloramphenicol, or aminocyclitol class of antibiotics.

The data demonstrate a very strong selective pressure imposed by anthropogenic activities on the transfer of resistance determinants among bacteria.
PLACKETT-BURMAN STATISTICAL DESIGN TO IMPROVE THE CULTURE MEDIA COMPOSITION AND THE ENZYME ACTIVITY OF TWO RECOMBINANT LACCASES IN PICHIA PASTORIS

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Background
Laccases are multicopper oxidases widely distributed in nature and catalyze the transformation of aromatic and non-aromatic compounds with reduction of molecular oxygen to water. We cloned the syntetic optimized genes, *GilCC1* and *POXA* 1B from *Ganoderma lucidum* and *Pleurotus ostreatus* respectively in *Pichia pastoris*. In expression experiments we obtained enzymatic activities of 451.08±6.46 UL⁻¹ and 0.13±0.028 UL⁻¹ for *GilCC1* and *POXA* 1B respectively. We are planing to employ recombinant laccases for the degradation process of kraft pulping or dye removal from effluents.

Objectives
Objective was to improved the enzymatic activity of each cloned genes by improving the culture media composition.

Methods
We use a Plackett-Burman statistical design to evaluate media volume, inocula percentage, copper, glucose, NH$_4$SO$_4$, peptone and yeast extract concentration, each one of them with two levels, to detect the positive or negative influence and contribution percentage of each one.

**Conclusions**

In both cases we improved the enzymatic activity by optimizing culture media composition. In the case of *GLCC1*; treatments 1 and 9 surpassed 1.5 UL$^{-1}$; which exceeds the maximum activity obtained in previous trials 0.13 UL$^{-1}$; being treatment 1 which achieved higher laccase activity 4.69 UL$^{-1}$ meaning an increase of 36.07 times. In the case of *POXA1B*; treatments 4 and 11 surpassed 1200 UL$^{-1}$; exceeding the maximum activity obtained in previous trials 451.08 UL$^{-1}$; being treatment 11 which achieved higher laccase activity 1373.72 UL$^{-1}$ meaning a 3-fold increase enzyme activity. For both laccases genes the culture time of *P. Pastoris* was 168 hours.
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Environmental microbiology - 2

SEQUENTIAL STATISTICAL DESIGNS FOR IMPROVEMENT A RECOMBINANT LACCASE PRODUCTION IN PICHIA PASTORIS IMMOBILIZED CELLS
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Background

We cloned the syntetic optimized gene, POXA 1B from Pleurotus ostreatus in Pichia pastoris. In expression experiments we obtained an enzymatic activities of 451.08 ± 6.46 UL⁻¹. We are planning to employ a packed bed column, with the recombinant immobilized cells for dye removal from effluents.

Objectives

Objective was to improved the enzymatic activity of recombinant POXA 1B by improving the culture media composition in a bath culture of immobilized cells of the recombinant strain.

Methods

We use a Plackett-Burman statistical design to evaluate media volume, copper,
glucose, NH_4SO_4, peptone and yeast extract concentration, each one of them with two levels, to detect the positive or negative influence and contribution percentage of each one. After that, a Box-Behenken design allow us to optimize the more influent factor by analyzing the three levels of factor interaction through a response surface.

**Conclusions**

We improved the enzymatic activity from 14.4U/L at 156h (first immobilized cell assay) to 1300U/L at 168 h of immobilized cells bath culture (after statistical) optimization, meaning a 90.3-fold increase enzyme activity at 168 hours. The recombinant activity of the recombinant enzyme POXA 1B produced by using immobilized cells; exceed the maximum activity obtained in previous free cells trials 451.08 UL^{-1}; meaning a 2.88-fold increase enzyme activity at 168 hours.
MICROBIOLOGICAL STUDY OF SEWAGE SLUDGE COMPOSTING USING MICRO-POURUS LAMINATE MEMBRANE COVER SYSTEM TECHNOLOGY

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Background
Treatment of wastewater involves generation of large amounts of solid wastes worldwide, known as sewage sludge. Composting of sewage sludge from wastewater treatment plants (WWTPs) is considered a successful strategy for its sustainable recycling.

Objectives
The aim of this study was to investigate the bacterial community structure and population dynamics during composting, using both cultivation-dependent and cultivation-independent techniques, to gain knowledge contributing to the improvement of the efficiency of the process.

Methods
The composting process was conducted during 180 days in an industrial composting plant. Samples were retrieved from a full-scale pile with semi-permeable Gore-tex covers, in combination with an air insufflation system. Waste sludge from an urban WWTP was mixed with vegetal pruning wastes as bulking agent. Microbial characterization included: enumeration of viable (platables) cells, measurements of enzyme activities, phylogenetic identification of isolates by 16S-RNA gene sequencing, and characterization of bacterial communities by a next-generation, Illumina-based sequencing approach.

Conclusions
Composting of sewage sludge under microporous-laminate membrane covers enhanced the degradation process, compared to other composting technologies. Changes in enumeration of cultivable microbiota took place during the 15 first days of composting. High activities of enzymes dehydrogenase, protease and arylsulphatase were observed during the first five days, suggesting a high rate of substrate degradation. Pyrosequencing data demonstrated that bacterial diversity was strongly influenced by the stage of the process. The greatest bacterial diversity was observed between days 15 and 30. These results suggest that composting under microporous-membrane covers can improve the process performance and reduce the operating cost.
COMMUNITY COMPOSITION AND DIVERSITY OF N FIXING CYANOBACTERIA ASSOCIATED WITH MOSSES IN SUB-ARCTIC ECOSYSTEMS

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Background

Moss associated cyanobacterial communities (MAC) are thought to be the major drivers of the Nitrogen (N) budget at high latitudes. Most studies have been carried out in the boreal forest and the High Arctic, whereas biological N fixation in other moss-rich regions e.g. sub-arctic alpine ecosystems may also be largely MAC-based.

Objectives

To evaluate diversity, abundance and N fixation activity of cyanobacteria associated with the four moss species Racomitrium lanuginosum, Hylocomium splendens, Pleurozium schreberi and Sanionia uncinata, all abundant in Icelandic ecosystems.

Methods

Moss samples were collected from three moss-dominated ecosystems in Iceland. Cyanobacterial identification and quantification was carried out by phase-contrast, fluorescence and confocal scanning microscopy. Estimation of cyanobacteria relative abundance was performed by amplification and sequencing of nifH genes. N fixation was assessed with the acetylene reduction assay (ARA).

Conclusions

The cyanobacterial strains identified appeared to be from the orders Stigonematales and Nostocales. N fixation varied over time, also it was responsive to microclimatic/micro-topographic gradients. Simulated climate warming and grazing negatively affected N fixation activity. Moss water content and type of vegetation were the most influential parameters on potential N2 fixation activity. Our finding about MAC in moss-widespread sub-arctic ecosystems may have substantial impact on the understanding of the N cycle in this terrestrial environment.
CHARACTERIZATION OF BACTERIAL COMMUNITY ASSOCIATED WITH THE DEGRADATION OF PIG CARCASSES

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Background
Foot-and-mouth disease virus causes a highly contagious vesicular disease of domesticated and wild ruminants and pigs. A significant volume of infected livestock carcasses should be disposed of safely, economically and environmental-friendly.

Objectives
Composting is one of strategies for the safe disposal of infected livestock carcasses. In this study, the bacterial community associated with the degradation of pig carcasses was characterized in the composting process of them.

Methods
The compost bin was constructed with concrete frames for the walls and floor. Two-layer stainless steel grids were installed to put the carcasses of 4 pigs (80~90 kg/individual) in the bin. The bin was filled with well-mature compost mixture. To collect compost sample, perforated stainless steel tubes, which have been specially designed to collect sample without disturbance, were installed in the bin. Bacterial community was analyzed using 16S rDNA-PCR and pyrosequencing.

Conclusions
The results of volatile organic compounds (VOCs) monitoring indicated that the easier degradable fractions of pig carcasses decomposed during initial 50 days, and most of pig carcasses decayed during 170 days. The VOC concentrations at the top layer of compost bin were lower than VOCs at the bottom, indicating the top layer of compost can play a role as biofilter for the removal of malodor gases. Dominant bacteria at the initial stage were Alcanivorax, Aquimarina, Atopostipes, Bacillus, Cerasibacillus, Pseudomonas, Psychrobacter, Terribacillus and Ulvibacter. At the active degrading period of pig carcasses, Alcanivorax, Bacillus, Clostridium, Ignatzschineria, Lentibacillus, Pelotomaculum, Sporanaerobacter and Tissierella were dominated. Caldicoprobacter, Clostridium, Sedimentibacter, Syntrophaceticus and Virgibacillus were followed during final composting period.
POTENTIAL USE OF YEASTS ISOLATED FROM FORESTED WETLAND IN AZO DYES DECOLORIZATION

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Background

Forested wetlands are interfaces between terrestrial and freshwater systems and play an important role in the hydrology and nutrient filtration, acting as carbon sinks and in biodiversity regulation and maintenance. Also, despite microbial biodiversity in these systems is high, the biotechnological potential of this microbial diversity has been almost ignored, particularly yeast communities.

Objectives

The main purpose of this study is to evaluate the potential exhibited by yeast isolates in decolorizing different azo dyes, for future use in bioremediation.

Methods

In this work, wetlands have been screened for wild yeast isolates exhibiting ability to degrade five different azo dyes, a major group of synthetic dyes highly recalcitrant to current degradation processes. From the universe of isolates selected based on different colonial morphologies, 92 yeast isolates were randomly chosen for assessing their decolorizing abilities. Hierarchical clustering, based on medium color changes during incubations, were used to evaluate differences in the performance of azo dyes decolorization.

Conclusions

Yeast isolates differed markedly in their ability to decolorize the tested dyes. Among the best eleven isolates selected and identified to the species level by molecular methods, the basidiomycetous yeasts Cryptococcus laurentii AGG729 and Cryptococcus podzolicus AGG691, showed the highest dyes removal capacity. Our
results pointed out that yeasts occurring in forested wetlands, have potential to be used in biotechnological processes, being azo dyes bioremediation a case in point.

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CHARACTERIZATION OF MICROBIAL COMMUNITIES WITHIN PETROLEUM REFINERY WASTES AND THEIR CATABOLIC POTENTIAL IN BIOREMEDIATION

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Background
Autochthonous microbial communities within the highly hazardous petroleum contaminated sites could play significant role in in situ bioremediation. Unlike oil spill and other hydrocarbon contaminated environments, microbial diversity and their remedial potential is least studied within petroleum refinery wastes.

Objectives
During the present study microbial diversity and their catabolic potential relevant for in situ bioremediation of petroleum refinery waste were studied.

Methods
Gas chromatography mass spectrometry, ICP, and other standard methods were used to characterize the waste sludge. Orion multiparameter meter was used to determine DO, pH, ORP, temperature on site. 16S rRNA gene sequences were obtained from metagenomes through Illumina (GAIIx) sequencer. Clone library based analysis was done to reconfirm the archaeabacterial populations. 65 bacterial strains were isolated through plating on R2A medium. BIOLOG eco plates were used to ascertain community level physiological profiles.

Conclusions
Dominance of anaerobic, moderately thermophilic, fermentative, sulphate-, iron-, nitrate- reducing, bacterial genera like Fervidobacterium, Coprothermobacter, Thermodesulfovibrio, Deferrribacter, Nitrospira, Anaerolinea, Thermoanaerobacterium, etc. was observed. Presence of methanotrophic and methanogenic, acetoclastic, hydrogenotrophic, archaea (Methanoseata, Methanolinea, Methanocella, etc.) was confirmed by both NGS and clone library analysis. Bacterial strains isolated from the samples showed their affiliation to genera Pseudomonas, Bacillus, Burkholderia, Enterobacter, Kocuria, Microvirgula, Pandoreae etc. The isolates were screened for their ability to grow in presence of various hydrocarbons (Naphthalene, Dodecane, etc.) crude oil, oily sludge, biosurfactant production, varied temperature (15⁰C - 45⁰C), pH (3.0 - 9.0) and heavy metals. The overall study illustrated microbial community structure and their potential in survival and hydrocarbon biotransformation, which may be exploited for in situ bioremediation strategies.
CHARACTERIZATION OF THE FUNGAL IN MACHINERY BY MEANS OF FUSARIUM ACUMINATUM

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Background
Freezing of water above -36°C is based on ice nucleation activity (INA) mediated by ice nucleators (IN), and biological particles are a potentially important source of IN in the atmosphere. INA of the fungal genus Fusarium was already described about 30 years ago and INA of Fusarium as well as INA of other fungal genera is assumed to be mediated by proteins or at least to contain a proteinaceous compound. However, to date the precise INA machinery of Fusarium remains unidentified.

Objectives
In this study we use analytical technics to identify INA protein in surface protein extract from Fusarium acuminatum. The ultimate Goal is to elucidate the biological and environmental impact of fungal IN.

Methods
We use liquid chromatography, customized freezing assay, SDS-page, mass spectrometry and molecular methods to subsequently analyze the fungal IN machinery.

Conclusions
Preliminary results show that the INA protein of F. acuminatum is contained in the early size exclusion chromatography fractions indicating a high molecular size. Moreover we could identify a single protein band from IN active fractions at 130-145 kDa corresponding to sizes of IN proteins from bacterial species. To our knowledge this is for the first time an isolation of a single protein from in vivo samples, which can be assigned as IN active from Fusarium.
POPULATION ANALYSIS OF BACTERIA IN A LOW TEMPERATURE WATER CATCHMENT SYSTEM

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Background

Drinking water supplies are based on groundwater resources all over the world. At some locations problems with higher concentrations of iron and manganese in groundwater can occur. The principle of subsurface iron and manganese removal are periodically injections of aerated water into an anoxic aquifer through a well, resulting in the precipitation of iron and manganese hydroxides. However, the precipitation of insoluble oxyhydroxides caused by chemical and microbial processes may result in the clogging of the aquifer and pumps.

Objectives

Bacterial communities of water wells from a water catchment system with subsurface iron and manganese removal were determined to identify key bacteria. Phylogenetic diversity and relative abundance of bacteria in water samples were compared between wells already in operation to those not yet operated.

Methods

To estimate well-clogging, qPCR with universal primers was performed for all samples and specific primers targeting (i) Rhodoferax spp., (ii) Crenothrix polyspora and (iii) Gallionella spp. were applied. Furthermore, PCR-DGGE technique and 454-pyrosequencing were used to observe bacterial alterations within water samples.

Conclusions

Our results showed that wells in operation contained a higher alpha-diversity, and bacterial communities were dominated by Rhodoferax and Methylotenera. Abundant groups in not yet operated wells were Gallionella spp. and Crenothrix polyspora. Redundancy analyses were performed to determine a relationship between samples and environmental factors. The parameters nitrate, iron(III) oxide, iron(II) oxide and ammonia create differences between bacterial communities in operated and not yet operated wells.

The diversity and composition of native microbial populations in water samples differed fundamentally from those influenced by operation.
MARINE HYDROCARBONOCLASTICUS BACTERIA AS WHOLE-CELL BIOSENSORS FOR N-ALKANES

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Background

Whole-cell biosensors are useful systems for the in-situ monitoring of seawater for hydrocarbons derived from oil spills. Currently available biosensors for hydrocarbons show limitations derived from the low solubility in water of hydrocarbons or the high ionic strength of seawater.

Objectives

We compared the performance of a biosensor system for the detection of short chain alkanes in seawater, hosted in either E. coli (commonly used in many whole-cell biosensors but not optimized for alkane degradation) or different marine hydrocarbonoclastic bacteria specialized in assimilating alkanes.

Methods

A set of reporter strains containing a plasmid harbouring the reporter circuit was constructed. Fluorescence assays were carried out to study the time-dependent response of the reporter strains to different alkanes at different concentrations, and the response of reporter strains to real samples such as seawater contaminated with petrol or crude oil.

Conclusions

Some of the marine hydrocarbonoclastic bacteria tested performed better than E. coli. These strains showed very good response to seawater containing pure alkanes, petrol and crude oil in just 2 hours. Alcanivorax borkumensis proved to be the best host for the biosensor system.
HOW THE PRESENCE OF ORGANIC POLLUTANTS AND METALLOIDS CAN INFLUENCE DIBENZOTHIOPHENE-DEGRADING BACTERIAL STRAINS

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Background

In an oil polluted site, several components of environmental concern can be found including Polycyclic Aromatic Hydrocarbons (PAH), Dibenzothiophene (DBT) and metals. Studies showed that the naturally occurring soil bacteria are able to transform them to safe end products (bioremediation).

Objectives

1- To find tolerance of *Burkholderia fungorum* DBT1 (B.DBT1) and *Burkholderia fungorum* 95 (B.95) to DBT and PAH mixture and in planktonic and biofilm forms
2- Evaluating the growth and transformation ability of these strains in presence of selenium and tellurium as sample inorganic pollutants.

Methods

The growth as biofilm in 96 well plate was established for both strains using Calgary Biofilm Device (CBD) with the method described by Ceri et al., 1999. Growth curves were drawn based on CFU counting on YMA plates. Biofilm photos were taken by Confocal Microscopy. Selenite and tellurite reduction efficacy assessment was performed based on protocols described by Kessi et al, 1999 and Turner et al, 1992 respectively.

Conclusions

Both strains can tolerate high concentrations of DBT (2048 ppm) in both planktonic and biofilm forms while the mixture of PAH can only be tolerated by planktonic form for both strains. B.95 is able to tolerate the maximum used concentration of selenite (2 mM) and transform more than half amount to elemental selenium. On the other hand B.DBT1 can only convert 0.5 mM of selenite, while is able to transform completely initial 0.1 mM concentration of tellurite to elemental tellurium.
VINASSE AS SUBSTRATE FOR GROWTH MICROBIAL CONSORTIUM-ALTERNATIVE FOR TREATMENT AND RE-USE WASTE.

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Background
Vinasse showed to be appropriate substrate for yeast growth in single culture. The use of vinasse as a substrate for microbial biomass production can be an alternative for the treatment and reuse of this waste.

Objectives
The objectives of this study were (i) to evaluate the biomass production by Bacillus subtilis and two strains of Saccharomyces cerevisiae in mixed culture using the vinasse as a substrate, (ii) investigate the nutritional quality of the product generated, (iii) to evaluate the effect of biological treatment on physico-chemical parameters and level of toxicity.

Methods
B. subtilis (CCMA 0087), S. cerevisiae (CCMA0137) and S. cerevisiae (PE2) were initially grown in 100 ml vinasse medium (49% v/v vinasse, 49% distilled water, 1% glucose and 1% yeast extract). The microorganisms grown in a consortium, two by two, and each consortium was conducted in three different cell concentration. During the incubation period were monitored the cell number and pH. After that, the dry biomass and the reduction in Chemical oxygen demand (COD) and biochemical oxygen demand (BOD) of treated vinasse was evaluated.

Conclusions
Based on these results, the best combination and concentration of microorganisms was selected for cultivation in bioreactor. During the incubation period the bioreactor same previous parameters were monitored. In addition to the COD and BOD, the treaty vinasse was evaluated for acute phytotoxicity. The data obtained showed that the two strains of S. cerevisiae in mixed culture with an initial population of 10⁷ cells/mL showed 176mg/L of biomass and reducing the power polluter of vinasse.
DISTRIBUTION AND GENOMIC ADAPTATION OF METHANOMASSILIICOCALES ECOTYPES IN WETLANDS AND ANIMAL INTESTINAL TRACTS

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\textsuperscript{2}Laboratory of Food Biotechnology, Institute of Food Nutrition and Health, Zürich, Switzerland
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Background
Methane (CH\textsubscript{4}) is a very potent greenhouse gas, mainly produced by methanogenic archaea as an endproduct of their anaerobic respiration. Methanogenic archaea are phylogenetically diverse and living in a broad range of anaerobic environments, among others, in wetlands and ruminant animals, which represent the major natural and anthropogenic CH\textsubscript{4} sources. Recently, a novel order of methanogens, the Methanomassiliicoccales, was discovered.

Objectives
We aimed to assess the environmental distribution of Methanomassiliicoccales in different wetlands (arctic and acidic/neutral temperate peat) and animal intestinal tracts. Additionally, a metagenomic characterisation of Methanomassiliicoccales enrichments was intended.

Methods
Screening for Methanomassiliicoccales 16S rRNA and mcrA genes showed that Methanomassiliicoccales are widely distributed in wetlands and animal intestinal tracts. Phylogenetic analyses of 16S rRNA genes revealed two distinct animal and wetland clusters, pointing to two fundamentally different ecotypes. Ruminant enrichment cultures grow in an anaerobic medium supplemented with trimethylamine and H\textsubscript{2} as substrates. Methanomassiliicoccales represented one major methanogenic group in the bovine rumen performing H\textsubscript{2}-dependent methylotrophic methanogenesis. Two partial Methanomassiliicoccales genome bins assembled from rumen metagenomes represented the two ecotypes and enabled the characterisation of the underlying genomic basis of their adaptations. Ecotype differences were reflected e.g. by the smaller genome size of the animal-associated ecotype as well as a higher abundance of genes for oxidative stress response in the soil ecotype.

Conclusions
Our findings suggest that Methanomassiliicoccales represent a novel ecologically important group of methanogens with distinct animal and soil associated ecotypes adapted to these different environments. Especially the animal-associated ecotype potentially plays an important role in global CH\textsubscript{4} formation.
OZONATION FOR THE REDUCTION OF THE MICROBIAL LOAD OF WASTEWATERS

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Background
Treated municipal wastewater contains microbial densities up to $10^6$ CFU/100mL. Ozonation is an oxidation process widely used to disinfect water due to the strong ozone biocidal properties.

Objectives
This work aimed at assessing the effectiveness of ozonation on the reduction of the microbial load of treated wastewater.

Methods
A synthetic wastewater containing a mixture of fungi and bacteria, including antibiotic resistant and endospore forming strains, was submitted to ozonation assays performed in a 1L lab-scale reactor, with contact periods of 15, 30 and 60 min. The treatment efficiency was assessed based on the microbial enumeration and quantification of 16S rRNA, and selected indicator genes, before and after ozonation and after three days of storage at room temperature.

Conclusions
Ozonation for 15 min reduced the total heterotrophic bacteria and fungi by 3 log and 1 log, respectively. Coliforms and enterococci were extensively removed, being detected at densities of about 5 CFU/100mL. However, 3 days after water ozonation, a strong recovery was observed, with total heterotrophs reaching $10^9$ CFU/100mL and fungi $10^2$ CFU/100mL. Total coliforms and enterococci were still detected after 3 days. After 30 and 60 min of ozonation, viable cell counts were below the detection limit (1 CFU/100mL) and no recovery was observed after 3 days. Using culture independent methods, it was possible to confirm those results, being observed significant reductions of selected indicator genes after 15-30 min treatments. However, both 16S rRNA and $bla_{TEM}$ genes, but not $vanA$, were detected after 3 days recovery.
MIGRATORY HABITS, AREA AND HABITAT INFLUENCE THE MICROBIOTA OF BIRDS IN THE DANUBE DELTA NATURAL RESERVE

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Background
Wild birds from aquatic habitats could, as reservoirs and vectors for antibiotic resistant bacterial pathogens, directly affect the health of animals and pose serious risks to residents and tourists, the microbial load and its variety increasing with the broadening of the habitat and depending on the migration area.

Objectives
The research aimed to monitor the variation and distribution of pathogenic bacteria in birds with different migratory habits, areas and habitats in the Danube Delta.

Methods
A total of 112 rectal and oro-pharyngeal swabs from 26 species of migratory and sedentary birds from the Danube Delta Natural Reserve were collected and subjected to classical cultivation or enrichment techniques. 113 strains were identified on Chromogenic UTI medium Brilliance™ and TCBS Cholera medium (Oxoid) as belonging to Vibrio, Pseudomonas, Proteus, Escherichia, Staphylococcus and Enterococcus genera. Pathogenic bacteria were prevalent in migratory versus sedentary birds, with alarming proportions of Vibrio spp. (23.89%), followed by E.coli/coliforms (21.26%) (table 1):

<table>
<thead>
<tr>
<th>Wild birds</th>
<th>V. cholerae, V. fluvialis</th>
<th>V. alginolyticus, V. metschnikovii</th>
<th>Pseudomonas spp.</th>
<th>V. mimicus, V. vulnificus</th>
<th>Proteus spp</th>
<th>Coliforms</th>
<th>Staphylococcus spp.</th>
<th>Enterococcus faecalis</th>
<th>E. coli</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sedentary</td>
<td>11.1</td>
<td>37.5</td>
<td>37.5</td>
<td>30</td>
<td>28.5</td>
<td>35.7</td>
<td>28.57</td>
<td>44</td>
<td>30</td>
</tr>
<tr>
<td>Migratory</td>
<td>88.9</td>
<td>62.5</td>
<td>62.5</td>
<td>70</td>
<td>71.4</td>
<td>64.3</td>
<td>71.43</td>
<td>56</td>
<td>70</td>
</tr>
</tbody>
</table>

Bacterial pathogens (%) isolated from cloacal and oro-pharyngeal specimens collected from wild bird species in the Danube Delta Natural Reserve (Table 1).
<table>
<thead>
<tr>
<th></th>
<th>7.96</th>
<th>7.08</th>
<th>7.96</th>
<th>8.85</th>
<th>6.19</th>
<th>12.39</th>
<th>18.58</th>
<th>22.12</th>
<th>8.87</th>
</tr>
</thead>
</table>

**Conclusions**

Thus, the bacterial load but not the spectrum of isolated species is influenced by migration site and habits rather than habitat.
BIODEGRADATION OF PERCHLOROETHYLENE AND TRICHLOROETHYLENE UNDER AEROBIC CONDITION

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Background

Tetrachloroethene (also known as perchloroethylene; PCE) and trichloroethylene (TCE) are highly chlorinated hydrocarbons, are the common groundwater contaminants in industrial areas. Dechlorination of hydrocarbons is a complex process and biological dehalogenation under aerobic condition also required more clarification.

Objectives

The hydrocarbon degrading bacterial communities produces mono/di oxigenase enzymes and has the potential to degrade the chlorinated hydrocarbons with co-metabolic pathways. Soil samples were collected from contaminated site and enrichment of specific microbial communities were obtained using methanol and toluene as sole carbon source.

Methods

The enrichment cultures Ui-mix and MEOH-1 were examined for the ability to degrade TCE and PCE under aerobic condition. Growth and degradation rate are examined using spectrophotometer and gas chromatography equipped with ECD. The colonies on the enrichment cultures are isolated, purified and identified based on the 16S rRNA gene sequencing.

Conclusions

We have developed two consortiums which can degrade the PCE and TCE completely in aerobic condition. Ui-mix and monocultures from the consortium showed highest degradation effectiveness to co-metabolize TCE along with toluene as sole carbon source. Similarly, the MeOH-1 and monocultures degraded PCE under aerobic condition. The consortium Ui-mix contain diverse group of bacteria incudes \textit{Chryseobacterium}, \textit{Pseudomonas}, \textit{Enterobacter} and \textit{Microbacterium}; similarly \textit{Pseudomonas}, \textit{Hyphomicrobium}, \textit{Cupriavidus}, and \textit{Microbacterium} are isolated from MeOH-1. The strains M3-1, M3-3 and M3-7 showed 98.4%, 95.3% and 94.5% removal efficiency of TCE and PCE within 20 days, respectively. Further studies have to be performed to optimize and enhance the removal efficiency and it gives a gateway for the bioremediation of chlorinated ethane.
METHANOGENIC ACTIVITY AND DISTRIBUTION OF THE METHANOMICROBIALES IN TRANSITIONAL BOG AND SWAMP FOREST PEAT PROFILES (POLESKI NATIONAL PARK, POLAND)
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Background

Peatlands are the largest natural carbon storage. Under good aeration conditions carbon is released as a CO₂ but when there is lack of oxygen, anaerobic brake down of organic matter take place and CH₄ release occur. In most peatlands with neutral pH, the main substrate of CH₄ production is acetate while in acidic, the main precursors are CO₂ and H₂. CH₄ production is attributed to the activity of methanogenic Archaea, which, in peat soils are represented in majority by the members of the Methanomicrobiales.

Objectives

The aim of presented work were: (1) to examine which layers of the peat soil profiles (from the surface top to a mineral base) in transitional peat bog and swamp forest surrounding eutrophic Lake Moszne in Poleski National Park (East of Poland) show the highest methanogenic activity; (2) to test the presence of the Methanomicrobiales in investigated peat materials.

Methods

Methanogens were detected in fresh samples via FISH technique using MG1200 probe. Soil samples were further incubated anaerobically (at 5-30°C) and CH₄ production (gas chromatography technique) was controlled.

Conclusions

The highest methanogenic activity (up to 39.46 mg CH₄ kg dw⁻¹ d⁻¹) below layers 45 cm (transitional bog) and 150 cm (swamp forest) were found. In all of the investigated samples the presence of coccus, bacillus and curved rods from the Methanomicrobiales were confirmed. Among investigated, morphologically diverse order, the dominant role plays coccus (up to 100%) while the rarest were representatives of curved rods (up to 9%).
RESPONSE OF THE COALBED ROCK METHANOTROPHIC COMMUNITY TO TEMPERATURE CHANGES

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The John Paul II Catholic University of Lublin, Lublin, Poland

Background

Methanotrophic bacteria are ubiquitous in the environment and play an important role in the global carbon cycling. Recently published research revealed that in spite severe oxygen limitation, aerobic methanotrophic species are also found in deep subsurface geological formations such as coalbed rocks. These rocks are excavated in huge amounts during coal exploitation and comprise a troublesome waste. The presence of the methanotrophic community creates an opportunity to reuse these materials for the purpose of methane biofiltration; however surface conditions vary much from these in situ, especially according to temperature, which in geological formations is stable (c.a. 30 °C).

Objectives

In the current study we investigated methanotrophic communities of the rocks originating from Upper Silesian (Poland) coal mines in terms of identity (16S rRNA sequences) and response to the temperature changes.

Methods

Assays were performed in hermetic microcosms where samples of 15 g of rock were incubated under mixtures of methane and air. Headspace gases were measured by means of gas chromatography. DNA was isolated using PowerSoil® DNA Isolation Kit (MoBio) and amplicons obtained with methanotroph-specific IF/IR, IIF/IIR primers.

Conclusions

The investigated microbial communities were heterogeneous and consisted of both type I and II methanotrophs. Their ability of to oxidize methane decreased along with temperature and in majority, yet not all, of the samples was inhibited at 10 °C. It was hypothesized that the potential of methane turnover in the coal bed rocks is related with identity of the methanotrophic inhabitants with type II methanotrophs (α-Proteobacteria) being responsible for methane oxidation at low temperatures.
UTILIZATION OF RHAMNOLIPIDS DURING BIOSURFACTANT-MEDIATED BIOREMEDIATION OF HYDROCARBON-POLLUTED SOIL

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Background

It was proven that rhamnolipids are environmental-friendly biosurfactants produced by \textit{Pseudomonas} species. There are many studies dealing with rhamnolipids-mediated bioremediation of hydrocarbons. However, it must be mentioned that both the environmental fate and the influence of rhamnolipids on microbial communities remain unexplained. We hypothesize that rhamnolipids could be more readily utilized by soil microbial communities than hydrocarbons.

Objectives

1. to investigate and compare biodegradation rates of rhamnolipids and hydrocarbons.
2. to investigate the effect of rhamnolipids on quantitative and qualitative composition of microbial community

Methods

Biodegradation rates – HPLC-MS, structure of the microbial community – Illumina.

Conclusions

In most cases at least 80% of monorhamnolipids and dirhamnolipids were degraded during field experiment. The biodegradation rates of hydrocarbons was significantly lower compared with rhamnolipids. Rhamnolipids are more preferentially degraded by soil microorganisms than hydrocarbon compounds. Thus, we suggest that the role of rhamnolipids during bioremediation processes may be negligible.

Ławniczak Ł, Marecik R, Chrzanowski Ł, Contributions of biosurfactants to natural or induced bioremediation, \textit{Applied Microbiology and Biotechnology} (2013), 97. 2327–2339.
CHARACTERIZATION OF A PSEUDOMONAS SP. ISOLATED FROM BRAZILIAN SOIL AND ITS ABILITY TO DEGRADE ATRAZINE.

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Background

Atrazine is a selective and systemic herbicide used for controlling weeds in several crops. It is considered an environmental pollutant due to its high potential to contaminate the surface water and groundwater. Pseudomonas sp. ADP, the model strain for atrazine biodegradation, mineralizes this herbicide using it as a nitrogen source.

Objectives

Characterization of a strain isolated from a Brazilian soil sample and its ability to degrade atrazine.

Methods

The soil sample was collected in Ribeirão Preto, São Paulo, and processed according to the method describe by Gargouri et al. (2013), with modifications. The strain was selected from ATZ-R medium, in accordance with the formation of clear zones around the colonies. The identification was performed by 16S rRNA sequence analysis. PCR reactions for detection of atz genes (atzA, atzB, atzC, atzD, atzE and atzF) was carried out according to the methodology of Devers et al. (2004) and, then, sequenced. Plasmidial DNA extraction was performed according to the alkaline lysis technique, previously described by Birboin & Doly (1979). The degradation test was performed in solid and liquid ATZ-R medium.

Conclusions

The strain was identified as Pseudomonas sp., presented all atz genes and a 35 MDa plasmid, however, only the atzA gene was detected in this plasmidial DNA. The strain was capable to mineralize this herbicide after 48 hours in solid medium, indicated by the clear zones around the colonies. The analysis of the ATZ-R liquid medium in spectrophotometer (225 nm) showed a significant decrease of atrazine after 48 hours of incubation.
THE MICROORGANISM THAT NEVER READ THE LITERATURE - FASTEST GROWING PHOTODAMAGE TOLERANT ALGA ISOLATED FROM DESERT CRUSTS, DEPENDS ON PIONEER FILAMENTOUS CYANOBACTERIA SPECIES TO SURVIVE DESICCATION

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²Chemical Engineering, Shamoon College of Engineering, Beer-Sheva, Israel

Background
With the premise that unique capabilities can be found in microorganisms facing extremely harsh conditions, we focused our efforts on desert biological sand crusts (BSC), one of the harshest environments to support life. Recently, we have isolated a small green alga (named Chlorella ohadii), which does not conform to some of the basic fundamentals in microbial ecology and photosynthesis:

Objectives

1. Acquired ability to acclimate to extreme environments is usually accompanied by reduced performance under optimal conditions. We intend to show that C. ohadii does not obey this rule. In its natural habitat, C. ohadii is facing diurnal desiccation-hydration cycles, vast temperature amplitudes and extremely high illumination intensities, yet when grown under optimal laboratory conditions it exhibits the fastest growth rates ever reported for an alga.

2. After many years of research, some feel that we have elucidated the functioning of the photosynthetic machinery, and what sets the upper limit for algal growth. However, the unparalleled fast growth, extremely high photosynthetic rates and resistance to photodamage, suggest this may not be the case. Namely, C. ohadii has been shown to be completely resistant to photoinhibition, and its productivity was unaffected by irradiances as high as twice full sun light.

3. Survival of C. ohadii in its BSC habitat depends on close association with filamentous cyanobacteria; a unique and novel mode of interspecies interaction.

Methods

We wish to present detailed physiological analysis of the unique properties supporting this phenomenal growth and resistances, combined with insights from their genomic, transcriptomic and metabolomic characterization.

Conclusions
Background
As water travels from treatment plant through the pipelines to consumer’s taps, the diverse drinking water microbiome varies in abundance and in composition as a result of a large number of variables. A quantitative understanding of relevant variables is essential towards ensuring better control of drinking water quality. Increasingly, DNA sequencing approaches such as 454 pyrosequencing and Illumina MiSeq sequencing are being utilised to this end. Yet, little has been done to understand the influence of sample collection protocols on the data collected through DNA sequencing approaches. The ability to accurately understand the factors that affect the change in the drinking water microbiome is critically reliant on the robust approaches for sample collection.

Objectives
The goal of this study was to understand the effect of sample volume and sampling flow rate on the structure and membership of drinking water bacterial communities. Understanding these effects is especially critical in drinking water systems, which represents a low biomass but high diversity aquatic environment.

Methods
To do this, we sampled at five different residential sampling locations in the City of Glasgow, UK. This sampling was conducted such that five different sample volumes, ranging from 1 to 20 litres, were filtered to harvest microbial cells at each sampling location and this exercise was repeated at two different flow rates at the faucet (laminar and turbulent flow regimes).

Conclusions
We will provide a detailed overview of these two variables on the reproducibility and reliability of DNA sequencing based investigations of the drinking water microbiome.
Background
Members of the genus *Nitrospira* are dominant in many natural habitats and of vital importance for wastewater treatment. They are chemolithoautotrophic organisms capable of growth with nitrite and CO$_2$ as sole energy and carbon source. It further has been demonstrated that some *Nitrospira* can utilize simple organic carbon compounds and molecular hydrogen as alternative substrates and can switch to nitrate reduction under oxygen limitation. However, this metabolism has been assumed to be a survival strategy and in most systems studied so far *Nitrospira* was outcompeted by denitrifying organisms when anoxic conditions prevailed.

Objectives
Here, a community of nitrogen cycle bacteria was enriched from the anaerobic compartment of a biofilter connected to a recirculating aquaculture system.

Methods
The culture was fed with filtered water from the aquaculture system, supplemented with ammonium, nitrite and nitrate.

Conclusions
A stable enrichment culture was obtained which anaerobically converted ammonium, nitrite and nitrate into dinitrogen. The culture was dominated by anaerobic ammonium-oxidizing *Brocadia* species, a denitrifier (*Denitratisoma*), and a *Nitrospira* species. Anammox bacteria and *Nitrospira* co-aggregated, while the denitrifiers formed separate clusters. This culture shows that anaerobic ammonium oxidizers and nitrite oxidizers, who were considered to be mutually exclusive, can be grown as a stable co-culture. Furthermore, this shows that some *Nitrospira* species can be competitive under nitrate-reducing conditions. This study demonstrates another unexpected lifestyle for an organism believed to be only competitive under aerobic lithoautotrophic conditions. It further indicates that *Nitrospira* in combination with anammox can be of interest for anoxic wastewater treatment systems.
OXALATE-DEGRADING ABILITY IS UBIQUITOUS IN DIAZOTROPHIC AZOSPIRILLUM
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¹Dept. of Crop Science, Agricultural University of Athens, ATHENS, Greece

Background
Oxalate-degrading bacteria are known from a wide variety of ecological niches and different taxonomical origins. They play important roles in local and global carbon-cycling and, when present in the guts of mammals, they seem to provide protection against excess oxalate uptake and renal stones.

Objectives
In the present study, we aim to investigate the ability to grow at minimal medium with Ca-oxalate or Na-oxalate as a sole carbon source of 30 diazotrophic Azospirillum strains isolated from the rhizosphere of various field grown gramineous plants species grown at different geographical Greek locations, molecular characterized at our previous research works, and reference type strains. Most of these strains tested for the ability to grow at minimal medium with oxalate crystals isolated from Amaranthus blitum (family Amaranthaceae) as a sole carbon source. Additionally, we investigated for the presence of formyl-CoA transferase (frc) gene a molecular marker used for the specific detection and identification of oxalotrophic bacteria involved in oxalate degradation.

Methods
Minimal media with Ca-oxalate/Na-oxalate or oxalate crystals from plant were used as a sole carbon source for bacterial growth. Molecular biology techniques and phylogenetic analysis for the frc genes.

Conclusions
The majority of the examined strains display an ability to use oxalic acid as a sole carbon source. This result and the presence of the frc genes shows that these strains can metabolize oxalate via the oxalyl - CoA - decarboxylase pathway to produce formate and incorporate it in their metabolism.
DETOXIFICATION AND BIODEGRADATION OF ENVIRONMENTAL POLLUTANTS BY MARINE MICROBIAL CONSORTIA

G.A.L. VIEIRA¹, L.A. DUARTE¹, L.D. SETTE²
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Background
The high industrial activity causes several environmental problems, mainly due to the discharge of wastes. Marine environments are susceptible to contamination by industrial waste and also may represent a target niche for microbial prospecting for bioremediation. Marine microorganisms are adapted to saline conditions and have potential for being used in many processes.

Objectives
To evaluate the potential of microbial consortia in detoxify and degrade environmental pollutants.

Methods
Microbial consortia were structured in different combinations using four ligninolytic fungi from marine environments, two bacteria from oil reservoir and two lipolytic yeasts from Antarctica, previously selected based on their capacity to produce enzymes and to degrade hydrocarbons. Erlenmeyer flasks containing 50 ml of mineral medium, the microbial consortium and the pollutant (RBBR 500 ppm and Diesel oil 1% vv) were kept in incubators for 7 (RBBR), 14 and 21 (diesel oil) days at 140 rpm and 28 °C. Samples were analyzed in Microtox for acute toxicity and ligninolytic enzymes (Lac, MnP, LiP) were quantified.

Conclusions
Consortium 7 showed promising results in the studies with RBBR dye, presenting discoloration up to 70% and a very low rate of mycelial adsorption (11%). Enzymatic production of Lac was up to 141 U/L. LiP and MnP production were very low. All samples containing RBBR were detoxified. Samples containing diesel oil presented no detoxification and the production of ligninolytic enzymes was higher for MnP (114 U/L). A longer period of incubation and/or a lower concentration of oil will be tested, since this compound is highly complex.
EFFECTS OF DUST STORM EVENTS ON TUBERCULOSIS INCIDENCE RATE IN NORTHWEST OF CHINA

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1Key Laboratory of Desert and Desertification,
Cold and Arid Regions Environmental and Engineering Research Institute CAS,
Lanzhou, China

Background

Tuberculosis (TB) is a major public health problem in China. Gansu has very high annual attendance rate of TB in China, and the province is also famous because of its severe dust storms. The epidemic timing starts in February and ends in July, and the dust storm mainly distribute throughout the spring and early summer, which strongly indicate a close linkage between causative agent of TB and dust storm events.

Objectives

We investigated the general impact of dust storms to TB across time by analyzing variation in weekly clinic visits in Gansu during 2005-2012 in relation with the climatic event.

Methods

We applied Mann-Whitney-Pettitt test and regression model to detect the seasonal periodicity of TB and dust storm in a time series, and assess relationships between the meteorological variables and weekly TB clinic visits.

Conclusions

By comparing the information on cases of TB of Gansu weekly reported with dust storm events, we found a clear link between population dynamics of the disease and the climate disaster: the onset of epidemics and the dust storm shared the almost same mean week. Particulate matter might be the direct reason for the outbreak of TB in dust storm days. To our knowledge, this is the first population-based study that provides a clear demonstration that epidemic of TB was affected by dust storm in China, which will help understanding the association between this environmental problem and the evolution of epidemic disease.
GENETIC INSIGHTS INTO THE SULFUR OXIDATION PATHWAY OF SULFUR CHEMOLITHOTROPHS LIVING IN FRESHWATER LAKES

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Background
Chemolithotrophic sulfur-oxidizing bacteria are capable of oxidizing inorganic sulfur compounds as sources of energy for growth. A number of genomic studies have been taken place focusing on sulfur oxidizers living in habitats where a sufficient amount of reduced inorganic sulfur compounds are available. These studies revealed that taxonomically diverse sulfur chemolithotrophs use different enzymes for the oxidation of various sulfur compounds. On the other hand, sulfur chemolithotrophs living in freshwater lakes, where the availability of reduced inorganic sulfur compounds is generally low, have been overlooked. Recently, we revealed the dominance of betaproteobacteria in the sulfur-oxidizing bacterial community of freshwater lakes, and isolated several of them in pure culture.

Objectives
The aim of this study is to gain genetic insights into the sulfur oxidation pathway of sulfur chemolithotrophs isolated from freshwater lakes.

Methods
The genomes of the new isolates were sequenced and then analyzed focusing on the composition and phylogeny of genes for sulfur oxidation. In addition, their genomes were compared with available complete genomes of other sulfur oxidizers.

Conclusions
As the first comprehensive comparative genomic analysis focused on sulfur chemolithotrophs from freshwater lakes, the results of our study revealed that they possess common sets of genes putatively involved in sulfur oxidation. One of these gene sets was shared only by freshwater sulfur-oxidizing betaproteobacteria.
NITROTOGA - A MAIN NITRITE OXIDIZER IN LOW TEMPERATURE ENVIRONMENTS

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²Microbiology, Radboud University Nijmegen, Nijmegen, Netherlands

Background

Candidatus Nitrotoga arctica is a cold adapted nitrite-oxidizing bacterium (NOB), which was cultivated from the Siberian Arctic. Since 2007, Nitrotoga-like NOB could be detected in different natural and technical environments e.g. beneath the Antarctic ice sheet, in cave systems, in wastewater treatment plants and in aquaculturing facilities. Thus, they are more distributed than previously assumed and of importance for the global nitrogen cycle as well as for a successful nitrification in wastewater processing.

Objectives

The aim of this project is to gain a deeper understanding of the phylogenetic diversity within the genus Nitrotoga belonging to the Betaproteobacteria and to compare representatives derived from natural and artificial ecosystems.

Methods

The 16S rRNA gene was analysed combining specific and semi specific primer pairs for Eubacteria and Nitrotoga. Additionally, the beta subunit of the key enzyme of nitrification in Nitrotoga and other NOB, the nitrite oxidoreductase B (nxrB), was sequenced.

Conclusions

So far, Nitrotoga-like bacteria from widespread habitats do not differ noticeably in their 16S rRNA gene sequence from each other, not even those from extreme or technical origin (similarity > 98%). Although they might possess various physiological features, different species cannot be recognized on basis of the 16S rRNA gene. A higher resolution was obtained by comparing the sequences of the gene nxrB. In contrast to the ubiquitous and phylogenetically ancient NOB genus Nitrospira, Nitrotoga does not seem to be very diverse so far. However, its high abundance in various environments makes it an important player in nitrification.
NAPHTHALENE DEGRADATION BY SULFATE-REDUCING MARINE COMMUNITIES

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Background
Polycyclic aromatic hydrocarbons (PAHs) are ubiquitously distributed to the environment through the extensive petroleum and chemical industries. These toxic chemicals are of concern because of continual, widespread release.

Objectives
The bicyclic hydrocarbon naphthalene undergoes aerobic and anaerobic transformations, however our understanding of the details of anaerobic naphthalene degradation lags behind that of aerobic processes.

Methods
Sediment was obtained from Tuckerton, NJ, where PAHs are introduced through nonpoint deposition to an otherwise relatively pristine ecosystem. Anaerobic enrichment cultures were established with 10% anoxic estuarine sediment under sulfate reducing conditions and amended with 500μM naphthalene as the sole carbon source. Naphthalene concentrations were monitored using GC/FID.

Conclusions
Naphthalene loss was observed in 139 days in primary enrichments, followed by 48 days in transfer cultures. Subsequent transfers receiving 30-50% active cultures are sediment free and degrade naphthalene within 21 days. Stoichiometric loss of 3mM sulfate per 500μM naphthalene is observed. In the presence of molybdate, which inhibits sulfate reduction, no naphthalene degradation occurs, linking sulfate reduction to naphthalene metabolism. Distinct bands unique to naphthalene degrading cultures are seen in 16S rDNA analyzed by DGGE, indicating the enrichment of naphthalene degrading bacteria. By incorporating ¹³C-Naphthalene into DNA and RNA via stable isotope probing (SIP), the active microbes responsible for naphthalene degradation can be determined. Additionally, the recovery of 2-naphthoyl-CoA-reductase from ¹³C-DNA as well as identification of ¹³C intermediates can help to describe the metabolic pathway. This investigation of a robust sulfidogenic marine enrichment culture advances our understanding of the diversity of microbes able to anaerobically degrade naphthalene.
QUANTIFICATION OF SOIL MICROBIOLOGICAL ACTIVITIES BASED ON CROP TYPE (OAT, TRITICALE)*

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²Microbial Biochemistry, Institute of Biochemistry and Biophysics PAS, Warsaw, Poland
³Microbial Biochemistry, Warsaw University of Life Sciences, Warsaw, Poland

Background

Agriculture in Poland has been an important part of the country's economy (about 60% of Poland's total area is used for crop cultivation). It is known, that agricultural types of soil contain much smaller number of microorganisms, in comparison with natural soils. Thus we hypothesised that agricultural soils are biologically degraded and not being able to become naturally regenerated which may lead to their inability of regaining the satisfactory level of fertility.

Objectives

The aim of the study was to indicate which type of crop: oat or triticale is more favourable for sustaining soil microbiological activity.

Methods

Eight different soil units agriculturally used, and the same number of control soils not agriculturally exploited were studied. Soils were collected in April 2014, using Egner’s bow from the surface layer. Under laboratory conditions the following analyses were performed: pH, electric conductivity, total carbon, phosphorus and nitrogen content, microbial biomass (chloroform fumigation), respiration (GC), soil dehydrogenase activity (TTC) and DNA isolation (chloroform extraction method with CsCl₂ gradient). Spearman’s rho correlation coefficient was used to assess relationships between chemical and microbiological soil properties.

Conclusions

It was demonstrated that cultivated soils are biologically degraded, what was evidenced by lower values of all microbiological parameters in relation to control sites. Comparing crop type we found that triticale is more favourable for sustaining soils microbiological activity rather than oat. Determined positive correlations proved that pH, carbon, phosphorus and nitrogen content are the most important factors determining soil microbiological activities.

*Project was financed by the National Science Centre (Poland), granted on the basis of decision DEC-2013/09/D/NZ9/02482.
ENVIRONMENTAL FACTORS INFLUENCED ON SOIL MICROBIOLOGICAL ACTIVITY*
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Background
Many of recent studies have focused on documenting how soil bacterial communities are affected by specific environmental changes or disturbances. It was indicated that vegetation type, carbon and nutrient availabilities, and soil moisture may influence microbial activity at local scales, whilst soil pH seemed to be a better predictor at the continental scale.

Objectives
The aim of the study was to determine effect of selected environmental factors: pH, redox potential (Eh) and total carbon (TC) on soil microbiological activity, expressed as microbial biomass (MB), DNA content, potential ammonification (PA) and most probable number (MPN) of ammonifying bacteria.

Methods
Fifth representatives of Eutric Cambisols collected in April 2014 from the surface layer of agricultural sites were investigated. Spearman’s rho correlation coefficient was used to determine relationships between environmental and microbiological soil properties.

Conclusions
Positive correlations among pH, TC and all microbiological properties were reported. pH was found to be proportionally related with MB (0.73**), DNA (0.84***), PA (0.65**) and MPN (0.73**). Analogical trend demonstrated TC, what was confirmed by values of r coefficients, as follows: MB (0.64**), DNA (0.78**), PA (0.76**), MPN (0.83***). In the case of Eh it was noted that its lower values have stimulating effect on microbiological activity in Eutric Cambisols, as evidenced by following assigned rho: MB (-0.69**), DNA (-0.83**), PA (-0.75**) and MPN (-0.82**). Among investigated environmental factors only impact of moisture have insignificant character (p>0.05) in relations to tested soil microbiological properties.
*Project was financed by the National Science Centre (Poland), granted on the basis of decision DEC-2013/09/D/NZ9/02482.
THERMOPHILIC METHANOGENS REDUCE AMORPHOUS FE(III) OXIDES AND GENERATE CRYSTALLINE MAGNETITE ON CELL SURFACES

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²Bioproduction Research Institute, National Institute of Advanced Industrial Science and Technology, Sapporo, Japan
³Biomaterial Sciences, The University of Tokyo, Tokyo, Japan

Background
Methane production is affected by various environmental stimuli including the presence of Fe(III) oxides [1,5]. Inhibitory effect of Fe(III) oxides and Fe(III)-reducing ability in mesophilic methanogens has been reported [2,3]. There have been only limited information about thermophilic methanogens.

Objectives
Thermophilic methanogens were investigated for their ability to reduce poorly crystalline Fe(III) oxides (ferrihydrite) and the inhibitory effects of ferrihydrite on their methanogenesis.

Methods
*Methanothermobacter thermautotrophicus* strain ΔH², *Methanosaeta thermophila* strain PT¹ and *Methanosarcina thermophila* strain TM-1¹ and strain FE-1 were anaerobically cultivated in various methanogenic substrates supplemented media at 55 °C in the presence or absence of ferrihydrite. The amounts of methane and concentrations of Fe(II) were measured by gas chromatography and ferrozine method, respectively.

Conclusions
Fe(III) reduction was observed by the thermophilic methanogens only in the presence of H₂ as the reducing source [4]. While addition of ferrihydrite resulted in inhibition of methanogenesis, ferrihydrite reduction by the methanogens partially alleviates the inhibitory effects. *Methanosarcina thermophila* generated magnetite particles on its cell surfaces through ferrihydrite reduction. These findings suggest that interaction between Fe(III) minerals and methanogens has significant impacts on materials and energy cycles in anoxic environments.
References
OPPORTUNISTIC PATHOGENIC YEASTS ON THE SURFACE OF FRUITS
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Background
Infections caused by treatment-resistant non-albicans Candida species, such as C. tropicalis, have increased. It has become an emerging challenge in managing fungal infections. Previous researches have showed that C. tropicalis are opportunistic yeasts exist in environment.

Objectives
We are interested in whether the opportunistic yeasts may spread from environment into human body.

Methods
We isolated and characterized pathogenic yeasts on the surface of fruits from supermarkets. A total of 291 isolates of 83 species from 24 different types of fruits were recovered. We have determined their species and drug susceptibility.

Conclusions
Of the 83 species, 7 common pathogenic Candida species were detected. They included 16 C. guilliermondii, 15 C. famata, 3 each of C. parapsilosis and C. tropicalis, 2 each of C. krusei, C. lusitaniae and C. orthopsilosis. The drug susceptibilities of 162 of the 291 isolates were determined. Totally, 158 (97.5%), 104 (64.2%), and 102 (63%) isolates were susceptible to amphotericin B (MICs ≤ 4 mg/l), fluconazole (MICs ≤ 8 mg/l), and triadimenol (MICs ≤ 8 mg/l), respectively. One C. tropicalis isolate (F91) from wax apple had MICs at 64 mg/l for both fluconazole and triadimenol. It belongs to diploid sequence type (DST) 149, a genotype found in isolates from human as well as soil. Hence extra caution shall be taken when providing fruits or juice to severely immunocompromised patients since drug resistant pathogenic yeasts may be on the surface of fruits.
BIOREDUCTION OF TELLURIUM AND SELENIUM OXYANIONS BY UNACCLIMATED ANAEROBIC GRANULAR SLUDGE

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Background

Microbial reduction of chalcogen oxyanions such as tellurite and selenite is promising for development of innovative biotechnological processes for applications in bioremediation, wastewater treatment and production of functional nanomaterials.

Objectives

Here, we demonstrate rapid and stable bioreduction of tellurite, for the first time, using unaclimated anaerobic granular sludge, obtained from an operating upflow anaerobic granular sludge blanket reactor treating paper mill wastewater. In addition, bioreduction of tellurite in the presence of selenite was determined.

Methods

Bioreduction experiments were performed in serum bottles under different growth conditions using anaerobic granular sludge as source of microorganisms. Bioreduction of selenium and tellurium oxyanions profiles were determined at regular time intervals. Biochemical and microbial characterization of selenium and tellurium oxyanions reducing granular sludge was determined.

Conclusions

Tellurite was rapidly reduced to brown and then to black colored elemental tellurium by anaerobic granular sludge. The reduction of tellurite occurred at a much faster rate than the selenite when supplied individually. Complete reduction of 1 mM of tellurite and selenite was observed in 8 and 96 h, respectively. When both tellurite and selenite were present in the medium, the reduction of tellurite was not significantly affected. But, selenite reduction was found to be slower in the presence of tellurite. It is interesting to note an overlap between the reduction of both tellurite and selenite, which is needed for production of Se-Te functional nanomaterials. Repeated bioreduction of 1 mM of tellurite was sustainable during multiple cycles of feeding in a fed-batch experiment, indicates the applicability in biological treatment and formation of Te based nanomaterials.
MICROBIAL ASSOCIATIONS RELATED TO METHANE PRODUCTION IN AN UP-FLOW ANAEROBIC SLUDGE BLANKET TREATING MOLASSES WASTEWATER

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Background
Anaerobic digestion of organic wastes involves complex microbial interactions. The understanding of microbial interactions in anaerobic digestion can be utilized to construct strategies for methane production stably and effectively.

Objectives
In this study, methane production was evaluated in the up-flow anaerobic sludge blanket (UASB) reactor using molasses wastewater, and the active bacterial and archaeal communities were characterized using rRNA-based ion torrent sequencing.

Methods
The UASB reactor achieved a stable process performance at an organic loading rate of 1.7~13.8 g-COD·L⁻¹·d⁻¹ (87–95% COD removal efficiencies), and the maximum methane production rate was 4.01 L-CH₄·L⁻¹·d⁻¹ at 13.8 g-COD·L⁻¹·d⁻¹. Community analysis revealed that bacterial and archaeal communities shifted along with OLR, and Lactococcus and Methanosaeta, comprising up to 84% and 80% of the respective communities, drove the changes. Microbial network analysis was applied to identify associations among the relative abundances of bacterial and archaeal operational taxonomic units, environmental parameters, and reactor performance. The constructed network (average shortest path, 2.261; clustering coefficient, 0.514) revealed the prevalence of bacteria–archaea associations (50% of microbe–microbe associations) and contained seven microbial hub nodes including both bacterial and archaeal nodes. Interestingly, the Lactococcus and Methanosaeta were network hub nodes and positively correlated. In addition, they shared the other microbial hub nodes as neighbors and were positively correlated with methane production.

Conclusions
The results indicate that the close association between Lactococcus and Methanosaeta is responsible for the stable production of methane in the UASB reactor using molasses wastewater.
INFLUENCE OF SOIL PHYSICAL AND CHEMICAL PROPERTIES ON FUNGAL AND BACTERIAL COMMUNITIES IN 12 VINEYARD SOILS FROM NORTHERN ITALY

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Background

Soil chemical and physical properties, which are determined both by pedo-climatic factors and by management, can significantly influence microbial communities.

Objectives

Scope of this work was to assess whether given features can specifically influence defined microbial taxa in a man-managed environment such as the vineyard.

Methods

Soil samples were collected in 12 vineyards located in two neighbouring groups of hills, both renowned in Italy for their wines: the Euganean Hills, whose soils are of prevalent volcanic origin, mainly from trachyte and riolite rocks, and the Berici Hills, with a marine sedimentary genesis, mainly from clay-limestone. Soil bacterial and fungal communities were determined through 16S and ITS 454 pyrosequencing analysis, their relative taxa abundances were calculated and their levels of correlation with soil characteristics were tested.

Conclusions

In order of abundance, the dominant bacterial taxonomic groups across all samples were Actinobacteria, Alphaproteobacteria, Acidobacteria, Bacteroidetes, Betaproteobacteria, Chloroflexi, Gemmatimonadetes, Planctomycetes,
Deltaproteobacteria, Gammaproteobacteria. Among fungal phyla, Ascomycota largely dominated followed by Basidiomycota. Prevailing fungal classes were Dothideomycetes, Sordariomycetes, Agaromycetes, Leotiomycetes, Eurotiomycetes and Tremellomycetes. Some of these taxa and less abundant microbial groups were found to be significantly influenced by soil pH and texture. Moreover, a considerable number of significant correlations were found between the relative abundance of microbial taxa and the amount of several different chemical parameters such as total carbon, nitrogen, phosphorus, exchangeable bases and microelements.

This study provides novel insights into how soil structure and management can affect soil microbial community composition.
EFFECT OF BIOAUGMENTATION WITH BACTERIAL STRAINS ISOLATED FROM THE SOIL OF RICE FIELDS ON COMPOSTING PROCESSES USING RICE STRAW AND SEWAGE SLUDGE

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Background
Composting is a biotransformation process in which solid organic matter turns into mature and stabilized material by the action of microorganisms in an aerobic process. Microorganisms play a key role in composting; the performance of the process and the quality of the final product depends on the quantitative and qualitative composition of microbial communities associated to the different stages of the process.

Objectives
In this work, we describe the use of two bacterial strains, \textit{Paenibacillus barcinonensis} and \textit{Bacillus} sp. BP7, isolated from the soil of rice fields in the Ebro river delta in Spain, as bioaugmentants in composting processes using rice straw and sewage sludge as starting material.

Methods
Composting piles were created by blending sewage sludge and rice straw at a ratio of 2.6:1 (w/w fresh weight) with a moisture content of 60\%. To test the effect of the two strains, individual piles were seeded with $10^8$ uf/c gram of either \textit{P. barcinonensis} or \textit{Bacillus} sp. BP7, and a series of parameters were determined during the composting process.

Conclusions
Among the parameters determined, the humification index after 25 days had increased 40\% in the piles supplemented with \textit{P. barcinonensis} and 20\% in those supplemented with \textit{Bacillus} sp. BP7, a clear indicator of the efficiency of the bioaugmentation. This was also accompanied, in the case of \textit{P. barcinonensis}, by a change in the humic/fulvic acids ratio that went from 3.2 after 25 days in the control pile to 5.2 in the case of \textit{P. barcinonensis}. These results suggest the potential application of bioaugmentation in composting.
ISOLATION OF IBUPROFEN DEGRADING BACTERIAL STRAINS FROM COMPOSTING PILES MADE UP OF RICE STRAW AND SEWAGE SLUDGE

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Background
Pharmaceutical and personal care products (PPCPs) discharged with wastewater treatment plant effluents are an emerging surface water quality concern. Ibuprofen is the most widely used member of a diverse class of pharmaceuticals termed non-steroidal anti-inflammatory drugs (NSAIDs), many of which share a phenylacetic acid (PAA) core. Ibuprofen has been detected in bodies of water worldwide. Investigations into its environmental impact have found that ibuprofen induced changes on fish, plants, algae and microbial aquatic communities at environmentally relevant concentrations.

Objectives
The aim of this research was the isolation of bacteria with characteristics for potential bioaugmentation to enhance ibuprofen degradation in wastewater treatment plants an in the sewage sludge resulting from the activity of these plants.

Methods
Composting piles spiked with ibuprofen were created by blending sewage sludge and rice straw at a ratio of 2.6:1 (w/w fresh weight) with a moisture content of 60% to which a concentration of 1g/l of ibuprofen was added. Samples were taken at different days in the different phases, up to 42 days when ibuprofen had been completely degraded. Microorganisms extracted from these samples were tested in minimal medium supplemented with 500 mg/ml of ibuprofen and incubated at 24°C or 55°C, to discriminate mesophilic from thermophilic microorganisms.

Conclusions
Ibuprofen degradation occurred only in the samples incubated at 24°C. Plating of microorganism from these samples allowed the isolation of some 30-40 different colony morphologies that were again tested for ibuprofen degradation. Two of the isolates were confirmed to degrade ibuprofen and were partially characterized as belonging to the genus Klebsiella.
DEVELOPMENT OF A TEST FOR THE DETECTION OF BACILLUS THURINGIENSIS VAR KURSTAKI IN ENVIRONMENTAL SAMPLES

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Background

Bacillus thuringiensis was isolated for the first time in 1902 from diseased silkworm larvae and its commercial use as an ecological insecticide started in 1958. B. thuringiensis var kurstaki (Btk for short) is routinely used to control caterpillar pests both at large extensions and at small gardens. The crystal toxin is only active when ingested by the caterpillars and absorbed in the alkaline digestive tract. The toxin has a relatively narrow spectrum of action and does not affect mammals birds or other insects and besides, the persistence of Btk on foliage is of around five days and does not accumulate on the environment. Because of its relative safety Btk has also been in use as a simulant by armed and security forces in exercises of response to biological or bioterrorist attacks.

Objectives

In this work we have developed rapid tests that can be used for the detection of the presence of Btk on air or surface samples after controlled release, and to measure the efficiency of decontamination measures.

Methods

Polyclonal antibodies were raised in rabbits by injecting them with autoclaved commercial Btk spores.

Conclusions

Immune response was confirmed by Western-Immunoblot and immunofluorescence assays and the specific IgGs, after purification by affinity chromatography using Btk spores, were coupled to polystyrene beads, to develop a slide agglutination test, colloidal gold, as part of lateral flow immunochromatography tests, or polystyrene ELISA plates, to develop ELISA based assays. Preliminary results confirm the validity of slide agglutination test whilst lateral flow and ELISA are under development.
MOLECULAR IDENTIFICATION OF MICROORGANISMS USED FOR „BACTERIOGRAPHIE“

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Background

The knowledge that bacteria use pigments to guard against UV radiation gave Erich Schopf the idea of using bacteria as unique tool for creating colors. Instead of painting artists' colors on different painting supports, he applies bacteria on agar-based media. Initially invisible, the picture gets visible due to the growth of the bacteria used. Erich Schopf has called this technique 'Bacteriographie' (www.bacteriographie.com).

Meanwhile the ensemble of bacteria used for 'Bacteriographie' comprises around 900 microorganisms from different regions of the world (e.g. Austria, Germany, Greenland, Iceland, India, Indonesia, Ireland). They were collected from air, food, water, soil and recently also from holy water. Next to their ability to form pigments a general criterion is their growth at room temperatures within a foreseeable time period. Additionally, they have to undergo strict screening tests to perceive their suitability.

Objectives

Although some classical microbiological tests were done to get more information about the microorganisms themselves, the identity of the isolates is generally not known. Thus, this study focused on their identification at species-level using molecular biological methods.

Methods

A subset of the most important isolates (n = 120) was identified by 16S rRNA gene sequencing.

Conclusions
About 40 different genera could be identified. The relation between gram-positive and gram-negative species was nearly balanced. However, also a few fungal species (e.g. Pseudozyma spp., Cryptococcus spp.) were identified. Different species of the genera Pseudomonas, Flavobacterium, Microbacterium, Rhodococcus, Kocuria, Chryseobacterium and Arthrobacter were frequently detected, which form yellow, orange and red pigments.
FEMS-0910
Extreme environments

RAPID RECOVERY OF CYANOBACTERIAL PIGMENTS IN DESICCATED BIOLOGICAL SOIL CRUSTS FOLLOWING ADDITION OF WATER
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Background
Cyanobacteria in biological soil crusts were shown to move upwards in to track water, however this it is not known how widespread this process is in other crusts.

Objectives
To examine soil surface colour change to green and hydrotaxis following addition of water to crusts

Methods
We used pigment extraction, hyperspectral imaging, microsensors and 13C labelling experiments coupled to matrix-assisted laser desorption and ionization time of flight-mass spectrometry (MALD-TOF MS)

Conclusions
The topsoil colour turned green in less than 5 min following water addition. The concentrations of chlorophyll a, scytonemin and echinenon rapidly increased in the top <1 mm layer while in the deeper layer their concentrations remained low. Hyperspectral imaging showed that, in both wet and dehydrated crusts, cyanobacteria formed a layer at a depth of 0.2-0.4 mm and this layer did not move upward after wetting. 13C labelling experiments and MALDI TOF analysis showed that Chl a was already present in the desiccated crusts and de novo synthesis of this molecule started only after 2 days of wetting due to growth of cyanobacteria. Microsensor measurements showed that photosynthetic activity increased concomitantly with the increase of Chl a, and reached a maximum rate approximately 2 hours after wetting. We conclude that the colour change of soil crusts to green upon water addition was not due to hydrotaxis but rather to the quick recovery and reassembly of pigments.
GEOMICROBIOLOGY OF JAZAN SABKHA, SAUDI ARABIA

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Background
Sabkha is a unique environment of economic and geologic importance with a valuable resource of distinctive features. Saudi Arabia possesses coastal Sabkhas which are extensively distributed along both the eastern and western areas.

Objectives
Therefore, samples (I and II) with depth of 50 cm were collected from a remote salt marsh at the southwest of Saudi Arabia, which is called Jazan Sabkha, to unveil the bacterial community that survive in such extreme environment.

Methods
Each sample was cut horizontally at interval of 1 cm for the first 10 cm and at 5 cm for the next 20 cm. The samples are considered salty and sulphurous, where sodium and soluble chloride at the range of 11,000-34,000 ppm and 11,000-99,000 ppm, respectively, while sulfate varied from 2,070 to 18,820 ppm. Next generation sequencing (454 pyrosequencing) was applied to determine the bacterial communities inhabiting each layer.

Conclusions
Overall, Pseudomonas Genus was dominated in both samples with 46% and 39% for I and II, respectively. Pseudomonas putida was identified in each cross-section at 1 cm with variable percentages. In addition, P. veroniim, P. cannabina and Herbaspirillum rubrisubalbicans were presented with less percentages. Leptospirillum spp. were detected at lower layers and at they were optimized between 15-20 cm while Methyllobacterium tardum appeared at their maximum around 7-9 cm. Most of genera saline species of Halobacteriaceae family like Halorhabdus spp., Halonotius spp., Halobaculum spp., Halomicrobium spp., Halorubrum spp., Halogranum spp., Haloplanus natans, Salinibacter spp. and Natronomonas spp. were detected within 1-2 cm near to the salty crusty surface of Sabkah.
Background
Investigations of extreme environments are important for the study of evolution relationships, discovery of new species and various ecological relations among organisms. Besides, new metabolites and novel enzymes, metabolic pathways of organisms' can be discovered from extreme environments.

Objectives
In this study, metagenomics approach is used to isolate potential biotechnological novel enzymes from Lake Acıgöl. Especially we have focused on proteases which are ubiquitous enzymes and they have crucial roles in different industrial applications.

Research area
Lake Acıgöl is located between Afyon, Denizli and Burdur city boundaries in Aegean region, Turkey. It is a good example for extreme environments due to its high salinity (about 200g/L NaCl) which makes the lake suitable for halophilic microorganisms.

Methods
Bacterial population of Lake Acıgöl have already been identified and Lake bacterial population consist of 38.8% Uncultured Bacterium, 30.8% Firmucutes, 15.3% Bacteriodetes, 7.6% Gammaproteobacterium and 7.6% Deltaproteobacterium. Because of the high population of uncultured microorganisms Lake Acıgöl is a potential candidate for novel enzyme exploration. To isolate extremophilic protease enzymes from Lake Acigol sediment samples were collected from different part of the Lake and Metagenom (DNA) isolation was carried out by manually according to Zhou et. Al.,1999 [1].

Conclusions
Metagenom library was constructed by using pUC19 plasmid. Screening studies for possible protease enzymes will be carried out using 1% of skim milk LB agar plates and Salt Tolerance Screening Method according to Culligan et. Al., 2013 [2].
Background
Microbe-mineral interactions have become of interest for space exploration as microbes can biomine useful elements from planetary regolith, which could serve as nutrients in a life support system.

Objectives
Therefore, this research aimed to assess the impact of space conditions on physiology of Cupriavidus metallidurans CH34 and identifying the molecular mechanisms behind microbe-mineral interactions on basalt, a lunar-type rock.

Methods
Survival of C. metallidurans CH34 was monitored after a 3-month period in mineral water with or without basalt by plate counts, flow cytometry, ICP-MS and microscopy. The effect of space conditions was studied via a flight experiment on board the Russian PHOTON-M4 capsule.

Conclusions
The results obtained from ground experiments showed that the cultivable fraction dropped to 10% indicating a transition to a more dormant state. In the presence of basalt, CH34 remained viable, attached and formed a biofilm. The space flight experiment indicated more viable cells compared to the ground experiment both in the absence and presence of basalt, indicating a positive effect of space flight on survival.
Additional physiological and molecular analyses are on-going to confirm these observations and to determine the molecular processes.

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MEMBRANE-BOUND AMYLOPULLULANASE IS ESSENTIAL FOR STARCH METABOLISM OF SULFOLOBUS ACIDOCALDARIUS DSM639

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Background
The growth of Sulfolobus shows that these species prefer starch or long-chain maltooligosaccharides to maltose as a carbon source and have both α-amylase and α-glucosidase activities. The annotation of the genome sequence of S. acidocaldarius assume that the gene encoding amyloolytic enzyme, Saci_1162, within the putative maltose/maltodextrin ABC transporter operon (Saci_1160 to Saci_1166) is involved in the degradation of starch or α-linked polymers into small maltooligomers.

Objectives
A putative amyloolytic enzyme, Apu (Saci_1162), was characterized whether it is involved in the starch metabolism in S. acidocaldarius.

Methods
The physiological role of Apu in starch metabolism was investigated by the growth and starch degradation pattern of apu disruption mutant as well as biochemical properties of recombinant Apu.

Conclusions
The Δapu mutant lost the ability to grow in minimal medium in the presence of starch, and the amyloolytic activity observed in the membrane fraction of the wild-type strain was not detected in the Δapu mutant when the cells were grown in YT medium. The purified membrane-bound Apu initially hydrolyzed starch, amylopectin, and pullulan into various sizes of maltooligosaccarhrides, and then produced glucose, maltose, and maltotriose in the end, indicating Apu is a typical endo-acting amylopullulanase which belongs to glycosyl hydrolase family 57. The maltose and maltotriose observed in the culture medium during the exponential and stationary phase growth indicates that Apu is the essential enzyme to initially hydrolyze the extracellular starch into small maltooligosaccharides to be transported into the cell.
FEMS-0495
Extreme environments

MICROBIAL COMMUNITY PROFILE OF THE IRON-SULPHATE RICH WATERS OF RIO SUCIO (BRAULIO CARRILLO NATIONAL PARK, COSTA RICA)
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Background
Río Sucio ('Dirty river') in Braulio Carrillo National Park (Costa Rica) is a river of volcanic origin that possesses a characteristic yellow color due to the presence of high amounts of mineral deposits from Irazu Volcano.

Objectives
To perform a physicochemical characterization of the Río Sucio water and to analyze the impact of these parameters on the microbial community that inhabits the river.

Methods
Physicochemical analysis was performed by ion-exchange chromatography (IC), inductively coupled plasma mass spectrometry (ICP-MS), scanning electron microscopy with electron dispersive spectrometer (SEM-EDS) and X-ray diffraction (XRD). Analysis of the microbial community was performed by analysis of the V5-V6 hypervariable regions of the 16S rRNA gene. PCR amplicons were sequenced on an Illumina MiSeq instrument and the sequences compared against the RDP database (Ribosomal Database project – http://rdp.cme.msu.edu/).

Conclusions
Physicochemical analysis revealed that Río Sucio is a slightly acidic river (pH 5.0) containing high concentrations of sulphate (502 ± 29 mg/L), calcium (96.6 ± 2.2 mg/L), magnesium (27.36 ± 0.16 mg/L), aluminium (17.1 ± 1.9 mg/L) and iron (5.20 ± 0.11 mg/L). The precipitated material in the bottom of the river was composed mainly of iron oxides, as indicated by SEM-EDS and XRD. Microbial diversity analysis showed that the most abundant organisms are closely related to sulfur and iron oxidizing bacteria such as Sulfuritalea spp., Ferrovum spp., Acidithiobacillus ferrooxidans, Leptospirillum ferrooxidans and Ferrithrix spp. These observations suggest that Río Sucio has an acid rock drainage environment where the chemical composition is biologically driven by sulfur and iron oxidizing bacteria.
Background
Biological systems are frequently exposed to microwave radiation. Many studies have investigated the influence of microwaves on these systems, but controversy over methods to distinguish between thermal and non-thermal microwave effects remains.

Objectives
To differentiate between non-thermal and thermal microwave effects on a physiology of a microorganism, a thermophilic bacterium was grown in a constant-temperature microwave or a convection oven. Comparing the growth properties of the thermophile in these conditions will reveal non-thermal microwave effects on cell growth and
physiology. Biophysical and biochemical analysis will demonstrate changes in morphology and chemical composition arising from microwave exposure.

**Methods**

Cell growth was analyzed by optical density (OD) measurements (supported by independent quantitative DNA analysis), and cell morphologies were characterized using electron microscopy imaging (SEM, TEM), dynamic light scattering (DLS), and atomic force microscopy (AFM). AFM was also used to probe the biophysical characteristics of the cells, in conjunction with nano-infrared spectroscopy (Nano-IR). Attenuated total reflectance infrared spectroscopy (ATR-IR) and fatty acid methyl ester (FAMEs) analysis were used to determine biochemical differences between cells grown in microwave and oven conditions.

**Conclusions**

Thermophilic bacteria were grown in a synthetic microwave such that thermal effects and microwave effects were distinguishable. These data demonstrate that there are physiological differences between cells cultured in a dielectric field and a convection oven, and that microwaves induce non-thermal changes to the structure, physiology, and chemical composition of the organism.
DO THE PROTEINS INDUCED BY HIGH HYDROSTATIC PRESSURE IN FUNGI PLAY A ROLE IN TOLERANCE TO PRESSURE?

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Background
Deep-sea is a fascinating environment, which inspite of the extreme conditions of pressure, temperature and nutrients, is home to many life-forms. In order to survive here, the organisms need to develop mechanisms to counter these conditions. Fungi, one of the most ubiquitous eukaryotic forms of life, have also been reported from deep-sea sediments. In order to survive, they undergo both morphological and biochemical adaptations. Induction of proteins is one such response elicited by fungi for survival.

Objectives
In our study to understand the mechanism of survival by fungi occurring in deep-sea conditions, we are looking for proteins expressed by fungi and their role in tolerance to high hydrostatic pressure using LCMS QToF.

Methods
To understand the mechanism of survival of fungi, in deep-sea conditions, we examine tryptic digests of proteins expressed under various hydrostatic pressure using LCMS QToF.

Conclusions
The results showed that some proteins were up-regulated whereas some were down-regulated at higher pressures. Some over-expressed proteins, probably playing a role in pressure tolerance were 30 KDa heat shock protein and heat shock protein SSC1 mitochondrial. Unfortunately, none of the up-regulated proteins could be considered as ones produced exclusively in response to high hydrostatic pressure as all are reported in response to other stresses also. Along with these proteins, there were several hypothetical proteins to which no function has been assigned so far. These could well prove to be the ones limited to pressure stress, which needs to be investigated further by either inactivating the proteins or using knock-down mutations.
Background
Urmia Lake in the Northwest of Iran is the second largest hypersaline lake in the world which has been explored with great haloarchaeal diversity. Extreme halophilic Archaea are widely distributed in hypersaline habitats.

Objectives
During the course of biodiversity studies in Urmia Lake, several new extreme halophilic Archaea were isolated. Amongst them we chose strain DC8^T for further characterization.

Methods
Strain DC8^T has been characterized taxonomically using polyphasic approach. According to minimal standards we performed Molecular, biochemical, morphological and physiological tests for characterization.

Conclusions
16S rRNA gene sequencing showed that strain DC8^T is a member of the family Halobacteriaceae; however, its similarity was as low as 90.1% and 89.3% to haloarchaeal taxa including Halosimplex and Halobaculum type species, respectively. The cells of strain DC8^T were non-motile and pleomorphic and needs at least 2.5 M NaCl and 0.02 M MgCl₂ for growth. Optimal growth was at 4.0 M NaCl and 0.1 M MgCl₂. The optimum pH and temperature for growth were pH 7.5 and 45°C. The G+C content of its DNA is 68.1 mol%. Polar lipid analyses revealed that strain DC8^T contains phosphatidylglycerol, phosphatidylglycerol phosphate methyl ester and phosphatidylglycerol sulphate. The only quinone present was MK-8 (I1-H₂). The physiological, biochemical and phylogenetic differences between strain DC8^T and other previously described genera of extremely halophilic archaea suggest that this...
strain represents a novel species in a new genus within the family *Halobacteriaceae*, for which the name *Halositis urmiana* gen. nov., sp. nov. is proposed.
FEMS-1534
Extreme environments

LIFE ON THE EDGE: PROTEOBACTERIA AND HALOARCHAEA DOMINATE THE MICROBIAL COMMUNITY OF AN EXTREMELY HIGH ALTITUDE LAKE, LAKE LLULLAILLACO (6170 M)
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Background

Despite the crucial role of microorganisms in lake ecosystem function, they are typically poorly studied in high altitude ecosystems. In general, there are few reports of microbial diversity in lakes located at altitudes >5000 m. Lake Llullaillaco is a previously undescribed lake located at 6170 m on the Chilean side of the Llullaillaco volcano, the second highest volcano in the world (6739 m).

Objectives

The aim of this study was to describe the basic morphology of Lake Llullaillaco and the microbial diversity from water samples taken from the lake in February 2013.

Methods

Bacterial cultures were obtained using different culture media and microbial diversity was analysed by the pyrosequencing of 16S rRNA genes of Bacteria and Archaea. At the time of sampling, the lake had an area of 0.9 ha and maximum depth of 6.8 m. Water temperature ranged between 2 and 5°C and pH was 6.5. 24 bacterial isolates were obtained belonging to the Alpha-, Beta-, Gammaproteobacteria and Actinobacteria. Bacterial diversity was dominated by Proteobacteria (86.5%), Actinobacteria (11.9%) and Bacteroidetes (1.4%), with Beta-, and Alphaproteobacteria being the most abundant taxa. Archaeal diversity was dominated by Euryarchaeota and the Halobacteriaceae, with a high contribution of unidentified sequences.

Conclusions

This study is the first description of microbial diversity in Lake Llullaillaco: diversity was low and characterized by a combination of phylotypes previously described from cold environments and a series of undescribed microorganisms.

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Background

One of the main directions of microbial ecology of extremophiles is the study of halophilic microorganisms' biocenoses, which are mainly found in saline lakes and other water systems as well as saline soils and salt mines. Microbial biodiversity of salt mines and saline-alkaline soils distributed on the territory of Armenia is still poorly investigated.

Objectives

The aim of the present study was to investigate the microbial composition of saline-alkaline soils of the Ararat Plain and Avan subterranean salt deposits in Armenia.

Methods

For this study, clone-library construction of PCR-amplified 16S rRNA genes, DGGE-PCR fingerprinting and cultivation-dependent methods were used. 16S rRNA gene clone libraries and PCR-DGGE fingerprinting sequences were generated from total DNA extracts using universal archaeal and bacterial oligonucleotide primer sets.

Conclusions

Sequence analysis of bacterial and archaeal clone libraries and DGGE-PCR products from the samples indicated a dominance of Firmicutes and Euryarchaeota from the bacterial and archaeal domains, respectively. Several aerobic chemoorganotrophic endospore-forming bacteria were isolated from the salt samples and identified as representatives of the genus Halobacillus, Piscibacillus, Virgibacillus and Streptomyces. Five halophilic archaeal strains were isolated from salt stone samples. Four of the archaeal strains were most closely related to members of the genus Haloarcula (97-99 % similarity) and one strain most closely to the genus Halarchaeum (<97% similarity), indicating that Avan salt mine harbors a unique community of possible novel species.
The work was supported by the CPEA-2011/10081 grant from the Norwegian Cooperation Programme in Higher Education with Eurasia.
COPQ, A NOVEL FAMILY OF SMALL SECRETED PROTEINS WITH PUTATIVE RADICAL-SCAVENGING AND METAL-DETOXIFYING ACTIVITIES, ARE STRICTLY CONFINED TO THE GENERA RALSTONIA AND CUPRIAVIDUS

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Background

The soil bacterium Cupriavidus metallidurans CH34 is tolerant to a wide range of metals (1). Whole-genome transcriptomics demonstrated a common response in this organism for metal and oxidative stress (2). Some of the highly induced genes are well conserved within 14 C. metallidurans strains (3) and encode short proteins (coined the COPQ family) that possess a distinctive signal peptide and up to ten repetitive motifs [GSRD]XXD[PV][YF]T[DE]G[ASG].

Objectives

Bioinformatic analysis of the COPQ family.

Methods

Transcriptomic procedures are in (2). Structural predictions were performed with ITASSER and PHYRE-2. Sequence similarity analysis was done by iterative BLAST against the non-redundant protein sequence database (nrdb – version 2014/09/10). Motif detection was done using BioGrep against 2,616 bacterial proteomes (refpep – version 2014/01/09).

Conclusions

BLAST analyses and BioGrep motif searches against reference protein databases showed that the CopQ family of proteins is strictly confined to the highly related species of Cupriavidus and Ralstonia. COPQ proteins are predicted to form straight, slightly helical polypeptides, probably owing to the abundance of chain-disturbing and evenly spaced glycines and prolines. The aromatic rings of the Tyr and Phe residues at the center of the motifs are fully exposed allowing easy access to reactive oxygen species (ROS).

We propose that COPQ-family proteins have a radical-scavenging function shielding cells from oxidative attack and/or metal toxicity. Highly purified COPQ proteins in single or mixed fashion are to be tested for their anti-oxidant and metal-binding
properties.
(2) Monsieurs et al. (2011). Biometals, 24(6) 133-1151
BIODEGRADATION OF AROMATIC HYDROCARBONS BY HALOPHILIC ARCHEAE ISOLATED FROM ALGERIAN SEBKHAS

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Background
Actualy, the biodegradation of hydrocarbons (specialy PAHs) has recieved great attention, however, such high and fluctuating salinity promotes the loss of cell wall integrity, protein denaturalization, and changes in osmotic pressure, the biological treatment of industrial hypersaline waste-waters and the bioremediation of polluted hypersaline environments are not possible with conventional microorganisms.

Objectives
The aim of our study is the identification of microorganisms that could answer these criteria for the biodegradation of extreme environments.

Methods
Three strains halophilic archaea Haloarcula sp.D21(AM982816), Halovivax sp.A21(AM982815) and Natrialba sp.C21(HG423210) were isolated from the salt lake close to Ain Salah, Algeria. These strains show good growth on extreme saline culture media (up to 25%NaCl) in the presence of diesel, naphthale and pyrene as the sole carbon source. The strain Natrialba sp.C21 has a better growth on these media even at 35% of salt. However, the highest rate of biodegradation of these three hydrocarbons is labeled in the presence of extreme halophilic consortium containing the three strains. This biodegradation is marked by the high rate of growth of the strains measured by the optical density, the production of biosurfactans estimated by the emulsifying index and by the lowering of the surface tension of the culture media (below 40mN/m). In addition, the biodegradation of these hydrocarbons in the presence of the halophilic consortium is correlated with the rate of the concentration of NaCl and Mg\(^{2+}\) in the medium.

Conclusions
Thus, our halophilic haloarchae strains, specialy Natrialba sp.C21, are potential candidates for the degradation of pollutants at high salt concentrations.
MICROCYSTIN VARIANTS IDENTIFIED IN ARCTIC CYANOBACTERIA USING A
NEW ANALYTICAL SCREENING METHOD

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Background

Cyanobacteria produce a large variety of toxic secondary metabolites. In the Arctic and Antarctic, many unusual variants of microcystins (MCs) - the most common type of cyanobacterial freshwater toxins - have been detected. These variants are generally difficult to detect by standard methods in background-rich samples.

Objectives

Therefore, there is a need to develop a screening method that can be used for a large number of environmental samples.

Methods

In this study, 25 cyanobacterial samples from diverse biotopes in Svalbard (Arctic) were analyzed for the presence of MC. In a preliminary ADDA-MC specific ELISA, 18 of 20 samples were tested positive, an unusually high percentage. In contrast, a liquid chromatography-tandem mass spectrometry (LC-MS/MS) precursor-screening, newly designed to identify candidate MCs from the ADDA fragment ions, suggested the presence of MCs in nine of the 25 samples. In three of the latter, MC variants could be subsequently confirmed and identified by complete MS/MS analysis as variants of MC-RR and MC-LA, similar to those previously reported for polar habitats. In five of the same nine samples, a gene involved in toxin production - mcyE - was successfully amplified, validating the analytical data. The amplified gene sequences were 93-98% similar to mcyE genes of Nostoc, indicating that this genus could be responsible for MC production in these samples.

Conclusions

The presence of the unusual variants in the polar regions could help to understand the evolution and ecological function of the toxins. Moreover, the toxicological risk of unusual toxins is not yet known and might be an undetected threat.
‘EX-SITU’ PRESERVATION AND CHARACTERIZATION OF ANTARCTIC CYANOBACTERIA IN THE BCCM/ULC COLLECTION

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Background
The BCCM/ULC public collection of (sub)polar cyanobacteria is funded since 2011 by the Belgian Science Policy Office. An ISO9001 certificate was obtained for the public deposition and distribution of strains, as part of the multi-site certification for the BCCM consortium. BCCM/ULC is currently holding 160 public cyanobacterial strains and the catalogue is available on http://bccm.belspo.be/catalogues/ulc-catalogue-search.

Objectives
Continuous maintenance of living cultures, some of which are also cryopreserved, ensure the preservation and the possibility to rapidly deliver strains to clients for fundamental and applied research.

Methods
The main holding of the collection concerns (sub)polar strains isolated from different biotopes and representative of a large taxonomic diversity. The molecular characterization is underway, on the basis of 16S rRNA and ITS sequences, but also Multiple Locus Sequence Analysis and genome sequencing. In addition, cyanobacteria are known to produce a range of secondary metabolites (e.g. alkaloids, cyclic and linear peptides, polyketides) with various bioactivities. The potential of the polar strains to produce cyanotoxins and other secondary metabolites is currently studied by ELISA, LC-MS and the detection of genes involved in their production.

Conclusions
Due to the geographic isolation and the strong environmental stressors of the habitat, the exploration of these metabolites in Antarctic cyanobacterial strains seems especially promising for biotechnology or biomedical applications.
THE CRENARCHAEON IGNICOCUS HOSPITALIS IS A POLYEXTREMOPHILIC ORGANISM WITH UNUSUAL HIGH RADIATION TOLERANCE

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Background

The hyperthermophilic crenarchaeon *Ignicoccus hospitalis* was isolated from deep-sea hydrothermal vents living there strictly anaerobic and chemolithoautotrophic. From an astrobiological point of view, hyperthermophilic organisms are promising candidates for early Earth inhabitants, when the absence of oxygen, occasional meteorite impacts followed by heated up oceans, and the missing UV-absorbing ozone layer enabled higher radiation dose of short wavelength to penetrate Earth’s surface.

Objectives

To ensure DNA integrity, *I. hospitalis* needs proper DNA repair mechanisms to cope with today’s harsh environmental conditions in its natural habitat. These highly efficient DNA repair mechanisms might result in a tolerance to other yet untested harmful environmental impacts of astrobiological relevance. Therefore, *I. hospitalis*’ radiation tolerance regarding ionizing and non-ionizing radiation has extensively been studied, and the DNA damage visualized on a molecular level. Future experiments will focus on repair kinetics and on the identification of proteins involved in DNA repair.

Methods
The survival after radiation exposure was determined by the most probable number technique (A), and PCR-based assays like RAPD (randomly amplified polymorphic DNA) analysis qualitatively confirmed changes in the genomic DNA after irradiation treatment (B).

**Conclusions**

*In hospitalis* cells showed a remarkable radiation tolerance although never exposed to it in its natural habitat. An exposure to 1000 J/m² monochromatic UV-C (254 nm) resulted in a reduction by 2-3 orders of magnitude which is in the same order as shown by *Deinococcus radiodurans*. An exposure to 24 kGy of ionizing radiation was survived (A); the DNA integrity was only slightly affected (B).

![Survival and RAPD profiles](image)

**A** Survival of *In hospitalis* after exposure to ionizing radiation. Cells were exposed to either γ-rays or X-rays. **B** RAPD profiles of genomic *I. hospitalis* DNA after ionizing radiation exposure.

**References**


Background

Ferroplasma acidarmanus dominated the acid mine draining biofilm community in accordance with metagenomic metabolic modelling (Chen et al., 2012). One of the most impressive features of F. acidarmanus was its ability to survive at such an extreme proton gradient with only a cell membrane. The S-layer protects the cell against mechanical and osmotic stresses of extreme pH conditions. A rigid cell walls or S-layer were not documented neither in F. acidarmanus Fer1 nor T. volcanium as yet.

Objectives

Here we present an evidence for function of a gene coding for S-layer and the cellular localization of the protein in F. acidarmanus Fer1.

Methods


Conclusions

The gene 638394352 was identified at 66-1422 (+) and coded for 452 amino acids with an isoelectric point of 10.08. The S-layer protein possessed a high content of acidic and hydrophobic amino acids. Protein played a structural role for the cell envelope and its biogenesis. We predicted two transmembrane domains of about 20 aa each anchored within the membrane while the majority of amino acids were located extracellular. The gene shared sequence homology with S-layer domain protein from Sulfolobus islandicus. The cellular localization model of 638394352 agreed with the cellular structure and localization of S-layer protein of orthologues Sulfolobus, which formed extracellular monomolecular crystalline mushroom-like
arrays of proteinaceous subunits on the cellular surface.

The peptide statistics, cellular localization, and protein homology, indicated that the gene was coding for S-layer domain.
THERMAL ADAPTATION OF ESTERASES THROUGH THE EVOLUTION OF LOOP REGIONS

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Background
Temperature exerts multiple effects on the structure and function of enzymes which are still poorly understood at the molecular level. The adaptation of microorganisms to extreme temperatures includes the evolutionary development of biocatalysts with a high catalytic efficiency under these conditions. Nowadays, such extremophilic enzymes are used as valuable tools to study relationships between protein stability, dynamics, and function.

Objectives
Here, we report on a comparative biochemical and molecular dynamics analysis of four homologous esterases belonging to the hormone sensitive lipase family which were isolated from bacteria living at temperatures ranging from 10°C to 70°C.

Methods
The recombinant esterases showed highly similar substrate specificities. Furthermore, their optimal temperatures and thermostabilities resembled the temperatures of the respective bacterial habitats. Apparently, the thermal properties of these esterases were optimized to sustain the temperatures of their respective habitats. Therefore, we have used this set of enzymes for the analysis of molecular determinants beyond thermal adaptation.

Conclusions
The structural analysis revealed significant differences between structures arising from surface exposed loop regions. The high flexibility of these structural domains was confirmed by molecular dynamics simulations performed at temperatures ranging from 15°C to 90°C. The results led us to propose four highly flexible loops (L2, L4, L10 and L12) distant from the active site as the structural regions most relevant for thermal stability. The link of such surface exposed loop-structures to thermal stability may indicate that natural thermal adaptation is achieved by tuning atomic interactions in “non-active site” loops without interfering with the catalytic function of an enzyme.
Background
Acid bioleaching environments are characterized by high levels of iron and heavy metals, which can increase the generation of reactive oxygen species and induce oxidative stress in microorganisms. Metaproteomic studies on *Leptospirillum* spp., a member of the bioleaching consortium, have shown that it contains high levels of cobalamin biosynthesis proteins. We have found that cobalamin exerts an antioxidant effect in *Leptospirillum* group II strains.

Objectives
The aim of this study was to clarify the possible mechanisms involved in cobalamin effect in *Leptospirillum* group II strain CF-1 through transcriptomic and proteomic approaches.

Methods
The sequencing and annotation of the genome were carried out using PacBio RS II technology and RAST software, respectively. Transcriptomic profile was obtained by RNA-seq using Illumina Hiseq 2500 sequencing. The mRNA abundance was calculated using Blast and expressed as RPKM (reads per kilobase per million) data. Proteomic assays were carried out using 1D-PAGE LC-MS/MS, and the LFQ intensities were normalized by MaxQuant software.

Conclusions
The genome analysis showed one circular chromosome with 2,709,324 bp and 3,070 coding sequences. Transcriptomic and proteomic analyses showed that cobalamin produces up-regulation of pathways involved in biosynthesis and repair of proteins, biosynthesis of iron sulfur clusters and amino acids, biofilm formation and energy metabolism proteins. Most of up-regulated pathways use S-adenosylmethionine or folate as cofactors which are products of cobalamin-dependent methionine and folate cycles. These data lead us to suggest that cobalamin could activate the central metabolism, improving the response to oxidative damage and restoring redox balance of the cell.
Acknowledgement: Fondecyt Grants 1120746, 1110203.
THE FE-S CLUSTER ASSEMBLY (ISC) SYSTEM IS FUNCTIONAL UNDER OXIDATIVE STRESS AND IRON STARVATION CONDITIONS IN THE ACIDOPHILIC BIOLEACHING BACTERIUM LEPTOSPIRILLUM GROUP II CF1

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Background
The synthesis and assembly of Fe-S clusters is a highly complex and coordinated process in living cells. The ISC system is the housekeeping [Fe–S] cluster assembly system whereas the SUF system is functional under harsh environmental conditions such as oxidative stress and iron starvation. Members of Leptospirillum genus are acidophilic iron-oxidizing bacteria that belong to the consortium of microorganisms that participate in the bioleaching of ores and metal recovery. Interesting, despite to inhabit in a highly oxidizing environment with high concentrations of metals, these microorganisms have only one [Fe-S] cluster assembly system, ISC

Objectives
In this work we evaluated the functionality of the [Fe-S] cluster biogenesis system ISC of Leptospirillum Group II CF-1.

Methods
Bioinformatic analysis and RT-PCR experiments were used to characterize the genetic clusters that contain isc genes and to evaluate the co-transcription of genes. The expression level of key genes iscR, iscS and hscB was determined by quantitative RT-PCR analysis when cells were exposed to oxidative stress and iron starvation. The functionality of the ISC system was evaluated measuring the activity of cysteine desulfurase enzyme (IscS).

Conclusions
The results showed that Leptospirillum Group II CF-1 responds to oxidative stress and iron starvation by up-regulating the [Fe–S] cluster assembly isc genes and increasing the activity of IscS.

Acknowledgement: Fondecyt Grants 1120746, 1110203.
MOLECULAR INVESTIGATION OF THE RADIATION RESISTANT CYANOBACTERIUM ARTHROSPIRA SP. PCC8005

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Background

The cyanobacterium Arthrospira sp. PCC 8005 has been selected by the European Space Agency (ESA) for producing oxygen and food during future long-duration manned space missions, as part of the bio-regenerative life support system ‘MELiSSA’. PCC 8005 must continue to produce oxygen and conserves high nutritive value while exposed to cosmic radiation in space.

Objectives

The tolerance and the response of Arthrospira sp. PCC 8005 to ionizing radiation were investigated.

Methods

Metabolically active planktonic cells of were exposed to Cobalt-60-generated gamma radiation and Helium and Iron particle radiation. The molecular response to radiation was investigated via photosynthesis, pigment, antioxidant, proteome and transcriptome analysis.

Conclusions

The cells of PCC 8005 were able to survive doses of 6400 Gy of gamma, and 1000 and 2000 Gy of He and Fe particle radiation. During irradiation, cells switched quickly from an active growth state to a growth arrest mode, via a shut-down of photosynthesis and carbon fixation. Resources were rerouted to cellular protection and repair. Various antioxidant systems were activated, such as glutathione, to protect lipids, proteins and DNA. The cells activated also ssDNA repair systems and systems to remove damaged amino and nucleic acids from the cells. During recovery, the cells induced the arh genes, a new cluster of genes with unknown function. Finally, energy and metabolic pathways were restarted, and full recover of photosynthetic proliferation was obtained. These results show that Arthrospira sp.
PCC 8005 is a peculiar radiation resistant bacterium, useful for biotechnological applications, in space and on Earth.
Background

Carbon dioxide (CO₂), methane (CH₄) and nitrous oxide (N₂O) are three major greenhouse gases (GHG) which are naturally recycled mainly by microbiological mechanisms including respiration- CO₂ fixation, nitrification -denitrification and methanogenesis-methanotrophy. Microbial groups involved in GHG recycling have been detected through molecular surveys in high altitude wetlands such as Salar de Huasco. This site is a high altitude wetland characterized by its extreme physical-chemical environmental conditions (high radiation, daily temperature shifts, salinity gradients) and by the presence of novel groups within Bacteria and Archaea domains and complex microbial structures (mats) in freshwater and salt-saturated sites.

Objectives

The aim of this study was to determine atmospheric and dissolved GHG concentration in Salar de Huasco wetland and dry Altiplanic adjacent areas (3,800 - 4,000 masl) and the potential communities involved in its recycling during the austral dry season (November 2014).

Methods

GHG were measured in discrete samples from air, water and gas bubbles below microbial mats inhabiting fresh water areas of Salar de Huasco wetland by standard chromatography. Also molecular analyses of the communities were studied using massive sequencing (metagenomic and 16S rRNA barcode).

Conclusions

This study indicates that Salar de Huasco microbial communities process GHG very fast since despite notorious accumulation of CH₄ bubbles (over N₂O) in cyanobacterial-microbial mats from fresh-water areas were found, the wetland was a net sink of GHG based on undersaturated values in the water and low ground air GHG levels compared with dry adjacent areas.
Acknowledgment: This work is part of the FONDECYT Project #1140356 and # 1140179.
HALOPHILIC FILAMENTOUS FUNGI FROM HYPERSALINE AND POLYHALINE ENVIRONMENTS

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Background

The existence of halophilic fungi in hypersaline environments has now been well established, negating the former belief that their presence was as simply ubiquitous fungi, or as being merely halotolerant. Several reports have indeed demonstrated the presence of these fungi in the Dead Sea and in solar salterns.

Objectives

Although hypersaline environments have been recorded as the primary econiche of halophilic fungi, the objective herein, was to ascertain the presence of this extremophilic group of fungi in polyhaline environments of the estuary and mangroves, in addition to solar salterns, and to characterise their nature of halophily.

Methods

Samples of water and sediment were obtained from these econiches and plated on to medium supplemented with high concentration of solar salt. The purified isolates so obtained were screened for their halotolerance levels, by growth on media containing increasing concentrations of salt.

Conclusions

Halophilic fungi were indeed isolated from polyhaline environments, as well as from hypersaline solar salterns. Most of the halophilic fungi screened were found to possess moderate halophily; some of the isolates were true halophiles, having an essential requirement of salt for growth. It is thus shown that this group of extremophilic fungi may be found in hypersaline as well as non-hypersaline marine environments.
DEVELOPMENT OF BIOLOGICAL SOIL CRUST ON THE BARE SOIL MOUNDS AFTER SHRUB DEATH AND DECOMPOSITION IN A SEMI-ARID ECOSYSTEM

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Background

Background: By the end of 2008, and following a prolonged drought period, a mass death of shrubs was observed in a semi-arid region of the Northern Negev Desert.

Objectives

Objective: This study followed the development of biological soil crusts (BSCs) on the soil mounds in the sites of the dead shrubs, spanning over a three year period, from the disappearance of the shrub skeletons until the mounds were flattened.

Methods

Methods: In addition to on-site physical measurements, BSC samples were collected and analyzed for their physical and biochemical properties (15-17 parameters) and their microbial community structure and compared to those of mature crusts.

Conclusions

Conclusions: Statistical analyses showed significant differences in the biochemical properties and the microbial community structure between the developing BSCs and the mature ones during five sampling campaigns. Interestingly, significant differences were observed also between the north-facing and the south-facing slopes of the mounds. The extent of these observed differences was linked to the sampling dates, implying an effect of the soil moisture. Significant differences persisted even after the mounds were totally flattened. This slow development of BSCs on the bare soil mounds after shrub death and decomposition in semi-arid ecosystems, may allow the establishment of new shrubs in these sites and the recovery of the affected ecosystem.
GDGT LIPID COMPOSITION OF THE THERMOPHILIC ARCHAEA

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Background
The archaeal lipids differ significantly from the bacterial and eukaryotic analogs. The most studied lipids of thermophilic archaea are ether-bonded glycerides, resistant to usual lipid derivatization methods. Thus, characterization of archaeal lipid profiles is often not provided with the description of novel species.

Objectives
In our work, we have analyzed the composition of polar lipids, cellular fatty acids and the GDGT lipids of the several thermophilic archaea (Thermoplasma sp. DSM-1728T, Desulfurococcus sp. Z-1312T, and Fervidicoccus sp. Kam-940T), among them are several recently described species.

Methods
Polar lipids were studied by two-dimensional thin-layer chromatography [1,2], GDGT lipids were analyzed by GC-MS after iodination with subsequent reduction by Zn/CH₃COOH [3]. Additionally, native GDGT lipids were analyzed by ESI-MS, and the polar groups of GDGT lipids were sylilated and analyzed by GC-MS.

Conclusions
The polar groups of GDGT lipids of DSM-1728T contained phosphate, mannose, arabinose and gulose. This is confirmed by glycolipids and glycoprophospholipids revealed by TLC.
The GDGT lipids of Thermoplasma acidophilum DSM-1728T are dominated by GDGT4, which is confirmed by GC-MS analysis of core hydrocarbons. The polar lipid profiles of archaea studied are strikingly different, however, there are common glycoprophospholipids for the DSM-1728T, Z-1312T, and Kam-940T strains. The further study of lipids of these archaea by LC-MS will clarify their structures.
This work is supported by Russian Foundation for Basic Research (grant #13-04-40328) and by President of Russia (grant MK-4530.2015.4).

References
USING A FUNCTION-BASED SCREEN TO IDENTIFY ACTIVE RUBISCOS FROM HYDROTHERMAL VENT ENVIRONMENTS: UNRAVELING REGULATION AND ACTIVATION OF A RUBISCO ENCODED ON A METAGENOMIC FRAGMENT

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Background

RubisCO (Ribulose-1.5-bisphosphate Carboxylase/Oxygenase) is a key enzyme in the Calvin Benson Bassham cycle, which accounts for most of Earth’s primary production. Although extensive work on RubisCO enzymes has been conducted in the last decades, we still know relatively little about the regulation and activation of RubisCO enzymes.

Objectives

We have developed a new activity-based screen that enables the identification of recombinant RubisCO active enzymes from metagenomic fosmid libraries. Using this screen we successfully harvested RubisCOs from a metagenomic fosmid library that was constructed from a deep-sea hydrothermal vent. After verifying the RubisCO activity, sequencing of one of the metagenomic fragments revealed a 35 kb DNA insert which encoded a RubisCO form I and form II.

Methods

We investigated the role of adjacent DNA regions on the RubisCO form I and form II regulation and activation by individually deleting each of the flanking genes and measuring the mutant’s activity. Additional experiments with the genes’ transcripts and band shift assays, to assess binding properties of the respective enzymes and genes, have furthered our understanding of how the RubisCO is regulated and likely activated.

Conclusions

This screen and this approach to work with the metagenomic fragments opens the door to directly studying RubisCOs from any environmental sample.
UV-RESISTANCE PROFILING OF A POLY-EXTREMOPHILIC GAMMAPROTEOBACTERIUM FROM HIGH-ALTITUDE ANDEAN LAKES

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Background
High-Altitude Andean Lakes (HAAL) at the South American Andes are exposed to high UV irradiation and high content of toxic elements. A wide variety of microorganisms named “extremophiles” were found to inhabit these environments as Acinetobacter sp. Ver31 and it was chosen as model poly-extremophilic HAAL’s microbe as it displayed high resistance to UV-B.

Objectives
The aim of this work is to integrate diverse experimental approaches to profile the UV resistance mechanisms of Ver3, herein called as “UV-resistome”.

Methods
DSMZ collection strains were used for comparison with Ver3. Strains were exposed to different doses of UV-B, the resistance was determined by colony forming unit counting2. Survival, proteomic profiling and biofilm development after the exposure was evaluated by incubating cells under dark (DR) or light (PR) treatments. Genomic data were analyzed identifying components related with the “UV-resistome”.

Conclusions
Our results shown a superior resistance to UV-B radiation of Ver3 than control strains, in all cases the recovery was more efficient after PR, and the genome sequence supported the phenomenological observations. Ver3 highlighted a number of unique genes, such as a novel cryptochrome. An “UV-resistome” was defined, mainly genes related to UV-damage repair3,4,5 on DNA and genes conferring an enhanced capacity for scavenging the reactive molecular species responsible for oxidative damage. The proteomic profiling of UV-challenged cells identified up-regulated proteins such as a specific cytoplasmic catalase, a putative regulator of quorum sensing, biofilm development and down-regulated proteins related to several energy-generating pathways. This is the first report on a genome from a polyextremophilic Acinetobacter strain.
BACTERIAL DIVERSITY OF GEOTHERMAL SPRINGS IN NAGORNO-KARABAKH

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Background
The ability of microbes to thrive in high-temperature environments has prompted researchers to study these microorganisms to better understand their physiological and molecular adaptations and eventual utilization in various biotechnological applications. Natural geothermal springs, including terrestrial hot springs are one of the habitats of thermophilic microbes. On the territory of Nagorno-Karabakh many geothermal springs with different geotectonic origins and different physicochemical properties are found.

Objectives
This research is focused on investigation of the bacterial community structure of previously unexplored geothermal springs of Nagorno-Karabakh, Karvachar (70°C) and Zuar (52°C).

Methods
16S rRNA gene clone libraries were constructed from total community DNA using universal bacterial oligonucleotide primer sets. Sequences were used for phylogenetic assessment of the communities.

Conclusions
Sequence analysis of clones indicated that clones obtained from Karvachar samples originated from phyla Proteobacteria (48.6%), Cyanobacteria (29.7%), Bacteroidetes (5.4%), Chloroflexi (5.4%), Verrucomicrobia (2.7%) and Planctomycetes (2.7%) and clones obtained from Zuar samples originated from phyla Proteobacteria (42.3%), Firmicutes (19.2%), Bacteroidetes (15.4%), Cyanobacteria (3.8%), Tenericutes (3.8%) and yet unclassified phylotypes (15.4% for Zuar and 3% for Karvachar). The majority of the phylotypes detected in the gene libraries shared less than 95% sequence identity with their closest matches in GenBank, indicating a unique community structure of these geothermal springs. These thermal springs can represent a resource for novel thermophilic organisms and biotechnological tools.

The work was supported by the CPEA-2011/10081 grant from the Norwegian Cooperation Programme in Higher Education with Eurasia.
LIFE ON THE FROZEN CONTINENT: DIVERSITY OF RUBISCO, NIFH AND PUFLM GENES IN SOILS AROUND THE PRINCESS ELISABETH STATION, SØR RONDANE MOUNTAINS, ANTARCTICA.

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Background
In Antarctica, photosynthesis by Cyanobacteria is generally thought to be the main primary source of organic carbon for complex microbial communities. Many cyanobacterial species are also able to fix nitrogen. Therefore, they can survive and prosper in almost every habitat, including Antarctica, Earth's most extreme continent. However, several studies of Antarctic microbial communities have shown that Cyanobacteria are not always highly abundant. We explored the hypothesis that other bacteria must take over their role and produce organic matter as well as fix nitrogen, in order to sustain the microbial community. Light is an abundant energy source during the Antarctic summer and some bacteria can use rhodopsin-type pigments to exploit this, whereas aerobic anoxygenic photosynthetic bacteria can use bacteriochlorophyll for photosynthesis.

Objectives
The presence and diversity of non-cyanobacterial prokaryotes that possess one or several of these properties was studied in terrestrial samples gathered in the proximity of the Belgian Princess Elisabeth Station (Sør Rondane Mountains, Queen Maud Land, East-Antarctica).

Methods
RuBisCO, nifH and pulLM genes were investigated by construction of PCR clone libraries and Illumina MiSeq sequencing.

Conclusions
Preliminary results indicate an extensive diversity of the genes coding for these processes in terrestrial Antarctica.
Background
Soil water repellency is a common phenomenon affecting the hydrological responses of many soil and land use types in different climates. This newly recognised ‘extreme’ environment leads to decreased water infiltration, reduced vegetation cover, fertiliser run off and soil erosion. The fundamental (biological) causes of soil repellency and its dynamic behaviour remain poorly understood.

Objectives
This study aimed to apply metaproteomic and high-resolution imaging approaches to model and predict switches between hydrophilic and hydrophobic soil surface responses.

Methods
Extreme, moderate and sub-critical water-repellent UK grassland soils, including Park Grass at Rothamsted Research, were sampled under wettable and repellent conditions. Soils were subjected to new extraction methods for determining the specific hydrophobic and the general metaproteomes and to Atomic Force Microscopy for determination of topological and adhesion properties.

Conclusions
Using our ultrahydrophobic extraction protocol, we identified novel ultra-hydrophobic microbial proteins, which likely play an important role in the development of soil water repellency. Such proteins could be extracted from moderate and extremely hydrophobic soils with medium-low soil moisture levels, but were absent in the comparable wettable soils. In control extractions, hydrophobic proteins were only extracted from wettable soil containing a positive control, demonstrating the specificity of our novel extraction method. Our newly developed metaproteomic method required only up to 0.5g of soil. Initial comparisons revealed differences in protein profiles, suggesting altered microbial ecophysiology in response to development of water repellency. Atomic force microscopy of repellant soils showed the nanoscale coverage of repellency on particles and aggregates by determining surface adhesion properties from force curves.
Background
Metagenomic studies carried out in salterns located in Alicante and Huelva, Spain, revealed that in intermediate salinities ponds (13-21% of salts) inhabits a dominant bacterial group belonging to Gammaproteobacteria, related to Alkalilimnicola, Arhodomonas and Nitrococcus.

Objectives
On the basis of these metagenomic studies we designed different media and growth conditions in order to isolate this bacterium that have shown to be dominant at intermediate salinities. Subsequently, we studied the strategy of osmoregulation of this successful halophilic bacterium.

Methods

Conclusions
We were able to isolate a microbe that has been cultured and described as Spiribacter salinus. Its complete genome has been sequenced and the recruitments against the available metagenomes of hypersaline aquatic habitats revealed its abundance in intermediate salinities, decreasing sharply at saturated and low salinities. S. salinus shows a simplified metabolic versatility, missing the chemolithotrophic and carbon fixation pathways.

We have combined physiological approaches and genomic analysis to derive a comprehensive picture of the molecular and cellular events that allow the adaptation of S. salinus to saline environments. A qualitative and quantitative assessment by natural abundance 13C-NMR spectroscopy and high-performance liquid chromatography (HPLC) analysis, respectively, have shown that S. salinus is able to synthesize ectoine when is exposed to high salinity, showing an essentially linear relationship between ectoine content of the cells and the external salinity. We have also tested representatives of multiple transporter families that have been proved to sense and respond to osmotic stress, from ABC transporter as OpuA, BCCT transporter as OpuD and from the TRAP- transporter family, TeaABC.
Background
Heshang Cave is a dissolution cave occurred in Cambrian dolomite. It is characterized by dark, oligotrophic, well-developed stalagmite and continuous dripping water throughout the year.

Objectives
Due to the great progress on the geochemical record and successful recover of microbial geolipids from stalagmite, it poses an urgent need to investigate microbial changes in terms of their relationship with environmental conditions which will help to decipher the microbial information in stalagmite.

Methods
Here we present a five year monitoring results about the variation of microbial communities in dripping waters in Heshang cave via 16S rRNA clone library construction and Biolog analysis of their carbon utilization. Meanwhile pH, temperature, conductivity, dripping rate, regional air temperature and precipitation were measured and collected.

Conclusions
Seasonal variations of bacterial communities were observed in composition, diversity and carbon utilization patterns. In autumn and winter bacterial community was dominated by Gammaproteobacteria, whereas in summer and spring Betaproteobacteria became dominant in water samples collected. Cluster and redundancy analysis indicated that regional air temperature strongly affect bacterial communities among the environmental factors investigated. Moreover the bacterial biodiversity was observed to increase with the temperature rising. These seasonal variation match well with the seasonal variation of microbial fatty acids in dripping waters in Heshang Cave which was also strongly affected by regional temperature. Our results show that microbial communities can response temperature change via multiple ways and the microbial variation recorded in stalagmite may be indicative of palaeo-temperature changes.
VARIATIONS BETWEEN MICROBIAL COMMUNITIES IN DEEP SUBSURFACE BOOM CLAY PIEZOMETER INSTALLATIONS

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Background

The subsurface Boom Clay layer is investigated as a potential host rock for geological disposal of radioactive waste in Belgium. The HADES underground research facility (EIG Euridice c/o SCK•CEN), located at 230 m depth under the site of SCK•CEN (Mol, Belgium), provides access to this clay layer for in situ geological, geochemical and geomicrobiological testing.

Objectives

In order to predict how microbiology will affect the biogeochemical processes in a disposal scenario, the resident microbial communities in the man-made structures within Boom Clay are being characterised.

Methods

In this study, water samples were collected from Boom Clay via various existing piezometers (diverse in depth, orientation, location, age, materials used). The aim was to assess differences or shared features of the microbial communities residing in piezometer boreholes, and to correlate variations to geochemical analyses.

Conclusions

Along the five piezometers, bacterial communities of the filters within one piezometer seem more similar to each other compared to those in other piezometers, despite a variety of filter materials or Boom Clay layers sampled within one piezometer. It shows that technical installations (such as piezometers) can introduce and promote local variations in the clay environment and the associated bioprocesses.

Further studies of other piezometers and of clay samples are needed, to pinpoint the source bacterial community underlying in situ enrichment, to unravel the mechanism that shapes such microbial community in different repository conditions and to outline the relevance of the (dominant) microbial classes in defining borehole water (and gas) chemistry.
Host manipulation and bacterial survival

PHENOL-SOLUBLE MODULINS ALPHA ALTER THE CELL CYCLE OF EUKARYOTIC CELLS.

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Background

Staphylococcus aureus (SA) is a Gram-positive bacterium responsible for a wide range of infections in humans and animals. We previously demonstrated that SA USA400 MW2 strain induces a G2/M phase delay in human HeLa cells. This phenomenon was associated with the accumulation of the cyclin-dependent kinase Cdk1/cdc2 and with the accumulation of unphosphorylated histone H3. Additionally we showed that the G2 phase was preferential for staphylococcal internalization and intracellular replication (1).

Objectives

The objectives of the study were

-to identify the active substances, which are responsible for the cell cycle alteration,
-to understand, how this alteration hijacks defense functions of the host cells.

Methods

Using size exclusion chromatography of MW2 supernatant, followed by mass spectroscopy analysis and FACS, we identified phenol-soluble modulin alpha (PSMα) peptides as the likely candidates for cell cycle alteration.

Conclusions

Synthetic PSMα1 and PSMα3 caused a G2/M phase transition delay. The implication of PSMα in cell cycle alteration was confirmed by comparison of wild type LAC wt
strain with the isogenic mutant (LACΔpsmA), lacking the psmA operon that encodes PSMα1 to 4, for its internalization efficiency in HeLa cells. The decreased internalization rate of LACΔpsmA suggested a role of PSMα in host cell invasion.

Furthermore, PSMα-induced G2/M-transition delay correlated with a decrease in the defensin genes expression suggesting a diminution of antibacterial functions of epithelial cells (2). Our results open new perspectives for the investigation of the mechanisms of the SA infection.

1 Alekseeva et al. Plos One 2013
2 Deplanche et al. FASEB. J 2015
Host manipulation and bacterial survival

LACTOBACILLUS ACIDOPHILUS1 MAY PREVENT DIARRHEAL DISEASE BY VIBRIO CHOLERAE IN NEWBORN RABBITS
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Background
Acute diarrheal disease-ADD is the second cause of death in the world in children under five years old and particularly in Latin America a good proportion is caused by Vibrio cholerae. The high incidence of ADD produced by V. cholerae is mainly due to antimicrobial resistance, being mainly to get control of cholera with probiotics our scientific interest.

Objectives
To assess the in vitro and in vivo capacity of Lactobacillus acidophilus1 to prevent diarrheal disease caused by V. cholerae 01OGAWA, in newborn rabbits. Our working hypothesis is to evaluate the use of lactic acid bacteria-LAB in the prevention and control of ADD.

Methods
Isolated and identified by biochemical and molecular tests L. acidophilus1, from 35 Colombian children stools. The amylolytic capacity and biomass formation by L. acidophilus1 were standardized previously. With L.acidophilus1, in vitro and in vivo antagonistic activities against the enteropathogen were performed and prevention of diarrheal disease was evaluated by V. cholerae in newborn rabbits.

Conclusions
The isolate was identified as L. acidophilus1, who in an inoculum of 35x10^6 bacteria/ml was able to exert the most antagonistic in vitro effect on V. cholerae. After the Kaplan–Meier estimator, rabbits faced to the pathogen without receiving probiotic had lower survival probability of 0.25 compared to the group of animals challenged to the pathogen and simultaneously fed with probiotic whose probability of survival was 0.95. L. acidophilus1 is considered a probiotic microorganism, able to survive passage through gastrointestinal tract in an animal model and prevent intestinal colonization by V. cholerae in newborn rabbits.
Host manipulation and bacterial survival

HETEROLOGOUS EXPRESSION IN BUDDING YEAST OF DIVERSE BACTERIAL TRANSLOCATED EFFECTORS THAT TARGET COMMON CELLULAR COMPARTMENTS

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Background
Facultative and obligate intracellular pathogenic bacteria subvert host cell functions to create an intracellular environment for their replication and persistence. This is achieved by injection of bacterial proteins into the host cell cytoplasm via specialized type 3 or 4 secretion systems, thus reprogramming cellular pathways. Heterologous expression in the Saccharomyces cerevisiae yeast model is often used to understand the function of translocated effectors.

Objectives
We report the expression in S. cerevisiae of over 30 candidate and cognate effectors from Chlamydia trachomatis, Coxiella burnetii, and Salmonella enterica as N-terminal GFP fusions, aimed to study their subcellular localization as a means to gain insight in their function within eukaryotic cells.

Methods
We present data on co-localization with fluorescent markers and the use of a battery of yeast mutants defective at specific membrane traffic stages to address differential cues recognized by these effectors.

Conclusions
Most effectors showed a ubiquitous cytoplasmic distribution, either diffuse or concentrated in foci. Sometimes, as in Coxiella CaeA, these foci were consistent with accumulates of unfolded or aggregated protein. However, a few particular effectors displayed specific localizations. Notably, Coxiella AnkB was imported into the nucleus and associated to the nucleolus; C. trachomatis CT696, C. burnetii CBU-77 and Salmonella SteA specifically localized to cellular membranes. Particularly, C. trachomatis CT696 decorated the plasma membrane, C. burnetii CBU-77 localized at vacuolar membranes, and Salmonella SteA targeted simultaneously both
membranous systems. The utility of the yeast heterologous system to study the determinants for recognition by bacterial effectors of subcellular eukaryotic compartments will be discussed.
Host manipulation and bacterial survival

C. CANIMORSUS AFFECTS COAGULATION BY REDUCING THE ACTIVITY OF VITAMIN K DEPENDENT CLOTTING FACTORS

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Background

Capnocytophaga canimorsus is a Gram-negative bacterium belonging to the oral flora of dogs. The bacterium causes rare but severe infections in humans that have been bitten or licked. Patients frequently develop severe sepsis with disseminated intravascular coagulation (DIC).

Objectives

Considering that several bacteria interact with coagulation factors or platelets, as well as the frequency of bleeding abnormalities in C. canimorsus infection, we were interested if C. canimorsus interferes with coagulation.

Methods

We incubated normal pooled plasma (NPP) with C. canimorsus strain 5 (Cc5) and monitored thrombin generation by a calibrated automated thrombogram (CAT) assay. Additionally, clotting times of Cc5 treated NPP were measured. Factor deficient plasma was used to assess the activity of individual coagulation factors in Cc5 treated NPP.

Conclusions

Cc5 inhibited thrombin generation and clotting times of NPP incubated with Cc5 were significantly increased. However, pre-treatment of Cc5 with the irreversible serine protease inhibitor AEBSF completely abolished this increase.

We observed a specific impairment of the Vitamin K dependent (VKD) factors, FX, FIX, FII and FVII. We could also show that FX was cleaved by Cc5.

To conclude, our findings suggest a proteolytic mechanism by which Cc5 affects coagulation and which causes degradation of FX.

We are now interested in finding the cleavage site, to see if the other VKD clotting factors are equally degraded and to identify the bacterial protease which mediates the cleavage.

Inhibition of coagulation could promote bacterial dissemination and also aggravate DIC associated bleeding.
FEMS-1087
Host manipulation and bacterial survival

V. VULNIFICUS DETECTED IN THE Spleen LEADS TO FATAL OUTCOME IN A MOUSE ORAL INFECTION MODEL.

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Background

Vibrio vulnificus causes rapid disseminating septicemia by oral infection in infected individuals who have an underlying disease, especially chronic liver diseases. Although the elucidation of specific risk factors for V. vulnificus infection in patients with liver diseases is of urgent importance, no appropriate experimental animal model that mimics the liver diseases in this bacterial infection has been available so far.

Objectives

Discover the risk factors for V. vulnificus infection in liver diseases patients.

Methods

To discover these risk factors, we generated a liver disordered mouse by performing bile duct ligation (BDL). The BDL mice were infected with the V. vulnificus by orogastric route.

Conclusions

Hepatitis developed in the BDL mice, however this did not affect mortality in mice after orogastric administration of V. vulnificus, suggesting that the liver disorders caused by the BDL were not risk factors for V. vulnificus septicemia. When the dead and surviving mice were compared, V. vulnificus could be detected from the spleen only in the dead group. Furthermore, significantly higher numbers of V. vulnificus were detected from the intestines in the dead group than in the surviving group (p<0.001). These findings suggested that proliferation of the challenge inoculum in the intestine was needed for the oral infection with V. vulnificus, and that the elimination of V. vulnificus in the liver and/or spleen plays a critical role in survival of the host.
Host manipulation and bacterial survival

EFFECT OF HYPOBARIC HYPOXIA ON MICROBIAL HOMEOSTASIS AND PATHOLOGICAL CONSEQUENCES IN GUT
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Background
At high altitude (HA), hypobaric hypoxia is a hallmark of environmental stress that faced by millions of people like pilgrims, trekkers, scientists and military personnel and they generally suffer from a group of non-specific gastrointestinal complications like anorexia, dyspepsia, nausea, acidity, haematemesis, peptic ulcers, etc. These diseases are mostly related to the alteration of intestinal microbes and their related phenomenon.

Objectives
This experiment was conducted to explore the relationship between the altered atmospheric pressure with the microbial composition, which is the major functional counterpart of GI tract.

Methods
Albino mice were subjected to hypobaric atmospheric pressures (55kpa) in a chemostat for 15 days and after dissection intestinal luminal contents were collected. The population of indicator bacteria were monitored by selective culture based methods and DGGE. The expression of inflammatory markers of intestinal epithelium were evaluated by qPCR and western blot.

Conclusions
The quantity of facultative anaerobes like Escherichia coli, other strict (Bacteroidetes sp. and Lactobacillus sp.) and obligate (Clostridium perfringens, Peptostreptococcus sp) anaerobes were increased in many folds after exposure to hypoxic environment. The increased level of bacteria and their endotoxins activated p38 MAP kinase and TLR4 pathways which are related to the over expression of inflammatory cytokines and mediators like IL23, IL17F, TNF a, INOS, COX2 and HIF 1a. As a result mucosal layers of intestinal wall was severely inflamed and perforated which also documented by SEM studies. These consequences are greatly related to the prognosis of different gastrointestinal ailments developed during hypobaric hypoxic stress at high altitude.
IRON ACQUISITION MECHANISMS OF THE CYSTIC FIBROSIS (CF) PATHOGEN BURKHOLDERIA CENOCEPACIA AND THE CORRESPONDING HOST RESPONSE

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Background
B. cenocepacia, is a highly antibiotic resistant CF pathogen which produces low molecular weight iron chelators known as siderophores, predominantly ornibactin, to sequester iron in the host. In response to bacterial and fungal infection, mammals produce lipocalins, some of which bind siderophores.

Objectives
To characterise ornibactin production by B. cenocepacia under different conditions and the resulting host lipocalin response to ferrated ornibactin and to investigate alternative iron acquisition mechanisms by this pathogen.

Methods
Ornibactin gene expression detected using real-time PCR and siderophore levels measured using the CAS assay. Utilisation of iron sources assessed using growth assays. Lipocalin 1 (LCN1) was expressed in E. coli using the Champion™ pET 100 vector system.

Conclusions
This study demonstrated that in iron-depleted cultures siderophore production by B. cenocepacia is significantly upregulated by 4 h (P<0.001) with maximal production by 5 h. Furthermore, B. cenocepacia can acquire iron from exogenous sources such as host iron binding proteins including ferritin and hemin, and from the Aspergillus fumigatus xenosiderophores, fusaridine C and triacetylfusaridine C demonstrating a multifaceted iron acquisition strategy. Host response investigations have determined that recombinant LCN1 binds ferric-ornibactin with high affinity and not unferrated ornibactin. Ongoing studies are examining cellular lipocalin responses and the utilisation of additional xenosiderophores. Understanding the iron acquisition strategies and host responses to this pathogen is warranted given the potential to prevent colonisation by this pathogen if iron acquisition can be compromised.
INFECTION OF CAMPYLOBACTER JEUNI REDUCES CFTR MEDIATED CL-SECRETION IN T-84
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Background

Campylobacter jejuni is the common bacteria cause acute gastroenteritis in humans. It has been reported that the C. jejuni-infection was caused by very low infective dose, and it suggested that C. jejuni might have a resistance system against bacterial clearance system in intestinal tract. The mucus layer in the intestinal tract is the first defense system against bacterial infection, and the condition is well maintained by water movement. Cl- transport is closely associated with water efflux on intestine. Cystic fibrosis transmembrane conductance regulator (CFTR) is the major Cl- secretory pathway which is activated by cAMP. Dysfunction of CFTR causes dehydration of mucus and accumulation of bacteria in the intestine.

Objectives

CFTR is tightly related with infection of pathogenic bacteria. However, it is not clear how CFTR related with the C. jejuni-infection. To elucidate the role of CFTR on C. jejuni-infection, we investigated the Cl- secretion in T-84 on infection of C. jejuni.

Methods

Cl- secretion was measured by ¹²⁵I- efflux on T-84 cells. C. jejuni-infection did not change ¹²⁵I- effluxes in steady state. Next, we confirmed forskolin or prostaglandine E2, agonists of cAMP-dependent Cl- secretion, activated ¹²⁵I- effluxes, which inhibited by CFTR inhibitor.

Conclusions

These indicated that C. jejuni-infection suppressed the activation of Cl- secretion dependent on CFTR. There were hypothesized that C. jejuni-infection effected on water movement and the mucus dehydration, which was related with bacterial clearance system in intestinal tract.
Host manipulation and bacterial survival

A STRATEGY OF PATHOGEN STREPTOCOCCUS ESCAPE MAMMALIAN PEPTIDOGLYCAN RECOGNITION PROTEINS KILLING

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Background
Our previous work showed that a new GI-type IVC secretion system location in the GI similar as previous descried an 89K GI-like structure of S. suis named S4GIs is popular in the genus streptococcus.

Objectives
To reveal an unknown mechanism of a novel family of peptidyl-prolyl isomerases (PPIases) named SP1 from S4GIs interact with mammalian peptidoglycan recognition proteins (PGRPs).

Methods
In this study, we show that the function of killing bacteria of innate immunity proteins PGRPs could be blocked by protein SP1. We confirmed intracellular concentration of SP1 from streptococcus depends on secretion channel of S4G and increased expression of SP1 could be regulated by the concentration of PGRPs in environment using ELISA and qRT-PCR.

Conclusions
In conclusion, this study indicated a new escape way of pathogenicity Streptococcus protects against mammalian innate immunity proteins killing.
Background

Biofilms are highly diverse and complex multispecies microbial assemblages with characteristic spatial, i.e. three-dimensional organization. Elucidating function of a biofilm community would require a detailed characterization not only of the spatial distribution of the sessile organisms but also their respective metabolites.

Objectives

Mass spectrometry (MS) of homogenized biofilms (and other methods) have shown the importance of bacterial metabolites in the corrosion process. However, as seen in the profilometry map of a corroded coupon (Figure 1), the corrosion and the biofilm that causes it are not homogeneous. In order to gain a better understanding of the MIC process, we need to develop an instrument that can combine MS and spatial information.
An ambient mass spectrometry imaging (MSI) system was developed specifically for imaging living biofilms on corroding lab or field samples. The Laser Ablation and Solvent Capture by Aerosol (LASCA) instrument uses an IR laser to collect a 200 μm spot from the sample and analyzes it using a high resolution QToF. A single spot is analyzed in less than 10 seconds, allowing for thousands of spots to be analyzed and built into an ion image (Figure 2). The laser can be tuned to remove a specific

Conclusions

We present a system that can create ion images of thick, uneven biofilms in 3 dimensions. We demonstrate the use of LASCA with the correlation of ion images to corrosion damage. Further, method to probe the penetration depth of biocides into a biofilm is presented.
FLAVIN-BINDING FLUORESCENT PROTEINS - ADVANCED IN VIVO ANALYSIS OF BIOLOGICAL PROCESSES WITH HIGH SPATIO-TEMPORAL RESOLUTION

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Background
Genetically encoded fluorescent proteins (FPs), like GFP from the jellyfish *Aequorea victoria*, became one of the most variable and popular in vivo reporters in cell biology, microbiology and biomedicine. They can easily be detected by in vivo fluorescence techniques thus enabling the non-invasive analysis of complex cellular processes. However, beside the large size and relative slow maturation, a major drawback of GFP and its variants is their strict limitation to aerobic biological systems. This is primarily due to the fact that the autocatalytic synthesis of the chromophore depends on molecular oxygen.

Objectives
To conquer these limitations, we developed a new class of small cyan-green fluorescent proteins which can be used in microbes and mammalian cells under aerobic, oxygen-limited as well as anaerobic conditions. These FPs carry flavin mononucleotide (FMN) as chromophore and are thus termed FMN-binding fluorescent proteins (FbFPs).

Methods
To evaluate the applicability of FbFPs, the photophysical properties including the fluorescence spectra and brightness have been characterized and further improved. Furthermore, in vivo studies demonstrated their broad usability as reporter in (facultative) anaerobic bacteria and for bioprocess engineering. In addition, FbFPs were also be used to generate novel genetically encoded biosensors allowing the ratiometric intracellular monitoring of essential environmental parameters including molecular oxygen and pH.

Conclusions
The unique properties of FbFPs render them particularly valuable as in vivo tools for fluorescence imaging, quantitative in vivo analysis of gene regulation, protein
localization and bio-sensing.
Background

Structures of macromolecular complexes are traditionally solved by X-ray crystallography, requiring crystals and bright X-ray sources to obtain resolutions at which side-chain densities can be seen. Last year, a revolution has taken place in the field of single-particle cryo-electron microscopy (EM), resulting in numerous high-resolution un-crystallisable structures. These breakthroughs have been made possible by the conjunction of a new generation of microscopes, detectors, and software.

In single-particle EM, thousands of noisy images of individual biomolecules are computationally combined into one 3D model. Best results are obtained when the sample of interest has been extensively purified and stabilized. During these steps, the biological context disappears, interacting partners dissociate, and unforeseen conformations could be induced.

Objectives

Ideally, single-particle like methods could be applied to biomolecules within their native environment. Unfortunately, almost all eukaryotic cells are too thick to be imaged by EM.

Our lab has set the long term goal to derive structures from biomolecules within their cellular environment. We label complexes with fluorescent tags to localize regions of interest (ROI) within vitrified cells. The coordinates of these ROIs are, together with the sample, transferred under cryogenic conditions into a dual-beam FIB/SEM system to trim the ROIs into ca 150nm thin lamella. Finally, the trimmed sample is cryogenically transferred to a high-end transmission EM for subsequent imaging at different orientations.

Methods

In here, we will report on the first results obtained with our cryo-correlative workflow on mycobacteria.

Conclusions
We discuss challenges and opportunities, the latter including developments such as phase plates and super-resolution microscopy.
AUTOMATED SEGMENTATION OF INDIVIDUAL CELLS AT PHASE CONTRAST MICROSCOPY IMAGES

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Background

Nowadays, computer processing of microscopic images is becoming more common as a part of research in the field of cell science, replacing slow and time-consuming manual processing by human experts. A challenging task is for example to process images of live mammalian cancer cells from a phase contrast microscope.

Objectives

When processing the time-lapse images of live cells, the main goal is to describe the movement and behaviour of individual cells. There are many methods available in the cases when individual cells are separated from each other. However, separating individual cells from colonies or even from cell monolayers is much more difficult.
Methods

We developed a novel algorithm for segmentation of individual cells. First part separate the cells from the background and it is based on the differences in time between consecutive images and a combination of sophisticated thresholding, blurring, and morphological operations. It is fast and precise. The second part of our algorithm separates individual cells in the clusters. It uses the halos between cells (thresholding and modified skeletonization) and fills the missing parts by connecting the hanging branches of the skeleton via Dijkstra algorithm.

Conclusions

We tested the algorithm on images of four cell types acquired by two different microscopes, evaluated the precision of segmentation against manual segmentation performed by a human operator. We created the software which implements our segmentation method. We added the possibility to modify the resulting segmentation. User can modify the result by merging or splitting the cell regions that was found by our algorithm.
STRUCTURAL STUDIES OF LEGIONELLA EFFECTORS

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Background

Legionella pneumophila is a Gram-negative bacterium and is the causative agent of Legionnaires’ disease. The Dot/Icm type IV secretion system (T4SS) is one of the key virulence factors of L. Pneumophila. By transferring effector proteins into the host cell, the bacterium is able to replicate inside human alveolar macrophages by rapidly altering the normal host endocytic pathway after uptake. Currently, about 300 Icm/Dot dependent effectors have been identified in L. pneumophila; however, the biochemical functions for most of these proteins remain unknown.

Objectives

Lpg1496 is an effector protein containing a conserved sequence of the SidE family in its C-terminal region. The SidE family members are present early during intracellular replication and localize to the cytoplasmic face of the replicative phagosome upon translocation. No information on other regions of lpg1496 has been available.

Methods

We are exploring the three-dimensional structure of lpg1496 to probe for possible biochemical functions using structural similarity to proteins or domains of known function.

Conclusions

Here, we identified two novel regions of sequence similarity in lpg1496 and showed that they form independently folded domains. Moreover, we have determined high-resolution crystal structures of these domains. Each domain consists of two α-helices flanked by four β-strands on one side and four β-strands on the other side. Furthermore, we obtained the structure of the conserved C-terminal domain that shows structural similarity to HD domains and possesses phosphodiesterase activity. These studies will lead to a better understanding of the functional role of lpg1496 in Legionella pneumophila, and may contribute to the development of novel therapeutic treatments for Legionnaires’ disease.
Background

The outbreak of mosquito vectors, the incidence of mosquito-borne diseases and the resistance of mosquitoes to conventional pesticides have recently caused a panic to the official authorities in the endemic countries. The bacterium Bacillus thuringiensis (Bt), a safe eco-friend entomopathogenic biocontrol agent, is widely used to complement the chemical control in integrated mosquito control measures.

Objectives

This study was conducted to identify native mosquitocidal Bt isolates to be utilized in the battle against insecticides–resistant mosquito populations.

Methods

Bt isolates have been recovered from different environmental samples collected from different locations throughout Saudi Arabia. Their mosquitocidal activities were tested against Culex pipiens 3rd instar larvae compared to that of the reference B. t. israelensis (Bti-H14) strain. Three isolates were almost as active as Bti-H14, showing LC50 of 4.2-4.8 µg/ml. Seven isolates showed 1.6-5.4 times more toxic than that of Bti-H14 (LC50 of 0.91-3.03 vs 4.88 µg/ml). Histopathological examination of the Bt-63 isolate-treated larvae showed cellular and subcellular alternations in midgut epithelia similar to that caused by Bti-H14.

Conclusions

Data of this study showed that spore-crystal mixtures of 7 native locally isolated Bt were higher mosquito larvicidal compared to that of Bti-H14. Molecular characterization of these potentially active isolates and their toxic parasporal protein crystals are currently being investigated. This study may lead to the identification of mosquitocidal Bt isolate(s) that could contribute to the battle against mosquito
vectors.
EFFECT OF GUT MICROBES ON THE EFFICACY OF BT MAIZE AGAINST LEPIDOPTERAN STEMBORERS

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Background
The evolution of resistance in agricultural pests to Cry proteins threatens the sustainable use of genetically modified Bt crops. The specific mechanisms of resistance are not well understood. According to the gut microbiota theory, extensive cell lysis caused by Cry proteins provide gut microbes access to the hemocoel where they germinate and reproduce, causing septicemia and death of the host.

Objectives
The main objective of this study was to determine whether microbes present in the mid-gut of Busseola fusca influence the efficacy of Cry 1Ab proteins.

Methods
Larvae were collected from maize fields and dissected to excise the mid-gut. Mid-gut contents (intestinal bacteria) were enumerated on general media. Different morphological types were selected to test the antibiotic susceptibility of the bacteria. The most effective bacteriostatic and bactericidal antibiotics were used and the morphological types were exposed to different concentrations of these to visualise the effects of the antibiotics. This consisted of growth curve studies on all the selected bacteria. A mixture of ciprofloxacin, ampicillin and doxycycline (500 µg/ml) was incorporated into an artificial diet. Stemborer larvae were allowed to feed on this for 7 days. These larvae were then placed on Bt maize (MON810) plant material expressing Cry proteins. Larvae actively fed on the plant material.

Conclusions
Results suggest that by placing antibiotic reared larvae on a Bt plant, the absence of the mid-gut microbes contributed to larval survival on Bt maize.
Insect-microbes interactions

GUT MICROBES OF BUSSEOLA FUSCA (LEPIDOPTERA: NOCTUIDAE)
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Background
Genetically modified maize plants are engineered to express insecticidal toxins derived from the bacterium Bacillus thuringiensis. However, field-evolved resistance of African maize stemborers (Busseola fusca) against Bt-maize has developed and spread throughout South Africa. Studies suggested (1) that gut bacteria are required for B. thuringiensis-induced mortality in most Lepidoptera species and (2) that the toxicity of B. thuringiensis depends on microbial community interactions within the gut.

Objectives
This study aimed to identify the diversity of gut-associated bacteria by both culture-dependent and culture-independent approaches.

Methods
A total of 133 bacterial strains were isolated from the midgut of B. fusca larvae collected from 30 different sites. Molecular phylogenetic analyses of 16S rRNA gene sequences revealed bacteria affiliated to Proteobacteria, Actinobacteria, and Firmicutes. Taxonomic distribution of these sequences placed the isolates into 20 different genera. The majority of bacteria identified were belongs to the genera Enterococcus, Klebsiella, and Bacillus. Culture-independent methods involved the denaturing gradient gel electrophoresis fingerprinting (DGGE), and sequence analyses of a 16S rRNA gene from the excised band supported culture recovery results. However, additional bacterial taxa not determined via culture recovery were revealed using this methodology and included members of the genera Chryseobacterium, Lactobacillus Lactococcus, and Spiroplasma.

Conclusions
Some sequences represent hitherto uncharacterized novel organisms that may have not been characterised yet. The Busseola fusca gut represents an intriguing and unexplored niche for analyzing microbial ecology, which will provide opportunities for research involving the impact of diverse and dynamic microbial communities on developing resistance against Bt-maize.
PRESENCE OF WOLBACHIA SP. IN DIFFERENT POPULATIONS OF Ctenocephalides felis: DEMOGRAPHIC HISTORY OR A WOLBACHIA SELECTIVE SWEEP?

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Background

The cat flea, Ctenocephalides felis, is the most common flea species found on cats and dogs worldwide parasitizing a wide range of hosts. It has long been recognized that many arthropods carry passenger micro-organisms. Wolbachia alone infects in excess of 20% of insect species at any point in time. Wolbachia-infected species have been found to harbour lower levels of mtDNA diversity in closely related uninfected species.

Objectives

At the present work we carried out a molecular and phylogeographical study of Ctenocephalides felis and C. canis from different geographical regions (Spain, South Africa and Iran). Furthermore, the presence of Wolbachia sp. and its influence in these populations has been tested.

Methods

Molecular studies were based on the amplification and sequentiation of ribosomal DNA (Internal Transcribed Spacer 2, and 18S gene) and mitochondrial DNA (cytochrome oxidase c-1 gene).

Conclusions

The presence of Wolbachia sp. was detected in the majority of fleas from different localities. The ribosomal data showed a great homology between all the populations of C. felis regardless the geographical origin but significative differences respect to C. canis. Nevertheless, cytochrome oxidase c-1 sequences revealed significative differences in the population of C. felis from South Africa, appearing these sequences with high homology with that of C. canis. The endosymbiont Wolbachia could be responsible for selective sweeps on mtDNA variability within species. We suggest an introgression of mtDNA and Wolbachia between C. felis and C. canis.
ANALYSIS OF THE IMPACT OF IRRADIATION ON SODALIS IN THE TSETSE FLY GLOSSINA MORSITANS MORSITANS

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Background

Tsetse flies are the sole cyclical vectors of the African trypanosomosis that causes sleeping sickness in humans and nagana in animals. Glossina species have established symbiotic relationships with three bacterial species: Wigglesworthia glossinidia, Sodalis glossinidius, and Wolbachia piipientis, mainly known as a reproductive symbiont. Asalivary gland hypertrophy virus that affects reproduction of infected flies has been reported. Vector control remains the most effective approach for sustainable management of African trypanosomosis. The sterile insect technique (SIT) has proven to be such an effective method and it is based on the mass production of the targeted insect, the sterilization by irradiation, and the sequential release of sterile males.

Objectives

Such releases in endemic areas might increase the disease incidence before achieving eradication. Therefore, the development of symbiont-based strategies to produce tsetse strains refractory to trypanosome infection would be ideal for SIT programmes. One approach which is currently under consideration is to modify Sodalis to produce anti-trypanosome factor(s) in the released sterile males.

Methods

We investigated the impact of irradiation on the establishment of Sodalis in the tsetse fly Glossina morsitans morsitans.

Conclusions

The results indicate that irradiating 5-7 days old male flies with 110 Gy does not increase the mutation rate, as assessed in 15 genes, but it does have a negative impact on the replication rate of Sodalis and the salivary gland hypertrophy virus. In contrast the irradiation treatment increased the prevalence of Wolbachia. These data are discussed in the frame of combining a symbiont-based and SIT-based control approaches.
QUALITATIVE AND QUANTITATIVE ANALYSIS OF MICROBIAL DIVERSITY BY PYROSEQUENCING IN TETRANYCHUS URTICAE

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Background

Because of its extreme polyphagous nature, Tetranychus urticae is a good model organism to study the rapid adaptation mechanisms to various kinds of host plant toxins and acaricides. The symbiotic microorganism has been known as a pivotal factor in such chemical adaptation of many invertebrates. In T. urticae, several symbiotic bacteria (Wolbachia, Cardinium, etc) have been identified but research has been limited to the elucidation of their functional roles in cytoplasmic incompatibility.

Objectives

Analysis of overall symbiotic bacterial composition in T. urticae is necessary as a cornerstone for the integrative study on its host adaptation and acaricide resistance development.

Methods

16S rRNA pyrosequencing was conducted for nine T. urticae populations with different acaricide resistance properties (7 green types and 2 red types) with GS Junior Sequencing system (Roche, Branford, CT, USA) and data analysis was performed by CLcommunity program (Chunlab Inc., Seoul, Korea).

Conclusions

The operational taxonomic unit by CD-HIT and TBC methods were estimated as 85.3±22 and 737±220 in average, respectively. Most bacterial species (>99%) were categorized in the order Rickettsiales mainly composed of Wolbachia spp. and Rickettsia spp. except for the acaricide-susceptible UD strain. The UD strain was composed of Wolbachia spp. (43.4%) and Flavobacterium (53.6%). Green- and red-type mites were grouped into different clades in the cluster analysis using the quantitative and qualitative traits, suggesting that the microbial diversity might be related with the phenotypic characters of body color. Further analysis would be
necessary to elucidate the functional roles of endosymbiont bacteria in regulating the intrinsic physiology of *T. urticae*. 
ANALYSIS OF YEAST-LIKE SYMBIOTE DIVERSITY IN THE BROWN PLANTHOPPER (BPH), NILAPARVATA LUGENS STÅL, USING A NOVEL NESTED PCR-DGGE PROTOCOL

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Background
Yeast-like symbionts (YLS) are endosymbionts that are intimately associated with the growth, development, reproduction of their host, the brown planthopper, Nilaparvata lugens (Hemiptera: Delphacidae). However, it is unclear how many species of YLS are found within N. lugens and how they are related to each other.

Objectives
In order to overcome the difficulty in detecting low numbers of YLS in BPH or culturing the YLS in vitro, in this study, we developed a novel strategy, nested PCR–DGGE approach, which was used to analyze the diversity of YLS.

Methods
The nested PCR protocol was developed as follows: firstly, the 18S rDNA gene and 5.8S-ITS gene were amplified using fungal universal primers. Subsequently, these products were used as a template in a second PCR with primers ITS1GC-ITS2, ITS1FGC-ITS2 and NFGC-NR, which was suitable for DGGE. Using this highly specific molecular approach, we found several previously detected fungi: Noda, Pichia guilliermondii, Candida sp., and some previously undetected fungi, such as Saccharomycetales sp., Debaryomyces hansenii and some uncultured fungi.

Conclusions
To the best of our knowledge, this is the first study to analyze the diversity of YLS in BPH by using the PCR-DGGE system. This methodological approach can be used to assess the relationship between variation in planthopper performance and YLS community.
GENETIC BASIS OF CYTOPLASMIC INCOMPATIBILITY CAUSED BY CARDINIUM

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Background

Endosymbionts of arthropods that cause cytoplasmic incompatibility (CI) manipulate host reproduction: bacteria in infected males sabotage the reproduction of uninfected mates, such that the relative fitness of uninfected females is depressed. Cardinium (phylum: Bacteroidetes) is the only bacterial lineage besides Wolbachia known to cause CI.

Objectives

Identify genes involved with CI and host interactions by transcriptional profiling of a CI-inducing Cardinium in male and female parasitoid wasps (Encarsia pergandiella) and genome sequencing of four Cardinium strains causing different phenotypes.

Methods

RNA was isolated from 1-3 day old male and female Encarsia pergandiella and bacterial mRNA was enriched using the Ribo-Zero™ Magnetic Gold Epidemiology Kit. Female and male samples were sequenced with Illumina Hi-Seq. The transcriptome was analyzed by mapping reads against the Cardinium hertigii genome. Cardinium genomes were also sequenced with Illumina HiSeq, assembled using reference-based and de novo assembly strategies and annotated using RAST.

Conclusions

First genome comparisons revealed evidence for distinct differences between Cardinium strains causing different phenotypes such as the absence of the biotin synthesis pathway in parthenogenesis-inducing Cardinium. Currently, the metatranscriptome is analyzed in more detail, focusing on sex-specific gene expression of candidate proteins for CI and eukaryotic cell cycle regulation. A ubiquitin-specific protease and a ubiquitin ligase, genes of the putative anti-feeding prophage, and other genes involved in host cell interaction were highly expressed. We expect that the results of the transcriptome sequencing and comparison of Cardinium genomes causing different phenotypes will reveal first deep insights into the mechanisms of reproductive manipulation in Cardinium.
DIETARY AND PHYLOGENETIC DETERMINANTS OF GUT COMMUNITY STRUCTURE IN HIGHER TERMITES

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Background

The microbial symbionts of termites play critical roles in the digestion of lignocellulose. Unlike wood-feeding lower termites, which are associated with cellulolytic flagellates, higher termites harbor an entirely prokaryotic gut microbiota. Simultaneously, they have considerably extended the range of their diet to lignocellulosic plant litter in various stages of humification. Since the same dietary specializations have evolved in different evolutionary lineages, higher termites offer the unique opportunity to study potential drivers of microbial community structure in the intestinal environment.

Objectives

To assess the influence of host phylogeny and diet on the composition of the termite gut microbiota.

Methods

We analyzed the bacterial microbiota in the hindgut of 19 higher termite species from different feeding guilds using Illumina sequencing of amplified 16S rRNA genes. Sequence reads were taxonomically classified using a curated reference database and subjected to phylogenetic and statistical analysis.

Conclusions

The high similarity in the bacterial gut microbiota among the wood-feeding and humivorous members of different host lineages identified diet as a strong determinant of microbial community structure in higher termites. At higher taxonomic resolution, however, individual bacterial taxa showed a strong specificity for certain host groups, suggesting they are coevolving with their respective hosts. Nevertheless, evidence of co-cladogenesis is scarce and most bacterial lineages may not co-speciate with their respective hosts over a longer evolutionary time. Rather, the observed patterns of
host restriction seem to be enforced by a combined selection by microhabitat and ecological niche, enhanced by a vertical transmission of symbionts facilitated by the social lifestyle.
Δ-ENDOTOXINS OF BACILLUS THURINGIENSIS FORM AMYLOID FIBRILS THERE ARE INVOLVED IN THE FORMATION OF PARASPORAL CRYSTALS

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Background

The δ-endotoxins of the entomopathogenic bacterium Bacillus thuringiensis are multifunctional, so they are not only with highly specificity destroy invertebrate sensitive cells, but have an antimicrobial effect, leading to the destruction of cell walls and cytoplasmic membranes of some microorganisms like activities of others antimicrobial peptides and proteins (1, 2). We discovered for the first time the amyloid fibrils (AF) relating with parasporal crystals of B. thuringiensis.

Objectives

The crystals of B. thuringiensis subspecies: kurstaki, israelensis, amagiensis were the objects of our study.

Methods

Crystals and AF we studied using transmission electron microscopy. To confirm the formation of AF observed their interaction with the dye Congo red in the polarization interference microscope. Molecular mass of proteins was determined by PAGE electrophoresis.

Conclusions

δ-Endotoxins are capable to forming the AF having different thicknesses (like the previously studied AF). The degree of ordering of the AF increased near the crystals. Our research have given us reason to believe that the B. thuringiensis AF are involved in the formation of parasporal crystals. We discuss the ecological role and practical importance of these AF.

TARGETING GUT MICROBES: NOVEL APPROACH FOR INSECT CONTROL

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Background

Several insects are significantly affecting global economy through insect borne diseases in Agricultural and public health sector. Developing resistance towards frontline insecticides is the major challenge in insect control. Metabolic resistance is one of the major forms of resistance mechanisms. Role of insect gut microbes in supporting normal growth and development of insect is well documented. However, the role of insect gut microbes in developing resistance & favoring insect adaptation has not yet been deeply investigated. Targeting insect gut microbes could lead to unforeseen avenues for effective management of harmful insects.

Objectives

To development novel strategy of insect control by targeting insect gut microbes.

Methods

Two model insects are used in the present study malaria vector Anopheles stephensi and cotton pest mealybug Maconellicoccus hirsutus. Insect guts were cured with selected antibiotic treatment. These cured mosquito larvae’s and Mealybug were observed for pesticide susceptibility. M. hirsutus were observed for fecundity, crawler and adult development wax content. While An. stephensi were tested for altered susceptibility against Bacillus thuringiensis israelensis treatment

Conclusions

We observed that gut microbes cured mosquito larvae and Mealy bug became more susceptible to pesticide and environmental factors. Hence we put forward interesting effective vector or pest control strategy by targeting gut microbes.
Background

Plant galls are abnormal vegetative growths that involve the participation of insect, bacteria, fungi and nematodes to induce growth. The role of microbial associations in gall formation are however, unknown.

Objectives

Our objective was to describe and compare the culturable diversity of bacteria and fungi from galled and non-galled apical branches in Haplopappus foliosus, an endemic flowering plant of the Asteraceae from Chile.

Methods

Fungal and bacterial isolates were grown in Sabouraud and enriched nutrient broth respectively. Molecular identification of the isolates was performed by analysis of 16SrRNA, 18SrRNA and ITS DNA genes analysis. Bioactive fungal secretions were analysed by HPLC-MS.

Conclusions

Our results indicate that there was a significant difference in microbial communities associated with cecidia and non-galled tissue, comprising 72% and 28% respectively. According to the Margalef index (10.25 galls and 1.95 non-galled) there was a high presence of fungi in galls 52% (n=31), bacteria 43% (n=26) and yeast 5% (n=3). Interestingly, some bacterial isolates have been described as PGPR, such as Massilia sp. and Bacillus simplex, and cellulose degrading Cellulomonas denverensis. Fungal isolates, have been characterized by their capacity to secrete bioactive compounds and we are currently using analytical techniques to determine potential function of these secretions in the formation of the gall association.
Knowing the microbial composition in galled and non-galled tissue will allow us to identify potential key roles in the formation of cecidia structures and develop future biotechnological strategies to prevent gall formation in endemic plants of the region.
Background
There are organisms of different complexity, which can survive complete water loss conditions. Among them are bacterial and fungal spores, plant seeds, nematodes, rotifers, tardigrada. The most complex known organism with ability to withstand severe dehydration is larvae of African chironomid P. vanderplanki. During water depletion, larvae cells pump out water and accumulate trehalose and a number of protective proteins – Lea, Hsp, thioredoxins, etc.

Objectives
Here we attempted to investigate the bacterial community inhabiting the gut of sleeping chironomid Polypedilum vanderplanki, an insect able to survive complete dehydration by induction of special metabolic state anhydrobiosis.

Methods
To assess the variability of gut microbiota in P. vanderplanki larvae, we performed 454 pyrosequencing of 16S rRNA gene.

Conclusions
Comparison of larvae maintained under controlled laboratory conditions for more than 10 years and freshly collected from wild allowed to identify the core microbiota. Interestingly, representatives of larvae core microbiota were absent in soil samples suggesting true associations with host. By production of germ free larvae we showed that the absence of bacteria does not affect the rate of successful anhydrobiosys. However, sterile animals are suffering from the fungal infestation. We revealed that some isolated bacterial strains have antagonistic activity against fungi. Moreover, we found that during rehydration millions of virus-like particles are accumulating in the interspace between gut epithelium and perithrofic membrane. These viruses could promote the development of anhydrobiosys phenomenon in P. vanderplanki insect. The work is performed according to the Russian Government Program of Competitive Growth of Kazan Federal University.
FEMS-1412
Insect-microbes interactions

METABOLIC ACTIVITIES OF STRICTLY AND FACULTATIVELY ANAEROBIC GUT BACTERIA AND THEIR INTERACTIONS IN A GERM-FREE COCKROACH MODEL
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Background
The diverse microbial communities in the guts of termites and cockroaches represent a complex metabolic network of individual populations. In view of the special microenvironmental conditions within the gut and possible interactions among the microbiota, it is essential to investigate in how far the metabolic properties of pure cultures reflect their activities in their natural environment.

Objectives
To develop a protocol for the inoculation of germ-free cockroaches with defined cultures of autochthonous gut bacteria and to compare the metabolic profiles of the isolates in pure culture (in vitro) with their activities and interactions in the gut environment (in situ).

Methods
Pure cultures of gut bacteria were isolated from the cockroach Shelfordella lateralis. After inoculation of germ-free cockroaches, the strains were quantitated and localized in situ using qPCR, GFP-fluorescence and FISH. Gut conditions and metabolites were analyzed using microsensors, HPLC and GC.

Conclusions
The germ-free cockroach model provides first insights into the factors affecting the metabolism of the gut microbiota in their native environment. The strictly anaerobic Fusobacterium sp. (strain FuSL) and the facultatively anaerobic enterobacterium (strain EbSL) exclusively colonized the hindgut. Both strains showed high cell densities in mono-association, but the abundance of strain FuSL was much lower when co-inoculated with strain EbSL. Oxygen strongly influenced the metabolic products both under in vitro and in situ conditions. The availability of oxygen in the gut would also explain why the anaerobic strain FuSL is outcompeted by the
facultatively anaerobic strain EbSL, which should achieve higher growth yields under microoxic conditions at the gut wall.
EVALUATION OF ANTIBACTERIAL EFFECT OF AMERICAN COCKROACH HEMOLYMPH ON SOME NOSOCOMIAL PATHOGENIC BACTERIA

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Background

Insects due to their long evolutionary history are able to recognize many pathogenic microorganisms and defend against them. Today, due to the development of resistance to synthetic antibiotics, researchers are trying to apply insect immune-derived products.

Objectives

This study was planned to investigate the antibacterial effect of the American cockroach (Periplaneta Americana) hemolymph on susceptible and resistant strains of nosocomial bacteria.

Methods

Adult cockroaches were reared in insectarium (25 ± 2 °C and 60% humidity) . To stimulate their immune system, 20\textsuperscript{µl} of 10\textsuperscript{6} Escherichia coli cells were injected into coelom . After 6 hours, induced hemolymph were collected . The antibacterial effect of extracted hemolymph were assayed in the agar diffusion method on susceptible and resistant bacteria to antibiotics of ceftazidime, imipenem and, methicillin including, susceptible E. coli strain "ATCC 25922" & "PTCCIBRC-M 10708", Staphylococcus aureus "ATCC25923", Pseudomonas aeruginosa "ATCC 27853" and, resistant strains of E.coli, Pseudomonas aeruginosa and, Staphylococcus aureus that were isolated from the hospital environment and stored in bank of bacteria. Also, non-induced hemolymph effect was evaluated on the same bacteria.

Conclusions

Evaluation of induced hemolymph effect on types of strains showed that induced hemolymph affected about 75% of susceptible bacteria strains (P-Value< 0.001), whereas it had no effect on resistant strains. Among bacterial tested, ceftazidime-sensitive E. coli (PTCC), ceftazidime-sensitive E. coli (ATCC 25922), and methicillin-sensitive S. aureus (ATCC 25923) showed sensitivity to induced hemolymph (P-Value< 0.001).
These results showed that stimulation of immunity system American cockroach leading to production of antibacterial proteins and peptides which have inhibitory effect on bacterial depending on the bacterial strains and their sensitivity.
THE ACTIVATION OF THE AUTOPHAGIC PROCESS BY EIEC PROVIDES A SMALLER SPREAD IN THE HOST CELLS.

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Background

Autophagy has been described as an intrinsic host defense system for recognizing and eliminating intracellular-invading bacterial pathogens. However, invading pathogens have evolved mechanisms to avoid autophagic detection. Enteroinvasive Escherichia coli (EIEC), an important diarrheagenic Escherichia coli, are closely related to Shigella, showing remarkable phenotypic and genotypic similarities. However, the disease induced by EIEC is generally less severe than that induced by Shigella spp.

Objectives

EIEC express much less icsB than S. flexneri, our hypothesis is that EIEC are being efficiently recognized and eliminated by the host cell autophagic process. Moreover, we examined the role of icsB in this process.

Methods

We generate an icsB EIEC mutant by pGEM-T easy Vector System, pJP5603 plasmid and DH5αλpir system. The EIEC∆icsB was confirmed by RT-PCR. HeLa cells were challenged with wild type EIEC and EIEC∆icsB for different time points. The delipidation of LC3B was analyzed by Western Blot. Bafilomycin was used as a positive autophagy control.

Conclusions

Differently from Shigella, EIEC induce autophagy in HeLa cells and this process seems to be independently of IcsB. Our results suggest that other virulence factors than IcsB are involved in the autophagic process induced by EIEC and that IcsB seems to not be a mechanism of EIEC camouflage against autophagic recognition. Additionally, the activated autophagic process could be involved with the slightest spread of EIEC in the host cell, providing a better control of infection.
SECRETORY PROSTATE APOPTOSIS RESPONSE-4 MIGHT REGULATE INTRACELLULAR SURVIVAL OF MYCOBACTERIA

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Background

Regulation of apoptosis has become one of the hot topics in mycobacterial infection. Recently, prostate apoptosis response-4 (Par-4), a tumor suppressor protein, is known to cause apoptosis in prostate cancer cells. Par-4 also participates in apoptotic pathway via interaction with glucose regulated protein 78 (GRP 78), which is activated by endoplasmic reticulum (ER) stress under stressful situation. Previously, we reported that ER stress-induced apoptosis plays a critical role as a host defense mechanism against Mycobacterium tuberculosis (Mtb). In this study, we investigated the role of Par-4 during mycobacterial infection.

Objectives

The aim of this study was to investigate the effect of Par-4 expression on mycobacterial apoptosis in macrophages.

Methods

We showed that mycobacterial infection induced expression of Par-4 and GRP78. Additionally in this study, strong interaction between Par-4 and GRP78 was shown at Mtb-infected macrophage surface through quantitative colocalization analysis. To determine if apoptosis could be initiated by GRP78 and Par-4 complex, we used siRNA for Par-4 before Mtb infection. Annexin V staining was used to identify apoptotic cell death. As expected, siPar-4 decreased apoptotic cell death induced by mycobacterial infection.

Conclusions

We found that Mtb infection activated Par-4 expression as well as ER stress induction. The interaction of extracellular Par-4 and cell surface GRP78 led to apoptosis via activation of caspase-8/caspase-3 pathway. The intracellular survival of
Mtb H37Ra was increased in Par-4 siRNA treated macrophages. These data suggested that Par-4 plays a crucial role in the intracellular survival of mycobacteria.
INITIAL MICROBIAL COMMUNITY INFLUENCES GUT MICROBIOTA
COMMUNITY STRUCTURE EVEN AFTER LONG-TERM DIETARY
INTERVENTION
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Background

Short and long term dietary interventions influences the structure and activity of gut bacterial communities. Recent literature suggests that the impact of dietary changes depends on microbial community structure at the time dietary changes are initiated.

Objectives

Investigate the effects of long-term dietary fibre intervention on the gut microbiota of rats from the same and different litters.

Methods

Forty-five, 21-day old female rats from six different litters were fed one of three diets: 5% fibre, 26% fibre (insoluble-cellulose) and a diet consisting of 50% of a basal diet and 50% cooked red kidney beans (rich in soluble-fibre). Each animal in a litter was randomly assigned to one of the three diets. Gut microbiota composition was determined before and after 14-weeks of treatment. Microbial communities were characterized using 16s amplicon sequencing. Taxonomic affiliations were determined using Mothur and SILVA bacteria database.

At the start of the experiment the microbial communities present in a rat varied with a rat’s litter membership. After 14 weeks on the experimental diets an animal's microbial community composition significantly varied with diet, but significant litter membership effects remained. After diets treatments fibre, short-chain fatty acid, nitrogen and carbon composition of the faeces depended on an animals litter membership and diet.

Conclusions

The results suggest that efforts to enhance the health of humans and other animals through the use of prebiotics may have limited success in general due to among individual differences in the composition of their microbiotas and differences in how these microbiotas respond to dietary manipulation.
Background

In a recently published paper (Schnorr et al., 2014), we provided the first characterization of the fecal microbiota for a community of human hunter-gatherers, the Hadza of Tanzania. Our data suggested a new mutualistic layout for the gut microbiome (GM), with an enrichment in opportunistic bacteria and a depletion in health-promoting Bifidobacterium.

Objectives

To obtain a deeper understanding of the GM-host mutualism in the Hadza, we explored the enterocyte-associated microbiome (EAM), whose layout of the EAM can inform about the functional influence on the host epithelium and other cell types in the mucosa, with critical implications for the maintenance of immune homeostasis. Microorganisms that directly interact with the enterocyte surface have a primary role in the microbiota-host cross-talk.

Methods

By means of a previously developed non-invasive ex-vivo minimal model, based on mucus-secreting HT29 cells (Centanni et al., 2013; 2014), we characterized the EAM of 21 Hadza compared to 9 urban living Italians by 16S rRNA barcoded sequencing, and inferred the functional profiles via PICRUSt.

Conclusions

Compared to Italians, the Hadza EAM were characterized by a greater amount of adhesive and opportunistic microorganisms, such as Enterobacteriaceae and Pseudomonadaceae, resulting in a functional enrichment in cell motility, signal transduction, interaction and biofilm formation. Our results depict an interesting mutualistic configuration of the intestinal mucosal microbiome in Hadza that is...
capable of enhanced microbiota-host cross-talk and interaction at the mucosal surface. These findings stress the importance of a deep microbe-host interaction at the intestinal mucosal interface along the course of human evolution.
FEMS-1634
(Human) microbiome

EFFECTS OF LINSEED AND HEMP SEEDS DIET SUPPLEMENTATION ON CAPRINE RUMEN BACTERIAL DIVERSITY.
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Background

The rumen, an heterogeneous microbial community including bacteria, archaea, protozoa and fungi, is a natural habitat that has evolved into an efficient system for lignocellulose degradation. Thanks to this microbiota, ruminants convert the plant materials into digestible compounds, such as volatile fatty acids, their main energetic fuels, and bacterial proteins. The equilibrium of ruminal microbiota is dependent on the diet, which carries fermentation substrates, and the efficiency of ruminal microbiota can be strongly affected by dietary changes.

Objectives

The objective of this study was to assess the effects of linseed or hemp seeds diet supplementation on bacterial diversity in the rumen content of goats using the Next Generation Sequencing.

Methods

In nine pluriparous Alpine goats fed the same pre-treatment diet for 40 days, ruminal fluid samples were collected, before feeding, using an aesophageal polyethylene probe. After this pre-treatment period the goats were arranged to three dietary treatment groups consisting of control diet (C), control diet supplemented with linseed (L) or hemp (H) seeds. Ninety days later, the same ruminal sample collection procedure was performed. The bacterial DNA was extracted using a protocol described in literature and 16S rRNA gene amplicons on V3-V4 region analyzed by Miseq (Illumina).
Conclusions

In the three dietary treatment groups, bacterial community was dominated by Bacteroidetes and Firmicutes with a high abundance of Prevotellaceae, Porphyromonadaceae and Veillonellaceae and a low presence of Ruminococcaceae and Lachnospiraceae. However, L treatment seemed to affect the bacterial population, reducing the microbial diversity.
Background

Many microbial infections involve formation of biofilms containing multiple species. Besides the difficult treatment, the taxonomic and functional characterization of the microbial communities is a major challenge and classic culture-dependent methods often fail to fully characterize the infection.

Objectives

Our aim is the development of Next-Generation Sequencing based methods combining analysis of 16S-amplicons with metagenomics to reveal both the taxonomic and functional diversity of infection related microbial communities with a focus on virulence and presence of resistance determinants.

Methods

We analyzed samples from chronic lung infections and acute infections of dental implants (peri-implantitis). Taxonomical composition was determined by sequencing barcoded amplicons of the 16S rRNA V1-V2 variable regions using Illumina technology. Sequences were compared with the Ribosome Database Project for the characterization of taxa. Selected samples were also used to isolate whole genomic DNA for metagenomic sequencing using Illumina technology. Sequences were assembled and annotated by bioinformatic gene prediction and alignment to the NCBI non-redundant nucleotide database.
The taxonomic composition of the samples both from chronic lung infections and peri-
implantitis revealed complex communities in both settings. The dominant taxa
included pathogenic taxa like *Pseudomonas, Staphylococcus* and *Streptococcus* in
the cystic fibrosis lung infections and several Bacteroidetes like *Tannerella* and
*Porphyromonas*, which are commonly described in such infections. Analysis of
Metagenomic sequences is ongoing to reveal the functional content of the
communities with focus on virulence factors and resistance determinants. Employing
metagenomics and amplicon sequencing combines taxonomic accuracy and cost-
efficiency with a deep insight in the functional genomic content of microbial
communities.
ANALYSIS OF MICROBIAL DNA SIGNATURES IN THE RODENT NARES AND OLFACTORY BULBS.
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Background

The nose is rich in bacterial species that produce outer membrane vesicles (OMV) known to participate in inter- and intra-species signalling. These structures, however, are highly reminiscent of eukaryotic exosomes and synthetic liposomes: all of these nanostructures involve lipid bilayers containing biomolecules such as DNA, RNA and/or protein. The blood-brain barrier has long scuppered efforts of drug delivery to the brain. Encouragingly, studies have revealed that the brain can be accessed across the olfactory nasal epithelium with biotherapeutics and nanoparticles.

Objectives

The main aim was to investigate the hypothesis that 1) the bacterial material is present in the brain of healthy rodents without causing disease and that 2) the possible entry route is via the nasal cavity across the olfactory epithelium. In addition, we analysed the microbial DNA signatures at the phylum and class level.

Methods

We first attempted PCR amplification of the V3 hypervariable region of bacterial rRNA genes from the mice and rats brain DNA. Produced amplicons were the subjects of 16S phylogenetic analysis by Ion Torrent Next-Generation Sequencing and RDP analysis.

Conclusions

The study proved existence of microbiomes in healthy mammalian brains. The comparison of the microbial signatures indicated that at least some of the bacterial DNA is common at both tissue loci, in support of the precept that nasal bacteria or their products might enter the brain through the olfactory epithelium. In addition, there is some similarity of the bacterial species between mice and rats, though some differences were down to host-species intricacies.
Background
Protein energy malnutrition is a potentially fatal body depletion disorder. The catastrophic effects of malnutrition include diarrhoea, malabsorption, increased intestinal permeability and alleviated immune response, thereby aggravating other pathological conditions and contribute to the global disease burden. Since effective treatment for malnutrition is lacking, therefore, metagenomic procedures allow to access the complex cross-talk between the gut and its microbial flora and understand how a different community composition affects various states of human health.

Objectives
The aim of the study is to develop protein energy malnourished (PEM) mice model and to evaluate the various changes in the body and organ weight, histological changes and the total protein content as compared to control. Moreover, a metagenomic approach has been employed for analysing the differences between gut microbial communities obtained from malnourished and apparently healthy mice.

Methods
Twelve mice were assigned to two groups (6 each/group) and fed either a malnourished (protein=2%, energy=295.2 calories) or a normal (protein=16%, energy=397.2 calories) diet. Body weight was taken till four weeks after which they were sacrificed. Weight of the visceral organs weight and their histological parameters were analysed. Faecal samples were further analysed for the differences in microbial communities between the two groups through RT-PCR. Serum samples stored under refrigerated conditions (-20ºC) were also analysed for total protein content between both the groups.

Conclusions
The present study reveals the development of PEM model by analysing various morphological, anatomical and histological parameters. Moreover, it characterizes the microbial community resident in the gut of malnourished mice as compared to control.
Background
As in mammals, studies on the intestinal microbiota of fish have revealed that the bacterial community may influence growth, feed conversion, epithelial development, and immunity of the host.

Objectives
This study aims at applying 454 pyrosequencing approach to study the autochthonous and allochthonous-associated microbiota from snow trout.

Methods
In total, 32 genomic DNA samples were obtained and then Roche 454 pyrosequencing was applied to the pooled set of 16S rRNA gene amplicons. Sequence data were prefiltered and analyzed with the Quantitative Insights Into Microbial Ecology (QIIME) v 1.8.0 pipeline using default parameters.

Conclusions
From our analyses, distinct microbial communities were detected between the snow trout gut-associated and intestinal content-associated microbiota, as has also been observed in human, pig, mice and alligators. According to the results Firmicutes phylum members composed the prominent phyla of the mucosa-associated microbiota regardless of the fish gender. However, in luminal-associated microbiota Fusobacteria phylum group made up the most dominant group in intestinal content of male and female fish. This suggests that the abundance and diversity of bacterial populations in gut mucus is, in general, quite different from the microbiota in gut contents, indicating that some microbial species poorly colonise gut mucosa layer. Many of these bacteria might be of high physiological relevance for snow trout as these groups have been implicated in vitamin production, nitrogen cycling and carbohydrate fermentation.
DETECTION OF SAPOVIRUS GV.2 BY THE NEXT GENERATION SEQUENCER IN THE STOOL SPECIMENS OF PATIENTS OF GASTROENTERITIS OUTBREAK FROM WHICH PATHOGEN HAD NOT BEEN IDENTIFIED

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Background

Causative pathogens have not been identified by ordinal examinations in some sporadic and outbreak cases of gastroenteritis received at Toyama Institute of Health. In contrast, some pathogens have been identified by metagenomic analysis using the next-generation sequencer (NGS).

Objectives

In this study, pathogens were searched by NGS in one outbreak of gastroenteritis from which pathogen has not been identified by conventional PCR analysis.

Methods

RNA-seq libraries were constructed from six specimens collected in the outbreak occurred in 3 facilities of health center for the elderly in April 2013, and were analyzed by MiSeq sequencer (Illumina).

Conclusions

About 20,000-50,000 reads were obtained in each specimen by NGS analysis, and 2 to 6 reads of nucleotide sequence of sapovirus (SaV) were identified from 3 specimens. These reads agreed with the part of the sequence of sapovirus Hu/Nagoya/NGY-1/2012/JPN (SaV GV.2) reported firstly in the food-borne gastroenteritis (Nagoya City in 2012) with 93-99% identity. Since multiple sequences of SaV were detected from the same specimen, the virus was suspected to be causative agent. Specific primers for SaV GV.2 were generated and PCR was performed. Products were obtained in all six samples by PCR, and their sequences
were revealed to be 99.8 % identity with that of SaV GV.2. Therefore, SaV GV.2 was concluded to be the causative agent of the present case. Pathogens that are not able to be detected by conventional primes because of low homology may be identified by metagenomics analysis that is independent on specific primers as in this case.
A 16S RNA ANALYSIS OF SMALL INTESTINAL BACTERIA COMMUNITIES IN ADULTS WITH/WITHOUT ACTIVE CELIAC DISEASE


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Background
Celiac disease (CD) is a chronic inflammatory disorder of the small intestine that presents only in a fraction of genetically predisposed individuals following gluten consumption. Recently, several studies have suggested that abnormalities in the small intestinal microbiota might be involved in the development or the pathogenesis of CD.

Objectives
The objective of this study was to characterize the specific composition of the duodenal microbiota of untreated CD patients and non-CD controls without mucosa damage in order to identify possible differences in bacterial communities in the upper small intestine.

Methods
Intestinal bacterial communities were identified by pyrosequencing of 16S ribosomal DNA extracted from duodenal biopsies. The study group included 9 untreated CD patients and 9 non-CD controls without duodenal mucosal atrophy.

Conclusions
Analysis of a total of 180,825 reads of the 16S rDNA showed that the majority of them were classified within two phyla: Firmicutes and Proteobacteria. Bacterial richness and diversity was higher in non-CD controls than in untreated CD patients. The principal coordinates analysis revealed that bacterial communities of non-CD controls and untreated CD patients were dispersed without forming a clear group according to the presence or absence of the CD. There is no characteristic pattern of bacterial communities associated to untreated CD patients. There are not differences in any bacterial group community frequencies in the upper small intestine between untreated CD patients and non-CD controls.
Background

Prostatitis is one of the most common urological diseases in men and accompanied by bacterial infection in the prostate gland which occurs in 5-10% of prostatitis patients and also, frequently causes recurrent urinary tract infection. Urogenital health and diseases rely on the interaction between the host and the urogenital microbial community.

Pyrosequencing a next generation sequencing (NGS) technologies allow us to characterize microbial communities using bacterial 16S rRNA sequences with high magnitude faster than previous Sanger dideoxy chain-termination method. Therefore, the aim of this study was to identify specific microbial descents that may play crucial roles in the development of prostatitis and to examine whether the presence or abundance of these bacterial changes in semen of prostatitis patients.

Material and Methods

Semen samples were collected from prostatitis patients and extracted genomic DNA for pyrosequencing by using DNA extraction kit.

Results and Discussion

In this study, some abundant pathogenic genus, such as Anoxybacillus, Staphylococcus, Corynebacterium, Pseudogluconobacter and Enterococcus in the prostatitis sample were found by using the pyrosequencing techniques.

In this study, in case of normal control, Streptococcus and Corynebacterium were abundance to 33.5% and 21.2%. On the other hand, the most abundant genera in semen of prostatitis patient were Enterococcus (20.7%), Pseudogluconobacter (16.2%) and Anoxybacillus (14.3%). These data provided the comprehensive report on bacterial communities between normal and prostatitis patients and also, it is concluded that prostatitis is related to substantial alteration in the composition and representation of the urogenital bacterial biota.
This identified variability in community membership and structure has important implications for understanding the etiology of prostatitis, for developing diagnostic tools and for treatment.
Background
Dandruff is a global common problem, up to 50% of both genders suffer from dandruff. However, although bacteria and fungi associated with dandruff are known, understanding the etiology of dandruff is incomplete.

Objectives
Correlation of the bacterial and fungal communities with dandruff is needed because the human skin is associated with many microorganisms with potential for both beneficial and detrimental interactions.

Methods
For all cases, samples from normal scalps and severe dandruff patients were collected and analyzed under protocols approved by Chung-Ang University College of Medicine IRB (Protocol #2012-02-01). From the collected samples, we amplified variable regions V1-V3 of the 16S rRNA gene (bacteria) and the D1/D2 region of the 26S rRNA gene (fungi) using universal primers and GS-FLX pyrosequencing.

Conclusions
Firmicutes were the most dominant in dandruff patients and showed an increase from 5.2% in cases defined as normal scalps to 91.0% in severe dandruff patients. Whereas, Proteobacteria decreased from 60.1% in the normal scalp to 6.2% in dandruff patients. Similarly fungi of the phylum Basidiomycota were associated with the normal scalp, decreasing from 66.6% to 24.7% in severe dandruff. In the Firmicutes Staphylococcus spp. were dominant in the dandruff scalp. In contrast, Malassezia spp. in the Basidiomycota were the most abundant fungus in the healthy scalp. This switch from Malassezia to Staphylococcus provides new comparative information on the microbiomes of the normal scalp and patients with exacerbated dandruff. Consequently, our results can be expected to lead to improved diagnosis and treatment.
GENOMIC ANALYSIS OF NOVEL CHITINOLYTIC SUBSPECIES OF CLOSTRIDIUM SARTAGOFORME FROM BUFFALO RUMEN REVEALS ITS IMPORTANT ROLE IN VOLATILE FATTY ACID PRODUCTION

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Background
The rumen of herbivores is an open fermentation chamber for digestion of complex plant biomass by highly efficient bacteria.

Objectives
Isolation of anaerobic rumen microbes from buffalo and its pheno-genotypic analysis. Correlate the genomic analysis with metagenome data for implication of the isolate’s role in rumen.

Methods
A strain of Clostridium sartagoforme species was isolated from buffalo rumen, involved in cellulose and chitin degradation. The isolate was cultured between 37°C and 42°C temperature with an optimum pH of 7.5 in Hungate’s medium supplemented with cellulose. Whole genome sequencing was performed using ion torrent PGM. Taxonomy study using local BLAST, showed highest phylogenetic relation with Clostridium sartagoforme DSM 1292 strain based on comparison of the 16S rDNA sequences. Comparison of sequences coding for housekeeping genes phosphoglycerate kinase and gyrase B confirmed its taxonomic novelty within the Clostridium genus. Biochemical assessment of strain’s fermentation ability using AN-Biolog plate revealed that the strain utilized more than 30 different metabolite precursors. Pathway analysis make known the organism to be highly active in the production of volatile fatty acids (VFAs).

Conclusions
The isolate represents a new subspecies, for which the name is proposed Clostridium sartagoforme AAU1 (GenBank Accession:ASRV00000000). Genomic analysis gives an insight about the role of bacteria in determining ruminant host physiology. VFAs produced by the isolate play a prominent role in animal health maintenance by contributing a major proportion of the ruminants’ daily energy requirement. A deeper understanding of genes involved and enzymes encoded allow further industrial applications and modulation of the rumen for enhanced farming applications.
METAGENOMIC ANALYSES REVEAL A CELLULOLYTIC AND FERMENTATIVE LIFESTYLE IN THE GUTS OF EPULOPISCIMUM-REPLETRE SURGEONFISHES

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Background

Epulopiscium fishelsoni, is a giant bacterium found exclusively in the guts of surgeonfishes. Three years after the publication of an incomplete draft genome of Epulopiscium sp. type B from Naso tonganus, we still have no clue into the metabolic potential of this hitherto uncultivated symbiont, and more so, the functions of the gut microbiota in reef fishes.

Objectives

Hence we were prompted to fill this knowledge gap, and thus provide a foundation for understanding the physiology and ecological roles of their microbiota.

Methods

We therefore employed a metagenome-sequencing approach to explore the phylogenetic diversity of Epulopiscium, and through binning of assembled metagenomic contigs, also shed insights into their metabolism.

Conclusions

Our data shows that distinctive feeding guilds of surgeonfishes harbour different Epulopiscium-like phylotypes, and that the low-to-high intestinal abundances of these bacteria correlate with their algal diet types. Metabolic reconstruction of the draft Epulopiscium-like genomes from these fishes revealed a capacity to catalyze hemicellulolysis and an anaerobic fermentation lifestyle that produces acetate and lactate. Our data thus provides the first in-depth analysis into their diversity and putative symbiotic role, which in turn illuminates the potential metabolic links shaping the symbiosis between the host and the giant bacteria. We further present more insights from the draft genomes, focusing on the nature of this symbiosis, and its interconnection to the trophic ecology of the fishes.
Background

Aromatic compounds comprise one-quarter of the Earth’s biomass and are the second most widely distributed class of organic-compounds in nature, next to carbohydrates. In tropical-countries, the buffalo are fed on lignocellulosic agricultural by-products like cereal-straws and tree-foliage. Ruminants digest such plant-materials by microbial-processes. The role of microbiota as reservoir of its unique functional capacity related to the metabolism of aromatic-compound needs to be explored in buffalo rumen.

Objectives

In the view of above concern, objective of the present study was to account the comparatives profiling of phylogenetic and functional-potential of genes related with the metabolism of aromatic-compound in Bubalus bubalis rumen fed with different diets.

Methods

The metagenomic analysis of liquid and solid-fraction of the rumen-biomaterial collected from Bubalus bubalis was carried out using high-throughput sequencing. The experimental animals received M1 diet for 6 weeks followed by M2 for 6 weeks and
then M3 for subsequent 6 weeks. On the last day of each experimental feeding period, rumen-samples were collected 3h post feeding using stomach tube.

Conclusions

The work presented here describes composition of overall functional-capacity related with the genes for metabolism of aromatic-compound and responsible taxonomic-communities of buffalo rumen-ecosystem. The higher % abundance was found to be related with genes involve in peripheral-pathway for catabolism of aromatic-compound in all treatment-groups. The taxonomic analysis of metagenomic reads indicated that rumen-microbiomes were dominated by Bacteroidetes and Firmicutes phyla in the all treatments-groups. The information obtained from this research will open new-horizons towards a full understanding of functional-genes related with metabolism of aromatic-compound and metabolic-capabilities of complex-compound degrading rumen-microorganisms.
INTAKE OF A LICHEN-BASED DIET ALTERS THE POPULATION OF METHANGENIC ARCHAEA IN THE RUMEN AND HINDGUT OF REINDEER

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Background
Reindeer are large Holarctic herbivores, which rely on a consortium of anaerobic microorganisms in the rumen and hindgut to digest plant cell wall carbohydrates. Reindeer may include a large fraction of lichens in their winter diet. Lichens are high in phenolic secondary compounds, which may affect the reindeer gut microbiome, and possibly alter the output of methane. Methane is produced by methanogenic Archaea mainly from CO₂ and hydrogen.

Objectives
In this project we examined the effect of lichens on the reindeer rumen and hindgut methanogenic Archaea.

Methods
Rumen and cecum samples were collected from two groups of Norwegian reindeer (Rangifer tarandus tarandus), one group (n=4) fed lichens and one fed a standard pelleted reindeer feed (n=3). The diversity of methanogens was assessed using 454 pyrosequencing. Real-time PCR experiments were performed to estimate the population densities. All animals were sacrificed in accordance with Norwegian regulations, Sections 9 and 10 in the Norwegian Animal Welfare Act.

Conclusions
Densities of methanogens and protozoa were higher in the cecum of lichen-fed reindeers but no differences were found in rumen samples. Methanobrevibacter (≥ 97% similarity) was the major genus of Archaea in all the samples, with M. thaueri as the dominant species in all reindeers fed lichens (>60% on average). Remarkably, M. ruminantium suffered a significant increase (up to 23.8%) with a lichen-based diet in both sampling sites. In conclusion, the intake of lichens affected to some extent the microbial diversity and density of the microbiome housed in the rumen and cecum of reindeer.
Background

Drinking water quality is a public health concern worldwide. Growing evidences depict drinking water as a complex matrix, in which a wide diversity of microorganisms interact in a dynamic network. Recent studies reveal that drinking water treatment process can affect the microbiome structure. Moreover the occurrence of antibiotic resistance genes in water is becoming an issue of great interest as the mobile resistome can easily spread among species. Molecular techniques can give a deeper knowledge, going beyond the limit of culture-dependent methods.

Objectives

In this study we evaluated and standardized a new pipeline for microorganisms concentration, DNA extraction and amplification, suitable for molecular analysis and optimized for High-Throughput Sequencing (HTS) approaches.

Methods

We collected samples in a water treatment plant in Milan (Italy), at different steps of the potabilization processes, from the aquifer to the tap. We analyzed the presence and the relative abundance of bacteria and eukaryotic microorganisms across the water treatment plant. Furthermore the presence of specific antibiotic resistance genes was detected and quantified with Real Time PCR, at each step of water treatment process. Since molecular techniques are unable to differentiate between viable and nonviable microorganisms, live/dead ratio was estimated using SYTO9/propidium iodide staining coupled with microscopy visualization. These analyses are integrated in a broader study characterizing microbiome structure variability using HTS techniques, in order to better understand this complex ecosystem.

Conclusions

The results agree with those obtained in the few recent studies published till now and can help to unravel the dynamics underlying water microbiome changes.
Background
Phototrophic and heterotrophic bacteria have been found directly inside halite evaporites that formed as bottom-growth crusts in a hyperarid core and inside gypsum minerals. The role of microorganisms in the precipitation of authigenic minerals like carbonates, sulphides and silicates, has been documented by some authors in several natural systems.

Objectives
In this study the interface between biological and geochemical components in the surface crust of a saline soil was investigated. The bacterial community structure and diversity as well as the possible role of the photosynthetic microbial community in the formation of the soil crust was evaluated.

Methods
A variable pressure scanning electron microscope equipped with energy dispersive X-Ray diffraction analysis was used for geochemical analysis, while pyrosequencing of the V2-V3 16S rRNA gene region was used for characterizing microbial diversity and community structure.

Conclusions
The organization and diversity of the microbial taxa was analyzed according to the physical and chemical heterogeneity of that peculiar habitat. Mineral structures made of gypsum and halite were documented as crystallized around the filaments of cyanobacteria. A univocal causal relationship could not been established between the crystallization of gypsum or halite and cyanobacterial or algal metabolism but their compartmentalization respect to the biological mat suggested that the organisms
could have at least a physical control over the distinctive structures produced. The results provided information on the type of distribution of different bacterial and cyanobacterial groups as a function of the biogeochemical interfaces, suggesting also possible forms of microbial interaction.
Background
Bacteriophages are the most abundant biological entities in the biosphere. They influence the evolution and dynamics of microbial populations. Although phage populations were studied extensively for the marine environment only few studies were dedicated to freshwater ecosystems.

Objectives
The main objective of this project is to study the phage population in Lough Neagh, the largest freshwater lake in the British Isles and to analyse interaction of these phages with bacterial hosts.

Methods
Both total microbial and viral subpopulations were investigated using standard protocols. Isolation of viral fractions was performed by common procedures. Pyrosequencing was used to identify bacterial communities and the dynamics of the bacterial population’s changes in Lough Neagh are compared with seasonal variation. Illumina sequencing was used to characterise viral communities in the same environment.

Conclusions
16S rRNA analysis detected a diverse range of bacterial phylum in both water and sediment. Proteobacteria, Acidobacteria, Bacteroidetes and Planctomycetes are found in the freshwater samples which are comparable to the sediment samples. The water samples taken from Lough Neagh have bacteria phylum which are not present in the sediment samples such as Chlamydiae. Analysis of viral metagenomes detected the presence of 16S rRNA sequences probably originated from transducing particles. For example Pelagibacteraceae was identified and this bacterial family is more commonly found in the oceans rather than a freshwater lake. Possibly this may indicate interactions between bacterial species from two different environments. Illumina analysis of viral metagenomes provided a diverse range of information and can be compared to other published metagenomes.
FEMS-2455
Metagenomics in the environment

OPTIMIZED AND STANDARDIZED SINGLE INDEXING SOLUTION FOR HIGHLY MULTIPLEXED MICROBIAL COMMUNITY PROFILING ON ILLUMINA MISEQ PLATFORM
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Background
The growing need to survey the tremendous microbial diversity in a culture independent manner, has led to the development of molecular methods through sequence profiling of conserved genes, in scientific field like ecology, agronomy, biotechnology, plant, animals or human health. Recently, the improvement of the Illumina MiSeq platform to a 2×300 bases paired-end version made it much more attractive for amplicons sequencing in metagenomics. Nevertheless, the lack of nucleotide diversity at initial sequencing positions due to sequence conservation is problematic on Illumina platforms. Moreover, validated Illumina amplicon sequencing protocols do not support more than 96 samples per run, which underutilized the overall capacity of a sequencing run.

Objectives
We therefore worked on the development of a standardized and optimized high-multiplexed metagenomic solution for the exploration of complex environments.

Methods
We developed an integrated solution combining (i) a custom and standardized amplification protocol for library preparation; (ii) a 192 custom single index validation strategy within a single run; (iii) the inclusion of a nucleotide sequence variability at the first sequencing cycles; (iv) an accurate paired-end reads assembly process and (v) a dedicated pipeline for taxonomic affiliation and microbial communities analysis.

Conclusions
Our high-multiplexing sequencing solution validated within complex metagenomics samples, provided high sequencing accuracy and technical reproducibility with no index biases based on taxonomic distribution analysis. The association of technical and bioinformatical improvements yields substantial cost reductions and provides greater target flexibility to investigate structure as well as functions of microbial communities through 16S rDNA, 18S rDNA, ITS and functional genes profiling.
METAGENOMIC ANALYSIS OF NATURAL MICROBIAL COMMUNITIES IN ELECTRO-ACTIVE MARINE SEDIMENTS

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**Background**

In marine sediments, a novel group of filamentous sulfide-oxidizing bacteria has recently been discovered, which induce electrical currents over centimetre-scale distances. These so-called cable bacteria grow from the oxic zone into the sulfidic part of the sediment up to a depth of several centimetres. Cable bacteria use a radically novel microbial metabolism, in which electron donor and electron acceptor reactions are spatially separated: the oxidation of sulfide in anoxic sediment layers is coupled to the reduction of oxygen at the sediment surface. This mechanism is only possible when electrons liberated from sulfide oxidation are transported from cell to cell along the length of the filament to the oxic sediment surface.

**Objectives**

To identify a marine sediment where cable bacteria are active in situ and to study the sediment microbial community using shotgun metagenomic sequencing.

**Methods**

Depth profiles of oxygen, sulfide and pH were taken with microsensors and cable bacteria distribution with depth was evaluated using fluorescent in situ hybridisation. Metagenomic analysis was performed for different depths of the sediment: the oxic surface sediment, the suboxic zone where neither oxygen nor sulfide was present, and the deeper sulfidic zone without cable bacteria. Extracted sediment DNA was sequenced using Illumina MiSeq with 300 bp paired end reads.

**Conclusions**

At the examined North Sea site, microsensor profiles indicated that metabolic activity of cable bacteria dominated the sediment geochemistry. The obtained metagenomic
data provide insight into the microbial community composition and functioning, and possible interactions between cable bacteria and other taxonomic groups.
INFLUENCE OF INOCULA SLUDGE VOLUME INDEX ON ANAEROBIC GRANULATION

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Background

Granulation is important for successful start-up of high-rate anaerobic reactors and usually takes long time. The influence of inoculum’s Sludge Volume Index (SVI) on granulation in high-rate upflow Hybrid Anaerobic Reactor (HAR) was studied.

Objectives

Understanding of microbial community changes due to variations in SVI will help in providing novel reactor setting up conditions to obtain faster granulation rate in high rate anaerobic reactor.

Methods

Ten HARs were operated with inoculum having different values of SVI (100 to 325), SVI were varied by adding different amount of inoculum (10% to 50% of the reactor volume). The methanogenic and bacterial community structures of anaerobic sludge obtained from these ten HARs were examined using culture-independent techniques like DGGE and qPCR.

Conclusions

Results showed that granulation was achieved in HARs with SVI values ranging from 150-210 or 18% to 27% inoculum of the reactor volume. DGGE identified 17 methanogenic and 22 bacterial operational taxonomic units in the ten reactors’ samples. DGGE analyses showed dominance of Methanosaeta and Methanosarcina-like species in reactors where granules were formed. qPCR analysis also confirmed that granulating reactors were rich in aceticlastic methanogens especially Methanosacetaceae. Bacterial rRNA genes obtained from DGGE results, were affiliated to four Orders - Clostridiales, Lactobacillales, Actinomycetales and Bacteroidales. But, no significant correlation was observed between granulation and bacterial abundance. The concentration of bacterial 16s rRNA genes was found to be less in granulating than non-granulating systems unlike methanogenic genes. Thus, this study suggests to make conditions favorable for growth of aceticlastic methanogens to decrease granulation time.
FEMS-1429
Metagenomics in the environment

MINING SLAUGHTERHOUSE DRAIN BIOFILMS FOR NOVEL ENZYMES AND BIOSURFACTANTS
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Background
Environmental DNA derived from uncultured bacteria has huge potential as a source of novel bioactive molecules as shown by recent screening campaigns of metagenome libraries which revealed a number of novel protein families, extremozymes and secondary metabolites. Before constructing a metagenomic library, the choice of the habitat should be carefully considered because it predefines the types and properties of biocatalysts to be discovered.

Objectives
In this study, we have constructed a metagenomic library in \textit{Escherichia coli} DH10b with DNA isolated from the microbial population of a slaughterhouse drain biofilm. By comparative screening for lipolytic, proteolytic and detergent activities we have identified number of clones producing bioactive molecules.

Methods
By applying a functional screening assay for biosurfactant activity, two haemolytic and surface-active clones (SA343, SA354) were identified. Sequence and structural homology analyses revealed that they contained genes encoding N-acyl amino acid synthases (NAS) located on environmental DNA fragments.

Conclusions
The respective NAS proteins (Nas343 and Nas354) were shown to be responsible for the haemolytic phenotype and the biosurfactant synthesis \textit{in vivo}. The molecular organisation of the NAS domain linked here to a c-di-GMP binding domain has so far not been identified in any other known protein. The produced biosurfactant was purified and identified by NMR spectroscopy as N-myristoyl-tyrosine. Its biophysical properties including the critical micelle concentration and the ability to reduce surface tension were compared with chemically synthesised N-myristoyl-tyrosine and shown to be in a similar range. Interestingly, N-acyltyrosine also showed antibiotic activity against various Gram-negative and Gram-positive bacteria.
MICROORGANISMS INVOLVED IN CARBON AND NITROGEN CYCLING IN THE ARABIAN SEA OXYGEN MINIMUM ZONE

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Background
Oxygen minimum zones (OMZs) play a major role in marine biogeochemical cycling. They are characterized by a high microbial diversity adapted to life along an oxic-anoxic gradient. In particular anammox and/or denitrification are dominant processes in these zones accounting for significant loss of nitrogen from the ocean. However, other processes within the nitrogen and also carbon cycling have not been explored well.

Objectives
To get insights into the organisms involved in carbon and nitrogen cycling in the Arabian Sea OMZ, we studied the meta-genomes of filtered ocean water recovered at two different depths, the upper zone (170m deep, PA2) and the central zone (600m, PA5).

Methods
We systematically screened for marker genes indicative for key nitrogen and carbon cycling processes by blasting the meta-genomes data against manually curated functional gene databases.

Conclusions
Both zones are characterized by a large fraction of alpha, gamma and delta-proteobacteria. In PA5, anammox contributed a significant amount to the overall diversity (about 8%) whereas Thaumarchaea were present in higher numbers in the upper zone (about 9%). The predominance of nitrifying archaea over bacteria could also be confirmed by amoA diversity. No bacterial amoA was detected in the entire dataset, and only very few sequences related nitrogen fixation were recovered. Furthermore, no canonical methane oxidizers could be found. However, pmoA sequences only distantly related to known diversity are present and could be linked to a published single cell genome, indicating the possibility of a cryptic methane cycle in the Arabian sea OMZ.
MICROBIOMES OF THE BUILT ENVIRONMENT ARE ALTERED BY DIFFERENT ROOM MAINTENANCE

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Background
The excessive removal of microbes from controlled built environments like intensive care units, operation theatres and especially cleanrooms is daily routine. However, effects of these stringent maintenance procedures on our surrounding microbiomes in indoor environments are unknown.

Objectives
Since recent studies showed the enormous potential of the human microbiome towards our health, this study aims to resolve the interactive interference of all microbiomes in the built environment and determines its potential for human well-being indoors¹.

Methods
High throughput deep sequencing technologies were applied for shotgun metagenomes and 16S/18S rRNA and ITS gene amplicons of Archaea, Bacteria, Eukaryota and fungi. Viability assays using ATP (adenosine tri-phosphate) and PMA (propidium monoazide) were deployed to identify potential viable microbiomes and supported by qPCR to investigate microbial abundance²,³. The biotechnological potential was further characterized by VOC’s (volatile organic compounds) assays of culturable indoor bacteria⁴.

Conclusions
These assays revealed high reduction of viable microbial abundance in cleanrooms⁵, higher similarities of viable microbiomes between controlled and uncontrolled areas compared to the total microbial fraction, distinct profiles of microbial communities from floors, medical devices and workplaces in an intensive care unit, and for all controlled built environments not only an overlap with the human, but also the plant microbiome. Hence plant leaves could be identified as a main source for microbial distribution on the surrounding indoor environment. The knowledge of certain key species and their ecological key functions in controlled as well as uncontrolled built environments will help to control indoor environments in a more sophisticated way for our health inside buildings in the future.
Background

Presence of heavy metals impedes use of sewage sludge as a fertilizer. Simultaneous sludge digestion and metal leaching (SSDML) combines conventionally distinct stages of metal leaching and sludge stabilisation into one, making the process economically attractive. Insights into the microbial community involved could enable further optimization of the process.

Objectives

To evaluate the performance of SSDML and understand corresponding microbial (bacterial) community shifts.
Figure 1. Bacterial DGGE profile during the process (0, 2, 4, 6, 8, 10, 12 and 15 are the n\textsuperscript{th} day of operation of experiment)
Figure 2. Unweighted pair group method with arithmetic mean (UPGMA) analysis of DGGE profile in Figure 1
**Methods**

Batch study was carried out at 37°C for 15 days after adding sulfur to heavy metal contaminated sludge, with aeration rate of 0.6 vvm. Samples for heavy metals, volatile suspended solids (VSS) analyses and DNA extraction were collected regularly.

**Conclusions**

Bio-oxidation of sulfur to H$_2$SO$_4$ lowered the pH aiding leaching of metals. Zinc, chromium, cadmium and lead reduced by 86.75%, 64.26%, 79.17% and 69.58% in the dry sludge respectively. Concomitantly, VSS reduction of 49.40 % was obtained making the sludge stable.

*Acidithiobacillus ferrooxidans* and *Acidiphilium acidophilum* were found to be the main bioleaching microorganisms through 16S rDNA PCR-Denaturing Gradient Gel Electrophoresis (DGGE) analysis, in contrast to the commonly reported sulfur-oxidising *Acidithiobacillus thiooxidans*. *Enterobacter, Alicyclobacillus cellulosilyticus* and *Pseudomonas* were the dominant heterotrophic organisms present across all samples. UPGMA analysis showed a significant microbial shift during the course of operation which could be mainly attributed to the change in pH and solids content.
Thus, after the single stage treatment of SSDML, sludge obtained was stabilised with reduced heavy metal load. PCR-DGGE analysis showed that both heterotrophs and bioleaching microorganisms were present and adapting well according to the dynamic pH.
FEMS-1771
Metagenomics in the environment

QUALITATIVE AND QUANTITATIVE BIAS ESTIMATION OF MICROBIAL COMMUNITY ANALYSIS USING BARCODED 454 FLEX+ METHODOLOGY FOR CLAY CONTAINING GEOLOGICAL SAMPLES
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Background
High throughput sequencing methodology can be used to determine the composition of environmental microbial communities. Generally, a barcoded PCR amplicon of the omnipresent 16S rDNA gene is used to unravel complex microbial communities. However, several biases and technical difficulties linked to this methodology, may jeopardize the outcome of the community analysis. For instance, it is often difficult to extract DNA out of environmental samples due to interferences caused by the geological matrix. This is especially true for clay containing geological samples. Beside, often containing highly reactive dissolved organic materials, clayey soils are also known for their high sorption capacity of (biological) macromolecules. Therefore, the presence of clay minerals may be responsible for additional biases when analyzing microbial communities of soil or geological samples.

Objectives
To quantify the biases, qualitatively and quantitatively, caused by the presence of clay minerals, during analysis of the composition of microbial communities in geological samples, with high throughput barcoded 454 Flex+ DNA sequencing methodology.

Methods
Four identical bacterial mockup samples were made, by mixing 18 different soil bacteria. Boom clay was only added to two samples. DNA was extracted, with an identical DNA extraction procedure. The 16S rDNA gene was PCR amplified with four different barcoded 16S₃₄₁-forward primers and one 16S₁₄₉₂-reverse primer. The amplicons were sequenced using the 454 Flex+ DNA sequencing methodology. The data were treated with our in house developed bioinformatics pipeline, including the tools, NoDe and CATCh.

Conclusions
At the start of the FEMS-symposium 2015 the results and conclusions will be ready to be presented.
BIOINFORMATICS ALGORITHMS TO GET ACCURATE 16S RRNA HIGH THROUGHPUT SEQUENCING DATA.

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Background
The revolution in new sequencing technologies has led to an explosion of possible applications, including microbial biodiversity studies via bacterial 16S rRNA amplicon sequencing.

Objectives
All 16S rRNA amplicon sequencing data suffer from the presence of erroneous sequences: (i) chimeric sequences introduced by the PCR reaction, and (ii) sequencing errors produced by the sequencing platform itself. As such, there is a need for efficient algorithms to remove those erroneous sequences in order to be able to accurately assess the microbial diversity.

Methods
First, a machine learning method called CATCh (Combining Algorithms to Track Chimeras) is able to integrate the output of existing chimera detection tools into a new more powerful method, outperforming all existing tools. Second, NoDe (Noise Detector) was introduced as an algorithm to correct 454 pyrosequencing errors, showing a significant improvement in reducing the error rate (reduction of 67 to 75%) and dramatic decrease in computational cost compared to state-of-the-art tools. Similarly, IPED (Illumina Paired End Denoiser) was introduced as first tool in the field for the denoising of MiSeq sequencing data.

Conclusions
Integrating both CATCh and NoDe/IPED into the preprocessing pipeline of 16S rRNA amplicon sequencing data significantly increases the reliability of the data, demonstrated by the positive effect on the clustering of the sequencing data in operational taxonomic units (OTUs). When tested on mock communities (MiSeq as well as 454 pyrosequencing platform), the number of OTUs closely approaches the actual number of species present in the mock samples. Both tools are freely available at http://science.sckcen.be/en/Institutes/EHS/MCB/MIC/Bioinformatics/.
MYCOSPHERE EFFECT OF THE PINE MUSHROOM (TRICHOLOMA MATSUTAKE) ON THE DIVERSITY, COMMUNITY STRUCTURE, AND FUNCTIONAL PROFILES OF SOIL MICROBES

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Background
The mycosphere is a distinct soil environment affected by fungal hyphal activities. Because mycosphere affects microbial activity, it is a valuable source of understanding microbial dynamics in soil ecosystem.

Objectives
The main objective of this study is to explore the mycosphere effect of T. matsutake by comparing bacterial and fungal diversity, community structure, and functional profiles in different zones of the fairy ring. We expanded on the design of past studies by comparing multiple fairy rings across two geographical locations.

Methods
We extracted soil DNA from mycosphere samples in fairy ring of T. matsutake and adjacent soil samples. After processing pyrosequencing data, we conducted metagenomic analysis for microbial diversity, community structure, and functional profiles.

Conclusions
Across samples, community was similar within but different between mycosphere and non-mycosphere samples—microbial diversity was lower in the mycosphere compared to non-mycosphere zones. Many ectomycorrhizal fungi were negatively correlated with the presence of T. matsutake. Analysis of bacterial functional profiles predicted distinct bacterial activity between zones; in mycosphere samples, xenobiotic biodegradation and metabolism, and amino acid metabolism were high, but carbohydrate metabolism, and glycan biosynthesis and metabolism were low compared to non-mycosphere samples. Overall, similar functional activity of mycosphere zones across fairy rings and geographic locations imply a mycosphere effect of T. matsutake, by recruiting mutualistic taxa and excluding antagonistic ones based on the functional trait.
Metagenomics in the environment

COMPARATIVE ANALYSIS OF METAGENOMES FROM FILTER UNITS OF FULL-SCALE HYBRID TREATMENT WETLAND

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Background
Treatment wetlands (TW) have become widely used for wastewater treatment over the recent decade for their cost-effective building and maintenance. While physical and chemical pollutant removal mechanisms in TWs are well known, understanding of microbial processes related to pollutant removal in TWs is still low. Metagenomics offers the ability to examine directly the genomic content of microbial communities, complementing taxonomic information with functional capability.

Objectives
(1) to characterize microbial communities in vertical and horizontal sub-surface flow (VSSF and HSSF respectively) filter units of a full-scale TW treating wastewater from a schoolhouse for over ten years; (2) evaluate the functional potential of bacterial and archaeal communities.

Methods
Water samples, including influent, effluent and intermediate well, together with filter media biofilm from both units were used for shotgun sequencing with Illumina MiSeq system. Metagenomes were assembled into contigs and used for microbial profiling, identifying functional genes and relating them to nitrogen, phosphorus and carbon cycle.

Conclusions
Domain distributions showed the dominance of Bacteria (>95%), a small fraction of Eukaryotes (>2%), Archaea (>1%) and Viruses (0.2%). Samples of filter material had similar bacterial dominant phyla, but different distribution of lower taxonomic ranks. Wastewater samples had diverse bacterial community distribution between samples. Over one-third of archaeal sequences belonged to methanogenic species. Nitrogen metabolism in VSSF and HSSF filter units was dominated by ammonia assimilation related genes followed by nitrate and nitrite ammonification, denitrification and nitrogen fixation related genes. Phosphorous metabolism was dominated by phosphate metabolism genes.
Metagenomics aims for sequencing and identification of DNA from communities of microorganisms in their natural environment. By estimating and comparing gene abundances between samples we can get a picture of the functional role of the metagenome, i.e. which genes and pathways are abundant at different conditions.

Methods for functional metagenomics typically rely on identification of functional domains such as PFAMs, TIGRFAMs and COGs. However, this results in a broad classification of gene products since many proteins can be classified into the same functional domain. The human gut gene catalog, for instance, is predicted to contain more than 5 million genes, but the number of PFAM domains is only around 10,000.

We present HierBin, a new method for functional annotation and quantification in metagenomes, giving a more detailed functional description of the metagenome and performing better in identification of changes between conditions.

First a supervised classification step is used to identify known functional domains in the data. Secondly, each domain is divided into subdomains using unsupervised clustering based on sequence similarity. Finally the abundance of each subdomain is quantified by mapping the sequence reads to the subdomain sequence and differentially abundant subdomains are identified.

We show, by evaluating on resampled data, that our method performs better in predicting differential abundant genes than methods using only functional domains for gene prediction and detects differences at higher resolution that would be invisible at lower resolution.

Finally, we apply our method both to metagenomics data from the human gut as well as environmental samples.
METAGENOMIC ANALYSIS OF MICROBIAL COMMUNITIES IN RAPID SAND FILTER TREATING GROUNDWATER. COMMUNITY DIVERSITY AND METABOLIC POTENTIAL

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Background
Rapid sand filtration is a drinking water production technology widely used in Europe and other countries around the world. This technique allows the removal of ammonia, manganese, ferrous iron, methane, sulfides and other compounds from groundwater by a combination of physical, chemical and biological processes. During many years, it has been recognized that microbial activity plays an important role in these processes.

Objectives
Therefore, it is essential to study the microbial composition in these rapid sand filters and how these communities interact and facilitate the removal of pollutants.

Methods
In this study, six samples from a rapid sand filter were analyzed through a metagenomic approach. By building a non-redundant gene catalog we identify the microbial diversity and the metabolic pathways occurring in this system.

Conclusions
Nitrospirae and Proteobacteria were the dominant phyla. The functional genetic potential included most of genes related to nitrogen oxidation (specially abundant in the nitrification pathway) and to methane oxidation. Genes involved in different CO2 fixation pathways were also abundant. Genera typically implicated in manganese oxidation as Hyphomicrobium and in iron oxidation as Gallionella were also abundant in the catalog. This investigation has so far revealed the main biological processes occurring in these ecosystems. In the next step, we will try to assemble genomes of the most abundant genotypes of the rapid sand filter community.
Metagenomics in the environment

INTRODUCING A COMPLETE METAGENOMICS ANALYSIS PIPELINE, BASED ON AN OPTIMAL COMBINATION OF ILLUMINA SEQUENCING TECHNOLOGY AND AN EASY TO USE ONLINE PLATFORM

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Background
Direct sequencing of metagenomes is commonly implied for bacterial community analysis related to food research, biotechnological processes and clinical diagnostics.

Objectives
Commonly the taxonomic classification of sequences is based on unique target regions such as the 16S ribosomal gene which includes a number of unique variable regions. In this work we present an overview of a new and comprehensive Metagenomics analysis pipeline that we are using to prepare, sequence and analyze metagenomics samples from a range of environments.

Methods
In this holistic approach we start with laboratory techniques that take into account and reduce as much as possible the sources of possible bias that can be introduced into metagenomics datasets by laboratory steps like sampling procedures and DNA isolation. We further include both biological and technical controls to continually validate the process.

Conclusions
The bioinformatic analysis is performed using overlapping paired-end reads which are annotated through the Greengenes database (De Santis et al., 2006). Finally we introduce the ‘BaseClear metaGenome Browser’ (BmGB)™, https://metagenomics.baseclear.com, an online platform in which researchers (with or without any bioinformatics training) can easily interpret the taxonomic composition of hundreds of metagenomes using intuitive interfaces and efficient analyses. The workflow is tested on a number datasets and results are discussed.
LIMITS TO ROBUSTNESS, REPRODUCIBILITY AND ECOLOGICAL CONSISTENCY IN THE DEMARCATION OF OPERATIONAL TAXONOMIC UNITS

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Background

The demarcation of Operational Taxonomic Units (OTUs) from complex sequencing datasets is a key step in large-scale characterizations of microbial communities. However, it remains unclear how well the various proposed OTU clustering algorithms approximate ‘true’ microbial taxa, and many conceptually distinct approaches continue to be used.

Objectives

We explored different aspects of OTU clustering, in particular the robustness, reproducibility and ecological consistency of OTU demarcation.

Methods

In a global and comprehensive survey of roughly one million microbial SSU sequences, we objectively quantified biases introduced by several widely employed algorithms.

Conclusions

We observed surprising trends in the robustness of sequence clustering to changing parameters, and with regard to the reproducibility of OTU-based findings. Moreover, we explored the ecological consistency of OTUs – based on the assumption that, like true microbial taxa, they should show measurable habitat preferences (niche conservatism). We systematically parsed sequence annotations to obtain broad ecological descriptions of sampling sites. Based on these, we observed that sequence-based microbial OTUs generally showed high levels of ecological consistency. However, different OTU clustering methods resulted in marked differences in the strength of this signal. We found that hierarchical average and complete linkage clustering provided the most robust and reproducible partitions with regard to a wide range of parameters. Moreover, complete linkage clustering provided the ecologically most consistent clusters. Assuming that ecological consistency can serve as an objective external benchmark for cluster quality, we
concluded that hierarchical complete linkage clustering, should be the default choice for OTU clustering.
Background: Turf algae holobionts (host macroalgae plus microbiome) are widespread in coral reefs worldwide. They are also the most abundant benthic structures in the Abrolhos reef Bank (Brazil), with more than 50% benthic coverage. Abrolhos reefs turfs are morphologically diverse regarding color and texture, but their microbial matrix is not well known. Objectives: Our aim was to perform a metagenomic characterization of the prokaryotic microbiome of the different morphologic types of turf algae holobionts from Abrolhos reefs. Methods: We used metagenomic approaches to obtain the whole communities associated with turfs assemblages, collected at different reefs during March and October 2013, totaling 22 samples. Results: Filamentous, non-heterocystous cyanobacteria were the dominant microbial phototrophs of the holobionts. Main macroalgal in the holobiont were red, green and brown seaweeds. 15,27 million reads were generated using Illumina sequencing. Proteobacteria (36.8%, N = 1,564,678), Cyanobacteria (28.8%, N = 1,222,446), and Bacteroidetes (10.2%, N = 434,897) were dominant among the turf algae prokaryotic microbiome. Major microbial components of the turf algae holobiont are Roseobacter sp., Silicibacter sp., Dinoroseobacter sp., Congregibacter sp., Maribacter sp., and Cyanobacteria. Genes belonging to aerobic anoxygenic photosynthesis (AAnP), sulfur metabolism (e.g. S oxidation, and DMSP consumption) and virulence potential (adhesins, fibronectin/fibrinogen-binding protein, type IV pili, superoxide dismutase) were found in the turf algae microorganisms. Conclusion: The complex assemblies of microbial guilds render turf holobionts important competitive advantages allowing them to outcompete other benthic organisms and dominate vast reef areas.
EXPLORING MICROBIAL DIVERSITY PATTERNS IN THE SØR RONDANE MOUNTAINS (EAST ANTARCTICA) USING NEXT GENERATION SEQUENCING AND ARISA

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Background
Antarctica is a continent of extremes. Low availability of liquid water and nutrients, extreme low temperatures and high levels of radiation exert high selective pressures on organisms. Consequently, most life forms are microbial. Estimations indicate that as little as 1-3 % of the continental surface represents ice-free regions. Relatively few biological studies focus on terrestrial samples which most often originate from the McMurdo dry valleys that make out the largest contiguous ice-free area. Mountain tops protruding through the ice sheets (nunataks) are less frequently studied although they might function as havens and reservoirs for terrestrial organisms. Geological data shows the presence of different kinds of bedrock in the Sør Rondane mountains (Queen Maud Land, East Antarctica). Patches of macroscopic organisms (lichens, mosses and arthropods) are scattered throughout these ice-free islands, indicating a high amount of variability in the presence of organic matter, and hence nutrients for microbial life. Here we present the results of a large scale sampling effort in the eastern Sør Rondane Mountains of such ice-free regions near the Belgian Princess Elisabeth station.

Objectives
We examined the composition and distribution of bacterial communities in these refugia, and investigated the possible impact of environmental parameters on these community compositions.

Methods
Samples were subjected to both a genetic fingerprinting technique (ARISA) and second generation sequencing (Illumina MiSeq 300PE).

Conclusions
Preliminary results indicate an effect of bedrock type and presence of macrobiotic organisms.
Background

Salmonella 4,[5],12:i:- has become a new epidemic serotype in Europe associated with human infections, being the identification/tracking of its clones crucial to contain their spread.

Objectives

We assessed the trends in S. 4,[5],12:i:- clones distribution and their association with antibiotic and metal resistance/tolerance genes in Portuguese isolates. Results were also compared with previous data from the last decade (2002-2010).

Methods

S. 4,[5],12:i:- isolates (n=158), confirmed by PCR (fliB-fliA/fliB), from different sources (clinical/food) and regions of Portugal (2010-2014) were analyzed. They were screened for sulfamethoxazole resistance genes (sul1-sul2-sul3) and other class 1 integrons genes (intI1, qac and gene cassettes) by PCR. Detection of metal tolerance genes [copper (pcoD), silver/copper (silA) and mercury (merA)] and other antibiotic resistance genes by PCR, susceptibility to 10 antibiotics [ampicillin (A), chloramphenicol (C), gentamicin (G), kanamycin, nalidixic acid, ciprofloxacin, streptomycin (S), sulfamethoxazole (Su), tetracycline (T) and trimethoprim (Tr)] (CLSI/EUCAST) and clonality by PFGE were performed in representative isolates.
Conclusions

We detected the presence of the 3 clones currently circulating in Europe: i) "European clone" (75%; sul2 and absence of intI1/sul1/qacEdelta1; mostly ASuT-blaTEM-strA-strB-sul2-tetB and carrying pcoD+silA+merA), which has expanded throughout this study period; ii) "Spanish clone" [6%; intI1; qacEdelta1+qacH; mostly AC(G)SSuTTtr-blaTEM-cmlA-floR-(aac(3)-IV)-aadA-sul1-sul2-sul3-tetA-dfrA12 and carrying merA-silA] and iii) "Southern-European clone" (1%; intI1; qacH; CSSuTTtr-cmlA-aadA-strA-strB-sul3-tetB-dfrA12) mostly with similar MDR and/or PFGE-types described since 2002. A marked decreased frequency of Spanish and Southern-European clones was observed contrasting with the expansion of the European clone characterized by ASuT-phenotype and copper/silver/mercury tolerance genes, which might facilitate adaptation and success of these strains.
BIOAUGMENTATION WITH METAL-RESISTANT PLANT GROWTH-PROMOTING RHIZOBACTERIA IMPROVE THE METAL RHIZOACCUMULATION POTENTIAL OF SPARTINA MARITIMA

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Background
As an adjunct to phytoremediation strategies and as part of an effort to make this technology more efficacious, a number of scientists have begun to explore the possibility of using plants and the microbial populations colonizing the rhizosphere.

Objectives
A glasshouse experiment using soil from a metal-contaminated estuary was designed to investigate the effect of a native bacterial consortium, isolated from Spartina maritima rhizosphere and selected owing to their plant growth promoting (PGP) properties and multiresistance to heavy metals, in plant growth and metal accumulation.

Methods
Plants of S. maritima were randomly assigned to three soil bioaugmentation conditions (without inoculation, one inoculation and repeated inoculations during experimental period) for 30 days. Growth parameters and photosynthetic traits, together with total concentrations of several metals, including As, Cu, Pb and Zn were determined in roots and/or leaves.

Conclusions
Bacterial inoculation improved S. maritima root growth, through a beneficial effect on its photosynthetic rate (AN) due to its positive impact on functionality of PSII and chlorophyll concentration. Also, bacterial inoculation favoured water use efficiency (WUE) of S. maritima, through the increment in AN, stomatal conductance and in root-to-shoot ratio. Moreover, this consortium was able to stimulate plant metal uptake specifically in roots, with increases of up to 19% for As, 65% for Cu, 40% for Pb and 29% for Zn. On the basis of these results, bioa augmentation of S. maritima with the selected bacterial consortium can be claimed to enhance plant adaptation and metal rhizoaccumulation during marsh restoration programs.
ANALYSIS OF ADAPTATION TO HIGH CONCENTRATIONS OF NICKEL IONS OF A NOVEL SPHINGOBIUM STRAIN BY RNA-SEQ

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Background
While bacteria belonging to the Sphingobium genus are generally known for their capacity to degrade poly-aromatic hydrocarbons, their response to high concentrations of metal ions has been largely disregarded. Only S. cupriresistens, a species isolated from a copper mine, has been reported to tolerate high concentrations (0.9 mM) of Cu²⁺ [1]. We have recently reported the whole genome sequence of a novel Sphingobium strain able to tolerate Ni ions up to 20 mM [2].

Objectives
In order to understand the molecular mechanisms by which this strain is able to adapt to high concentrations of Ni ions we are currently investigating its transcriptome profiles by RNA-seq approach.

Methods
Sequencing of rRNA-depleted RNA was carried out for independent triplicate cultures in the presence or absence of 10 mM NiCl₂, using the SureSelect strand-specific RNA library prep kit (Agilent) for Illumina MiSeq multiplex sequencing.

Conclusions
Transcriptomic data show the differential expression (fold change ≥ |2|, p-adjusted < 0.05) of about one-hundred genes. Most of them are up-regulated in the presence of Ni²⁺ and include genes for membrane proteins and metal efflux systems. Gene annotation was further enriched with description of operons and UTRs as identified by the EuGene integrative gene finder tool [3]. The observed metal resistance of this Sphingobium strain highlights its possible use for biodegradation of xenobiotic compounds in metal-rich environments.

ADAPTABILITY OF TRICHODERMA ASPERELLUM TO HIGH CONCENTRATIONS OF VARIOUS METALS
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Background
Trichoderma asperellum is a biological agent capable of removing heavy metals from the environment via biosorption and/or bioaccumulation. Most studies investigated the use of dead fungal biomass for metal biosorption. The potentials of using live cells to transform toxic metals into less harmful forms (bioaccumulation) is lesser known.

Objectives
In this study, we explored the adaptability of \textit{T. asperellum} to increasing concentrations (100-6000 ppm) of various heavy metals (Al, Cd, Cr, Cu, Pb and Zn).

Methods
Adaptive tolerance index (TI) of fungus in Potato Dextrose Agar was measured.

Conclusions
The results conveyed bioaccumulation potential of \textit{T. asperellum}. Results revealed that \textit{T. asperellum} has adaptability towards increasing concentrations of Cd (TI: 0.01-0.04), Cr (TI: 0.22-0.28) and Cu (TI: 0.24-0.31) at the beginning of the experiment. Decreasing TI values (from 1.08 to 0.01) with increasing Al, Pb and Zn concentrations indicated poor adaptability. The most tolerable concentration across all metals, except Cu, was 1000 ppm. In fact, \textit{T. asperellum} tolerated up to 6000 ppm Al. Concentrations of 1500-2000 ppm generally exerted inhibitory effects with Cu being most toxic (tolerable up to 300 ppm). The adaptability of \textit{T. asperellum} to high metal concentrations was evident with SEM micrographs showing hyphae growth, in spite of some structural deformities, metal precipitates and particles deposited on the hyphal surface. Nevertheless, in cases where growth was inhibited, the severely deformed hyphae displayed irregular, grooves and rough texture. Our study highlights the potential of live \textit{T. asperellum} cells in removing heavy metals from the environment in a cheap and environmental-friendly manner.
RECOVERY OF LITHIUM FOR NATIVE MICROORGANISMS FROM SALAR DE ATACAMA BRINES, CHILE.

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Background
Salar de Atacama (Chile) is the largest basin in Atacama Desert (2900 Km²) and is located at 2300 m.a.s.l. This hypersaline system possess an average concentration of lithium as brine of 1500 ppm. Lithium is obtained industrially through the process of solar evaporation ponds, reflecting the conditions present in this Salar, including high UV radiation, high evaporation and low rainfall; however, this process is non-selective and is pollutant. We propose the recovery of lithium from brines based on biosorption using Gram-positive bacteria selectively accumulate lithium

Objectives
The aim of this study was to describe microbial diversity present in brines of Salar de Atacama and to isolate native bacteria with lithium adsorbent potential.

Methods
Natural brines and concentrated lithium brines of Salar de Atacama were characterized. Such brines present hostile conditions for life, including high salt concentrations (0-100 g/L of Na⁺; 190- 450 g/L of Cl⁻; 0.2 -8 % of Li) and pH between 0.9 and 7. Simultaneously, microorganisms from the Salar were grown in a range of mineral mediums with or without organic carbon, and different concentrations of LiCl (0- 300 uM) through which the ability to absorb lithium was determined.

Conclusions
Bacteria and Archaea were detected in both brines. These microorganisms are likely naturally adapted to the differences in concentration of the elements present, especially lithium. Furthermore, bacteria capable of adsorbing lithium after 12 hours were obtained. Analyses of microbial diversity of the brines and development of a process for obtaining lithium by native microorganisms are currently underway.
MODELING ARSENITE OXIDATION BY SINORHIZOBIUM SP. M14 IN CHANGING ENVIRONMENTAL CONDITIONS

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Background
A complement to the chemical methods of treatment of water contaminated with arsenic are the biological methods i.e. biooxidation. Among the biological methods, technologies using bacteria, which oxidize arsenites, appear to have significant potential. In the literature, many bacterial strains able to oxidize arsenates are described, but most of them are not able to survive in new conditions. These disadvantages can be overcome if we use strains or microflora enriched by the gene pool localized on the plasmid pSinA Sinorhizobium sp. M14, carrying genes responsible for arsenic metabolism.

Objectives
The aims of this study were to (i) investigate the impact of biotic and abiotic factors, simulating the natural environment, on the growth kinetics and (ii) develop a kinetic model of arsenite oxidation by Sinorhizobium sp. M14.

Methods
The effect of different factors on arsenite biooxidation (i.e. start OD, substrate concentration, mixture of arsenic compounds, temperature and pH) on the efficiency of oxidation of arsenites and the changes of physiological condition of the studied strains was investigated. Cultures were grown on MSM supplemented with 0.04% yeast extract. As speciation samples were analyzed by HPLC/ESI MS, while the amount of biomass was monitored by OD/CFU determination. Based on the collected data, a kinetic model of the process was developed.

Conclusions
Culture of microorganisms in different concentrations of As (1;2,5;5 ppm), in mixture of arsenic compounds -As(III)/As(V) (2.5ppm/2.5ppm; 1ppm/4ppm), in different start OD (0.01;0.025;0.06;0.08;0.1;0.15), in various pH (4;6;7;5;10) and in different temperatures (10;16;22;30°C) allowed to develop a model of kinetics arsenite oxidation by Sinorhizobium sp. M14.
INVESTIGATIONS ON MULTIMETAL UPTAKE IN THE PRESENCE OF PESTICIDE BY FUNGAL ISOLATE

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Background
Wastewaters are often contaminated with complex cocktail of pollutants, out of which heavy metals and pesticides pose significant health risks. However, majority of the researches focus on bioremediation of individual metals or pesticides in isolation.

Objectives
Therefore, efforts are needed to obtain microbial strains which can simultaneously handle multiple pollutants.

Methods
Here a newly isolated fungal strain was assessed for its multimetal and pesticide uptake potential in the composite salts medium using glucose and lindane pesticide as the carbon and energy sources. This strain was able to grow in the presence of 1% glucose, 30 mg/L lindane and 30 mg/L multimetal (5 mg/L each of Cd\textsuperscript{2+}, Cr\textsuperscript{6+}, Cu\textsuperscript{2+}, Ni\textsuperscript{2+}, Pb\textsuperscript{2+} and Zn\textsuperscript{2+}) and could degrade maximum amount of lindane to 1.92 ± 0.01 mg/L in about 72 h. The cube root growth kinetic constant (k) of the strain was obtained as 0.0211 g\textsuperscript{1/3} l\textsuperscript{-1/3} h\textsuperscript{-1} in batch study. However, the presence of lindane caused an inhibition effect in the uptake of heavy metals and thus only Pb and Zn were brought down below the permissible mandates for irrigation. The results indicate that the growth kinetics and pollutant uptake is affected by the nature of contaminant matrix. More studies towards the elucidation of mechanisms of metal removal by the fungus in presence and absence of pesticide by various spectroscopic and microscopic techniques such as SEM, TEM-EDX, FTIR, XRD and XPS have been conducted.

Conclusions
These results reveal the inherent potential of this fungal strain to remove multimetals in the presence of pesticide from contaminated wastewater.
SEARCH FOR BACTERIA IN A MEDIEVAL ARCHEOLOGICAL SITE IN VERDUN (FRANCE) AND ISOLATION OF CUPRIAVIDUS NECATOR B9, A NEW METAL RESISTANT STRAIN

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Background
Remains of a well-preserved medieval foundry were excavated by archaeologists in 2013 in Verdun (France). Three to four ancient workshops specialized in brass and copper alloys were found with an activity between 13th to 16th c. Levels of Cu, Zn and Pb reached 20 000, 7000 and 6000 mg kg⁻¹ (dw), respectively, in several soil horizons.

Objectives
The objective was to examine the microbial community in this contaminated site.

Methods
A total of 8–22 10⁶ reads were obtained by shotgun metagenomics in each horizon (length : ± 220 pb). Bioinformatic analyses suggest the presence of very complex bacterial communities dominated by Proteobacteria. Using selective media three Cupriavidus necator/eutrophus strains were isolated. One of those strains, strain B9, differed from C. necator N1 and C.eutrophus H16 on several aspects. For instance, stain B9 was not able of autotrophic growth but was copper resistant (up to 1.2 mM). The strain features a plasmid that differs in size in comparison to the one observed in R. eutropha H16. Stain B9 was therefore characterized through genomic sequencing and a total of 516 contigs were obtained (Illumina sequencing, N50=25246 pb).

Conclusions
Preliminary analyses of the contigs confirm the absence of cbb genes necessary for autotrophy and show the presence of a ~70kb region containing metal detoxication genes (mainly copper but also a mercury resistance merRTPCA operon) also found in environmental Ralstonia pickettii strains and mainly an extended cluster of cop genes (cop KH3SRABCDIH2GFL1L2QH) largely syntenic with the cop region of C. metallidurans plasmid pMOL30.
METAGENOMIC PROFILING OF MICROBIAL METAL INTERACTION IN RED SEA DEEP-ANOXIC BRINE POOLS’ WATER

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Background
Different geochemical studies reported high metal abundance in Red Sea deep-anoxic brine pools, especially in Atlantis II Deep, which has the highest metals content. Brine pools showed wide diversity of biologically essential and non-essential metals. Several metals known for their toxicity to biological life were detected in these pools. Yet, our previous microbiome analyses of the pools demonstrated vast microbiological diversity.

Objectives
In this study, we compare metal-resistant prokaryotic microbiomes in different metal-rich brine water samples from; Atlantis II lower convective layer (ATII-LCL), Atlantis II upper convective layer (ATII-UCL), Discovery Deep (DD) and Kebrit Deep (KD).

Methods
Using 16S rRNA pyrotags and shot-gun 454-pyrosequencing, we perform a comparative analyses of a-taxonomic assignment of Operational Taxonomic Units to major bacterial and archaeal groups and b-metal resistant protein-coding genes, of the microbial communities and metagenomes. The ATII-LCL, ATII-UCL, DD and KD brine pools metagenomes protein-coding genes involved in microbial-metal interaction and resistances were assessed for abundance, diversity and novelty.

Conclusions
We report specific microbial diversity of these three brine pools. Functional analyses of the metagenomes revealed different metal resistance mechanisms. This was supported by the strong correlation between specific high metal/s concentration in selected brine water, where; metal resistance, enrichment of metals metabolism and transport were revealed. As expected, ATII-LCL showed the highest relative abundance of genes involved in microbial-metal interaction. Additionally, we report significant abundance of peroxidases-encoding genes, mainly in ATII-LCL, and we hypothesize that generation of H₂O₂ occurs through interaction of pyrite deposits.
Background
The brown rot fungi are important components of the coniferous northern hemisphere forest ecosystem and are characterized by their ability to colonize and degrade lignocellulosic materials; preferentially metabolizing hemicellulose and cellulose components and leaving a residue of modified lignin. Multiple aspects of the metabolism and degradative capabilities of these organisms are dependent upon cations, including but not limited to iron, manganese, and calcium.

Objectives
As wood degrades, it is characterized by decreasing pH and increasing concentrations of selected cations. The decay fungi, as they colonize wood, differentially translocate cations and actively modulate their ionic environment. The base cations are needed for many metabolic functions including membrane stabilization and they also function as enzyme cofactors and electrolytes. Transition metals are directly involved in lignocellulose degradation. We will examine the influence of cation concentrations on fungal sheath formation, and the activity of a biologically unique iron-based non-enzymatic degradative system in the brown rots.

Methods
The role of iron in the chelator-mediated Fenton (CMF) system will be examined as it is particularly crucial to the ability of the brown rot fungi to depolymerize lignocellulose. Methods used include but are not limited to ICP and XFM. Organisms used in this work include: Serpula lacrymans, Postia placenta, Fomitopsis pinicola and Gloeophyllum trabeum.

Conclusions
The role played by transition metals, pH, oxalate production and calcium oxalate crystal formation in brown rot physiology and cell wall breakdown mechanisms is discussed along with the potential ecological significance of cation mobilization and redistribution within the forest floor ecosystem.
ADHESIN COMPETENCE REPRESSOR (ADCR) FROM STREPTOCOCCUS PYOGENES CONTROLS ADAPTIVE RESPONSES TO ZINC LIMITATION AND CONTRIBUTES TO VIRULENCE

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**Background**
Altering zinc bioavailability to bacterial pathogens is a key component of host innate immunity. Thus, the ability to sense and adapt to the alterations in zinc concentrations is critical for bacterial survival and pathogenesis.

**Objectives**
- To understand the adaptive responses of group A Streptococcus (GAS) to zinc limitation
- To elucidate the regulation of adaptive responses by metalloregulator AdcR

**Methods**
- RNA sequencing
- Gel mobility shift assay
- qRT-PCR
- Animal infection studies
- X-ray crystallography

**Conclusions**

Genes involved in zinc mobilization and conservation are derepressed during mild zinc deficiency, whereas the energy-dependent zinc importers are upregulated during severe zinc deficiency.

We also demonstrated that transcription activation by AdcR occurs by direct binding to the promoter. However, the repression and activation by AdcR is mediated by its interactions with two distinct operator sequences.

Mutational analysis of the metal ligands of AdcR caused impaired DNA binding and attenuated virulence, indicating that zinc sensing by AdcR is critical for GAS pathogenesis. Together, we demonstrate that AdcR regulates GAS adaptive responses to zinc limitation and identify molecular components required for GAS survival during zinc deficiency.
MECHANISMS OF CHROMATE RESISTANCE AND REDUCTION BY KLEBSIELLA PNEUMONIAE ChroAq1 ISOLATED FROM A CHROMIUM CONTAMINATED AQUIFER

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Background

Chromium-reducing microorganisms as well as their enzymes chromate-reductases have taken great interest due to their potential application in bioremediation processes. They are phylogenetic and metabolically diverse; in some cases Cr(VI) reduction is result of the metal respiration as in Geobacter. However in other microorganisms like genus Klebsiella has little knowledge about the metal reduction mechanisms.

Objectives

The present study was aimed at the characterization the mechanisms of Cr(VI) resistance and reduction in Klebsiella pneumoniae ChroAq1, isolated from long-term Cr(VI)-contaminated aquifer located in Guanajuato, México.

Methods

It was determined the ability to reduce Cr(VI) to Cr(III) by growing cells and cell-free extracts of K. pneumoniae ChroAq1. The cellular fractionation was performed by ultracentrifugation. Moreover the complete genome of ChroAq1 was sequenced with illumina technology, automatically annotated and analyzed. Some putative genes involved in resistance and reduction were deleted by double recombination.

Conclusions

The isolated strain ChroAq1 was identified as Klebsiella pneumoniae, its resistance observed was 1.8mM of Cr(VI) in anaerobic conditions and 22mM in aerobic conditions and was able to reduce Cr(VI) only anaerobically. The cell-free extract from this strain showed a NAD(P)H-dependent chromate-reductase activity associated to the soluble cell fraction. The gen chrA is present in the genome of
ChroAq1 however its deletion did not abolish the chromate resistance. In other hand, we deleted four genes presumptively involved in Cr(VI) reduction, one encodes a protein 84.57% identical to YieF (ABJ74146), other encodes one 86.58% identical to NemA (P77258) from *E. coli*, however they had no effect on the chromate reduction capability, the other two are being studied.
PRODUCTION OF SELENIUM AND ZINC ENRICHED BIOMASS AND NANOPARTICLES USING LACTOBACILLI
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Background

Selenium (Se) and Zinc (Zn), are the trace elements which play a pivotal role in humans well-being. Symptoms resulting from deficiency of these essential metals are diverse but chronic and severe and if untreated, can prove fatal. Bioproduction of Se and Zn enriched biomass and nanoparticles by lactobacilli can be explored and exploited for development of dietary organic sources of these minerals to overcome the mineral deficiencies and to improve the human health.

Objectives
To produce Se and Zn enriched biomass and Se⁰ and zinc oxide nanoparticles using lactobacillus spp.

Methods
Fifty human origin lactobacilli were screened for their ability to accumulate Se and Zn by growing them on medium added with different concentration of sodium selenite and zinc sulfate respectively. Estimation of Se and Zn accumulated by cultures was carried out by ICPES and AAS, respectively. Selected cultures were used for biogenic production of Se and ZnO nanoparticles and their size were determined by Scanning Electron Microscope.

Conclusions
Among all the isolates, L. reuteri NCDC77 was found to have greater ability to accumulate Se (820 μg/g dry weight) coupled with potential probiotic properties while L. plantarum YM2C was able to accumulate large amount of Zn (1.32mg/g dry weight). In addition, nanoparticles of Se and Zn produced by the selected strains, were found to be in the range of 100-200nm. Harnessing these potential cultures can serve as a novel application of these cell factories for the production of Se and Zn enriched functional foods and as dietary supplement.
Metals and microbes

ACCURATE MINIMAL INHIBITORY CONCENTRATION PREDICTION FOR CUPRIAVIDUS METALLIDURANS CH34 USING A TOXICITY MODEL BASED ON TWO CHEMICAL METAL-ION CHARACTERISTICS

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Background
The inhibitory effect of metal(-ion)s on the growth of microorganisms is mainly determined by their chemical binding properties to essential biochemical molecules (e.g. thiols, proteins, enzymes, RNAs, DNA) and the effective intracellular concentration. Active control of the intracellular metal-ion concentration is one of the successful survival strategies microorganisms use against toxic metal-ion concentrations and they can harbor a variety of resistance systems to control the intracellular concentration of different metal-ions. Cupriavidus metallidurans CH34 is one of the model bacteria used to study survival in the presence of elevated metal-ion concentrations. Numerous studies have been performed using C. metallidurans CH34 to elucidate its various metal-ion defense mechanisms. As a consequence, a substantial amount of physical, chemical and biological data is available on the impact of several metal-ions on the behavior of C. metallidurans CH34.

Objectives
To test whether it is possible to establish a metal-ion toxicity model that is able to predict the Minimal Inhibitory Concentration for C. metallidurans CH34.

Methods
A two-variable metal-ion characteristics toxicity model, based on $\log K_{OH}$ ($K_{OH}$ = first hydrolysis constant) and $\chi^2_{m-r}$ (= covalent index), that efficiently correlates ($r^2=0.90$) with the measured MIC of 20 different metal ions, has been developed for C. metallidurans CH34 cultivated in chelating-free mineral medium. To test this chemical model for its applicability, the MIC of Pd²⁺ was first calculated and afterwards experimentally determined.

Conclusions
The model prediction, 0.032 mM, corresponded very well to the measured MIC of 0.029 mM, indicating that the two metal-ion characteristics, $K_{OH}$ and $\chi^2_{m-r}$, can be used to reliably predict MIC values.
SEVEN MERA ALLELES ARE SPREAD AMONG ENTEROCOCCUS FROM DIFFERENT SPECIES, ENVIRONMENTS AND CLONAL BACKGROUNDS (1926-2012)

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Background

Successful bacteria accumulated different genetic features shaping their evolution and fitness to diverse environments/hosts. Mercury (Hg) is widespread in Nature and possibly contributed for selection of particular strains.

Objectives

To evaluate the dispersion of diverse merA alleles among enterococci from several origins and clonal backgrounds.

Methods

merA alleles was searched in Enterococcus available genomes, used to construct a maximum-likelihood phylogenetic tree. A PCR scheme+RFLP+sequencing was developed to detect the six merA alleles identified, namely I-(GenBank-AECE01000068), IIA-(AIUL01000023.1), IIB-(ASDU01000008.1), III-(NZ_KB947199.1), IV-(AECE01000068.1) and VI-(NZ_KB030055.1), among 918 Enterococcus (Portugal; human/animal/environment/food; 1996-2012). Clonality was evaluated by PFGE/MLST, merA-transfer by conjugation and Hg-phenotype in m_Enterococcus+128mg/L HgCl₂.

Conclusions

merA were found in 4% (n=37/918) of Enterococcus studied. PCR+RFLP+sequencing distinguished the six alleles previously found at Genbank and a new one (type V). They were distributed among diverse sources and clones, in our and GenBank isolates: types I and IV-human E. faecalis (Effs; n=5; USA; 1987-88; ST9/ST206); IIA-human/pig/trout/feed/hospital-sewage E. faecium-
Enterococcus sp (n=17/1/1; Portugal/France/Denmark/Germany; 1989-2012; 12ST-Enterococcus including from ST78-ST17-ST18-lineages-CC17); IIB-human Efls/E. dispers (n=2/1; Portugal/USA; 1926-2001; ST206/ST105/ST107); III-human/animal/hospital-sewage Efls/E. casseliflavus (n=15/1; Portugal/France/USA/Canada/Japan; 1961-2012; ST64/ST9/ST30/ST159/ST245); V-human/trout/pig-manure Enterococcus (n=4; Portugal; 2001-2006; ST94/ST890); VI-human/hospital-sewage/piggery-soil/piggery-manure/trout Enterococcus (n=23; Portugal, Italy France, Germany, Denmark, Hungary, Switzerland, Ireland, Norway, Brazil, USA; 1961-2012; 14ST including ST78-ST17-ST18-lineages-CC17). The presence of merA among major Enterococcus/ST9-Efls lineages associated with human infections, suggests that mercurial-compounds potentially contributed for their selection/maintenance. merA+ Enterococcus grown in >128mg/L-HgCl₂. Transfer occurred for IIA and VI alleles. All but VI-merA alleles, were also identified in genomes of different species of Firmicutes (data not shown), suggesting genetic exchange of enterococci with bacteria sharing the same communities/environmental challenges.
THE SENSORY PROTEIN HBPS FROM THE SOIL BACTERIUM STREPTOMYCES RETICULI SPECIFICALLY INTERACTS WITH IRON, HEME AND AQUO-COBALAMIN

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Background
A novel type of redox-signaling system has been recently characterised in bacteria, the HbpS-SenS-SenR sytem from streptomycetes. The extracellular HbpS protein binds ferrous ions, heme and aquo-cobalamin.

Objectives
Elucidation of molecular mechanisms that modulate the function of HbpS-SenS-SenR, and analyses of binding sites as well as binding kinetics in HbpS-ligand interactions.

Methods
Disruption and site-directed mutagenesis; growth assays; crystallization; FRET, UV/Vis and CD spectroscopy; bioinformatics

Conclusions
HbpS-SenS-SenR is involved in the protection of Streptomyces against oxidative stress and is conserved in many other actinobacteria. HbpS acts as an accessory module of the two-component system SenS-SenR. Analysis of the HbpS crystal structure and biochemical studies revealed that HbpS assembles as an octamer that is crucial for interaction with the sensor kinase SenS. Under conditions of oxidative stress this leads to the autophosphorylation of SenS that, in turn, phosphorylates the response regulator SenR. This activates the transcription of anti-oxidative genes.

HbpS sequestrers large quantities of ferrous iron ions which might protect Streptomyces from the effects of iron-based oxidative stress. HbpS is also an unusual heme-binding protein in which a specific threonine apparently binds to the tetrapyrrole macrocycle. In vitro and in vivo studies have also shown that HbpS can degrade heme. This activity may be responsible for HbpS-mediated protection against toxic concentrations of heme. Moreover, HbpS binds aquo-cobalamin. The calculated $K_d$ of 34 μM suggests that HbpS might bind aquo-cobalamin in both bacterial cultures and in the Streptomyces natural environment the soil. The physiological relevance of this interaction remains, however, to be elucidated.
Calcium carbonate precipitation is a widespread phenomenon among bacteria, with relevant implications in natural processes and great potentiality in numerous applications. Nevertheless, the molecular aspects of the process are still unknown.

Objectives

We have been studying CaCO₃ precipitation in the model bacterium Bacillus subtilis by different multidisciplinary approaches to identify: a) bacterial genes, and b) cell structure(s), involved in biomineralization; c) analytical markers able to distinguish the "biosignature" of natural calcite.

Methods

For objective a), we produced B. subtilis mutants by insertional mutagenesis and checked their precipitation phenotype. For objective b), we tested bacterial dead cells and cell fractions in a suitable precipitation test in solution. For objective c), we analyzed calcite crystals produced by B. subtilis by X-ray powder diffraction (XRPD) and Electron Paramagnetic Resonance (EPR) spectroscopy.

Conclusions

By analysis of mutants impaired in calcite formation, we identified a gene cluster as involved in the process, and two genes of the cluster, etfA and etfB, as the minimal gene set necessary for precipitation. The B. subtilis cell wall fraction, called BCF, was able to induce calcite formation in solution. BCF was applied on stone specimens and on a monumental site as an eco-friendly biotreatment for stone consolidation. New calcite formation and a little cohesion increase were observed in the treated stones. Both XRPD and EPR revealed unusual spectral parameters, attributed to the effects of the bioprecipitation of the mineral. The resultant spectroscopic fingerprint of bacterial calcite would be useful for identifying traces of bacterial activity in fossil carbonate deposits.
Metals and microbes

ALUMINUM TOXICITY AND TOLERANCE IN MICROORGANISMS
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Background

In acid soils below pH 4.5, aluminum becomes more soluble and toxic to plants, a few micronutrients, such as manganese and iron, become more soluble and toxic; and most plant nutrients, such as phosphorus, become more limited. This situation can cause serious destruction of the environment and significant economic problems. To overcome these problems, understanding of the toxicity of acid soils and the tolerance mechanisms in plants and microorganisms, especially the Al-toxicity and tolerance mechanisms, is essential.

Objectives

We have isolated many plants and microorganisms well adapted to highly acidic soils. Although there have been numerous reports on molecular mechanisms of Al tolerance in plants and microorganisms, such as organic acid secretion, alteration of cell walls, and Al-sequestration, our isolates suggested the complexity of the mechanisms of Al toxicity and tolerance. In this study we studied the Al-toxicity and tolerant mechanisms.

Methods

Microorganisms: Acidocella aluminidurans NBRC 104303(T), Burkholderia bannensis NBRC 103871(T), B. acidipaludis NBRC 101816(T) and 103872, B. heleia NBRC 101817(T), Pullulanibacillus sp. CA42, Rhodotorula taiwanensis CGMCC 2.4753, and Saccharomyces cerevisiae BY4741 and their derivatives.
Al tolerance test: Low phosphate synthetic minimal medium containing 1% glucose was buffered with 5 M succinic acid at pH 4.5. Cells were cultured in the presence of different concentrations Al at 28\textdegree C.

Conclusions

The results suggest that in addition to the aforementioned mechanisms of Al tolerance, an extracellular polysaccharide, cellular functions, and alteration of
membrane lipids, are relevant with the Al toxicity and tolerance. The knowledge will greatly help to overcome the acid soil problems.
Background

Metal nanoparticles have been produced using chemical and physical methods for many years. However, the exploitation of strong reducing agents may lead to undesired toxicity issues. It is, therefore, important to develop alternative and ecofriendly methods. Recently biosynthetic methods employing microorganisms have emerged as simple and viable alternative to produce metal/metalloid nanoparticles. In particular, the chalcogens selenium and tellurium in their elemental forms exhibited interesting antimicrobial activity. Moreover, nanoparticles produced with these elements show intriguing optoelectronic and semiconducting properties.

Objectives

A microbial strain characterized as *Ochrobactrum* sp. MPV1, isolated from the highly metal-polluted site of Scarlino (GR), was studied for its ability of effectively reducing both the oxyanions selenite and tellurite to their elemental forms in aerobic conditions and consequently producing elemental selenium and tellurium nanoparticles.

Methods

The isolate is capable of reducing 2 mM SeO$_3^{2-}$ in 48 hours and 0.3 mM TeO$_3^{2-}$ in 96 hours. The intracellular accumulation of nanoprecipitates was demonstrated through SEM-EDX and TEM analysis. Moreover, several analysis were performed in order to shed light on the mechanisms involved in selenite and tellurite reduction to the elemental state. The results obtained suggest that selenite and tellurite are reduced through two different mechanisms in *Ochrobactrum* sp. MPV1. Glutathione seems to play a major role in selenite reduction, while tellurite reduction could be ascribed to the catalytic activity of intracellular NADH-dependent oxidoreductases.

Conclusions

In conclusion, *Ochrobactrum* sp. MPV1 is an ideal candidate for the biogenesis of Se0 and Te0 nanoparticles, with possible biotechnological and industrial applications.
INVESTIGATION OF FACTORS DETERMINING STRESS RESISTANCE OF SOLVENTOGENIC BACTERIAE STRAIN CLOSTRIDIUM ACETOBUTYLCICUM DERIVED BY ADAPTIVE SELECTION METHOD

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Background
Biobutanol is a promising type of motor fuel and feedstock for chemical industry.

Objectives
To elaborate efficient technology of biobutanol manufacturing from renewable materials adaptive selection technique was applied to derive solventogenic bacterial strain _Clostridium acetobutylicum_ producing butanol in concentration up to 18.7 g/l. It was shown that the selected bacterial strain when compared to the parent strain was capable to withstand 1.3 times higher butanol level and was distinguished by osmotolerance, resistance to temperature impact and antibiotic chloramphenicol.

Methods
To define the factors governing enhanced resistance of selected strain _C. acetobutylicum_ to stress exposure, lipid fatty acid composition of cellular membranes in the parent and butanol-adapted clostridial strains was analyzed. Using real-time PCR technique, we carried out comparative examination of expression of heat shock gene groEL and key gene adhE responsible for butanol biosynthesis in parent and adapted clostridial strains grown on butanol and solvent-free media.

Conclusions
It was found that the ratio of saturated to unsaturated acids is 1.84 times higher in butanol adapted strain _C. acetobutylicum_ than in parent strain, which leads to stabilization of structure, promotes membrane fluidity and induces resistance to stress factors.

It was established that expression of groEL gene in the parent strain declined 100-fold on butanol-containing media, while expression of gene adhE was not recorded, in contrast to adapted strain _C. acetobutylicum_ showing only minor decrease of gene expression under similar conditions. The obtained results indicate that cell metabolism of adapted bacterial strain was less sensitive to adverse butanol effects as compared to wild-type bacteria.
METHANOGENIC CONSORTIA STRUCTURE AND STABILITY CHANGES AFTER LONG-TERM STORAGE

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Background
The start-up of trial operation of biogas plant usually consists of adding leachate from other biogas plants or inoculation by cattle slurry. An alternative to the traditional start-up operation is the use of the freeze-dried microbial consortia that were previously selected in anaerobic digestion process.

Objectives
The aims of this work were investigate the effects of freeze-drying of methanogenic consortium selected from agriculture anaerobic digestion plant on: (i) biogas production efficiency and (ii) stability of microbial community structure.

Methods

Lab-scale anaerobic digestion process. Freeze-drying. Biogas yield: daily biogas production, methane content were analyzed by gas chromatography GC-MS. Physical and chemical analysis: pH, volatile fatty acids (VFA), chemical oxygen demand (COD), ammonium. Determination of total cell counts by fluorescence microscope (Japan) using the filter set WU for DAPI detection. DNA and RNA isolation. PCR amplification and 16S rRNA gene amplicons libraries.

Conclusions
The conducted analyses revealed that the consortium selected from anaerobic digester plant is active and able to effectively produce biogas after long-term storage. 16S rRNA gene amplicon libraries analyses showed that the microbial consortia have the same methanogenic community composition after long-term storage. These results demonstrated that described method for long-term storage and revitalization of methanogenic consortia could be used as a starter to quick start-up industrial process of biogas production.
Background

Hydrogen can be naturally produced by several groups of organisms, including cyanobacteria. As a biofuel, hydrogen has substantial advantages over carbon-based compounds as it does not generate carbon dioxide and other atmospheric pollutants, and has a higher mass energy density than any other fuel. However, the current cost of its industrial production and the low yield of hydrogen by living organisms are not conducive to increasing its use as an environmentally friendly alternative to the traditional fuels.

Objectives

Our goal was genetic engineering of cyanobacteria for the purpose of increasing the yield of hydrogen that was generated by these strains.

Methods

For this purpose, we employed homologous and heterologous expression of hydrogenases in cyanobacteria.

Conclusions

Genetic engineering of cyanobacteria allowed us to significantly increase their hydrogen-producing capacity without compromising the stability of the mutant strains. As the details of molecular regulation of hydrogen production in the cell become increasingly available, new steps are being taken to improve the level of hydrogen production by cyanobacteria. Due to a photoautotrophic nature and low maintenance cost of these microorganisms, they represent the most economical system for biological hydrogen production with a potential for commercial application.
NEW GENETIC APPROACH TO ENGINEERING OF MUTANT STRAINS OF CYANOBACTERIUM ANABAENA VARIABILIS ATCC 29413 WITH ENHANCED LEVELS OF BIOHYDROGEN PHOTOPRODUCTION

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Background
Heterocystous cyanobacteria are capable to efficient nitrogenase-dependent H₂-photoproduction in the case of mutational elimination of uptake hydrogenase activity. In this study we used previously isolated chemically induced H₂-producing mutants of *A. variabilis* PK84 and PK17 with impaired activity of uptake hydrogenase and also ammonium producing ethylenediamine-resistant (ER⁰) mutants ED21 and ED93 characterized by derepressed levels of nitrogenase activity and low levels of glutamine synthetase activity. Our sequencing data revealed the point mutations in conservative sites of genes *hypF* (PK84: Asp374Asn), *hupL* (PK17: Cys446Tyr), *glnA* (ED21: Ala382Thr, ED93: Pro465Leu).

Objectives
The construction of *A. variabilis* strains with mutations in both *hypF* (or *hupL*) and *glnA* genes and analysis of their H₂-producing capacities in comparison with reference strains PK84 and PK17.

Methods
The mobilizable plasmid vector pRL498 was used for cloning and transfer of DNA fragments with mutations into *Anabaena* cells via triparental mating with *E. coli*. Mutant strains PK84 and PK17 were used as recipients for cloned fragments with *glnA*-mutations (with selection of ER⁰ recombinants) whereas mutants ED21 and ED93 were recipients for *hypF* or *hupL* genes inactivated by insertion of spectinomycine resistance cassette (with selection of Sp⁰ recombinants)

Conclusions
In both variants of gene transfer we have selected some biotechnologically perspective double mutants with lower growth rate and higher level of H₂-production compared with strains PK84 and PK17 (up to 130 % per biomass unit) that might be explained by enhanced levels of nitrogenase activity accompanying by a partial block in biosynthetic or assimilatory processes caused by additional alterations in *glnA* gene.
Background
Biodiesel comprising of fatty acid alkyl esters is a promising alternative fuel. The biodiesel fuel can be produced under milder conditions and with fewer steps using biocatalysts instead of conventional alkali catalyst. In order to increase the biocatalyst stability and to ensure its easy separation from the biodiesel fuel, it is required to immobilize the biocatalyst on a suitable carrier. The comparison of biodiesel fuel production with different immobilized biocatalysts will be helpful for the process optimization and for further industrial implementation.

Objectives
In our work, we studied the immobilization of lipases on silica, diatomite, and ion exchange resins, and the immobilization of yeast cells and mycelium on both soft (polyurethane foam) and rigid (haydite) carriers to perform the comparison of different immobilized biocatalysts in stirred-tank and packed bed bioreactors.

Methods
*Pseudomonas fluorescens* lipase, *Rhizopus oryzae* lipase, *Lipase B Candida antarctica*, glutaraldehyde, tributyrine, and triglycerides standards were from Sigma-Aldrich. *Aspergillus niger* DSM823 and *Yarrowia lipolytica* DSM8218 strains were from DSMZ. Activity of biocatalysts was assayed by the hydrolysis of tributyrine with analysis of butyric acid by GC. Methanolsysis of triglycerides was controlled by HPLC. The immobilization of microorganisms was controlled by SEM (see Fig.1, DSM8218 on haydite, bar 10 µm).
Conclusions
The highest yield of biodiesel fuel (fatty acid methyl esters) was observed for the DSM823 mycelium immobilized on polyurethane foam in packed bed bioreactor. This suggests the whole-cell biocatalysis in continuous mode bioreactors is the most promising approach in biodiesel fuel production.

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THE USE OF MIXED HYDROLYTIC BACTERIA IN THE PRODUCTION OF SUPPLEMENTS ENHANCING BIOGAS PRODUCTION PROCESS

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Background
During the degradation of the lignocellulosic biomass by hydrolytic bacteria many organic compounds such as: volatile fatty acids, simple carbohydrates and many intermediate metabolites are released. These compounds are used by successive groups of microorganisms involved in the subsequent stages of anaerobic digestion process.

Objectives
The main aim of this study was (i) degradation of lignocellulosic biomass (maize silage) by mixed hydrolytic bacteria and (ii) production of supplements dedicated for enhanced biogas production.

Methods
(I) Preparation of the supplements: aerobic degradation of lignocellulosic biomass (3% d.m. of corn silage) by mixed hydrolytic bacteria for 72 hours at 30°C at pH 7. Centrifugation and extraction of the supernatant from the culture. (II) Analysis of influence of produced supplements on anaerobic digestion. Lab-scale anaerobic digestion process. Biogas yield: daily biogas production, methane content were analyzed by gas chromatography GC-MS. Physical and chemical analysis: pH, volatile fatty acids (VFA), chemical oxygen demand (COD), total solids (TS), volatile solids (VS).

Conclusions
In the presented work it was showed that the mixed hydrolytic bacteria can be used for preparation of supplements dedicated for the methane fermentation process. The results demonstrate that supplements produced by hydrolysis mediated by mixed hydrolytic bacteria, can increase the efficiency of biogas production up to 20-30%. Such supplements can be used in both well-working biogas plants in order to increase the efficiency of the biogas production process.
Background
Many organic residues are often used as a substrate in anaerobic digestion, but the efficiency of the process is determined by the level of biodegradability of waste materials. To increase digester gas production from hard-degradable compounds various co-digestion process are applied. If co-substrates are used in anaerobic digestion system it improves the biogas yields due to positives synergisms established in the digestion medium and the supply of missing nutrients.

Objectives
The main aim of this study was to verify if maize silage, which is the most popular substrate used in production of biogas in agriculture plants, can be also a valuable co-substrate used for utilization of hardly biodegradable substrates such as sewage sludge waste (SSW) or residues from coal mines (RCM).

Methods
Laboratory anaerobic digestion process in various ratio corn silage and utilization of waste (3%, 2+1%, 1+2%, 0%). Biogas yield: daily biogas production, methane content were analyzed by gas chromatography GC-MS. Physical and chemical analysis: pH, volatile fatty acids (VFA), chemical oxygen demand (COD), total solids (TS), volatile solids (VS). Determination of total cell counts by fluorescence microscope, using the filter set WU for DAPI detection.

Conclusions
Obtained results showed that maize silage can be used as valuable co-substrates during utilization of various industrial waste materials. The addition of only 1% of the dry matter of maize silage to sewage sludge waste and residues from coal mine increased the biogas yield (even up to 300%).
CHARACTERIZATION OF AN ANAEROBIC THERMOPHILIC GLYCEROL-DEGRADING ENRICHMENT CULTURE

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Background
The glycerol market was totally changed by the biodiesel industry, which resulted in the production of an excess of this compound as an industrial by-product. As a consequence, the price of glycerol dropped and a huge interest in alternatives for its valorisation emerged since then. In the field of Biotechnology research, glycerol is an attractive compound for the microbial production of chemical building blocks.

Objectives
The aim of this work was to investigate thermophilic anaerobic communities capable of conversion of glycerol.

Methods
Thermophilic sludge from a lab-scale anaerobic reactor fed with skim milk and sodium oleate (50:50% chemical oxygen demand) was incubated at 55°C in closed bottles containing bicarbonate-buffered medium supplemented with 10mM glycerol. Periodic successive transfers of the glycerol-converting enrichment culture, combined with serial dilutions were performed. After eight generations a highly enriched, low diversity (microscopic observations and 16s rRNA DGGE profiling) microbial culture was obtained.

Conclusions
The enriched culture converted glycerol mainly to methane (6mM) and acetate (7mM) within 6 days of incubation. A yet unknown organic compound was also produced. Sequencing results obtained on the Illumina platform showed the bacterial predominance of an uncultured Thermotoga species (75 % of the retrieved sequences), an uncultured Anaerobaculum species (13 %) and a close relative to Thermoanaerobacter pseudethanolicus (5 %). Isolation of the new uncultured Thermotoga and Anaerobaculum species is ongoing and their role in glycerol degradation will be assessed.
SIMULTANEOUS AND SEPARATED SACCHARIFICATION OF BANANA PSEUDO-STEM WASTE (MUSA X PARADISIACA VAR. SAPIENTUM L.) AND FERMENTATION OF THE HYDROLYSATE USING MICROBE CONSORSIUM FOR BIOETHANOL PRODUCTION

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Background
Bioethanol fermentation can using the high cellulose substrat such as banana pseudo-stem that has 35.96% of cellulose content. The research about simultaneous and separated saccharification of banana pseudo-stem waste (Musa x paradisiaca var. sapientum L.) and fermentation of the hydrolysate for bioethanol production to get an effective enzyme dosage of hydrolysis, consortium of microbes, method of fermentation has been carried out.

Objectives
The purpose of the research are to get the most effective consortium and method in bioethanol fermentation from banana pseudo-stem waste (Musa x paradisiaca var. sapientum L.) using Simultaneous Saccharification and Fermentation (SFS) and Separated Hydrolisis and Fermentation (SHF) methods.

Methods
The experimental method was used in fermentation process of banana pseudo-stem waste with Complete Randomize Design (CRD) consist of 2 factors and 3 replications. First factor were the combination of method and consorsium (K) and second factor were sampling time (T).

Conclusions
The result showed that the most effective dosage during dosage optimization of α-amylase enzyme was dosage 1 with Dextrose Equivalent (DE) about 23.1; hemicellulase enzyme was dosage 2/3 with DE about 34.9; cellulase enzyme and glucoamylase enzym were dosage 2/3 with DE about 57.4. The result showed that consortium of Pichia stipitis and Saccharomyces cerevisiae and Simultaneous Saccharification and Fermentation (SFS) method were the best consortium and method in bioethanol fermentation of banana pseudo-stem waste that produce the highest ethanol concentration about 6.376% in 72 hours fermentation.
Background

Vinasse is a liquid by-product of the ethanol production. Its high organic content (COD of 20-40g/L) means that its disposal into the environment is hazardous and has a considerable pollution potential. However, vinasse can be treated by fermentation process in bioreactors. In this case, thermophilic anaerobic digestion can be used to treat this wastewater from sugar cane industry.

Objectives

The aim of this work was to isolate thermophilic fermentative bacteria present in vinasse.

Methods

Thermophilic bacteria was isolated by anaerobic serial dilution technique on plates containing fermented vinasse with pH of 6.5 and at 50°C. Colonies were identified by MALDI-TOF mass spectrometry technique.

Conclusions

Bacillus thermoamyllovorans, facultatively anaerobic and amylolytic bacterium was isolated from fermented vinasse. This strain ferments carbohydrate producing lactate, acetate, ethanol, and formate but not hydrogen (Combet-Blanc et al, 1995). The presence of this bacterium shows that vinasse is a source of microorganisms that can be used as inocula in anaerobic treatment of vinasse.
Background

Endo-β-1,4-mannanases are important catalytic agent in several industries. The enzymes randomly cleave the β-1,4-linkage in mannan backbone and release short β-1,4-mannooligosaccharides and mannose. They are classified as glycoside hydrolases.

Objectives

The present study, the endo-β-1,4 mannanase was isolated from thermotolerant bacteria and physicochemistry properties of purified enzyme were characterized.

Methods

Microorganisms (320 isolates) were screened on minimum medium containing locust bean gum at 45°C. The species of selected strain was identified by 16S rDNA sequence analysis. An endo-β-1,4-mannanase was purified to homogeneity by using anion-exchange, hydrophobic, and size-exclusive column chromatographies. The molecular mass was analyzed by SDS-PAGE.

Conclusions

The 16S rDNA sequence revealed that the highest mannanase producing microorganism was Bacillus sp. The specific activity of purified enzyme was 14.5 U/mg. The apparent molecular mass was 38 kDa. The optimal pH and temperature for enzyme activity were pH 6.0 and 60°C, respectively. The enzyme was stable in a pH ranges from 5 to 9, after 16 h of incubation at 4°C and stable up to 60°C for 1 h, and more than 80 % of initial activity remained. This endo-β-1,4-mannanase can be applied in several industries such as animal feed, pharmaceutical, paper and pulp, coffee, and bioethanol.
ESCHERICHIA COLI [NI-Fe]-HYDROGENASES ACTIVITY DURING GLYCEROL FERMENTATION UPON FORMATE SUPPLEMENTATION

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Background

Escherichia coli encodes four [Ni-Fe]-hydrogenases (Hyd).

Objectives

H₂ producing Hyd activity was studied during glycerol fermentation at different pHs upon external formate supplemented (10 mM).

Methods

H₂ evolution was measured by suing pair of Pt and Ti-Si redox electrodes.

Conclusions

At pH 7.5 and 6.5 wild type cells showed similar H₂ production rate (V₁H₂) when in the assays glycerol was added, whereas in formate supplemented assays it was decreased ~1.5 fold at pH 6.5. V₁H₂ was lowered ~2 fold in selC (coding for formate dehydrogenases) single mutant at pH 7.5. At both pHs when formate was added in the assays, V₁H₂ decreased in the strains where Hyd-3 or formate hydrogen lyase 1 related enzymes were disturbed. When glycerol was supplemented in the assays, V₁H₂ was significantly lowered in triple hyaB hybC (coding for large subunits of Hyd-1 and Hyd-2, respectively) selC or hyaB hybC hycE (coding for large subunit of Hyd-3) mutants at both pHs.

Taken together it might be suggested that Hyd-3 becomes mainly responsible for H₂ production during glycerol fermentation when external formate is added. Besides, in the glycerol supplemented assays, three hydrogenases can work in H₂ producing mode and only deletion of three of them decreases the production of H₂. This effect might be due to disturbance of H₂ cycling. Therefore, responsible Hyd enzymes have been revealed under the conditions above.

All this is of significance in application of different carbon sources, especially mixed carbons, in H₂ production technology using bacteria.
PURIFICATION OF FERULIC ACID ESTERASE FROM BACILLUS MEGATERIUM

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Background

Ferulic acid esterase (FAE) are a subclass of carboxylic acid esterases (EC 3.1.1.73) that play role in complete enzymatic hydrolysis of hemicelluloses by hydrolyzing the ferulate ester groups in the cross-linking between hemicellulososes and between hemicellulose and lignin.

Objectives

To screen and purify ferulic acid esterase from thermotolerant bacteria

Methods

A total of 250 microorganisms collected from soil and rotten wood were isolated on minimum medium containing 1% rice straw at 45°C for strains that produced high FAE activity. The selected isolate was identified by 16S rDNA sequence analysis. Enzyme was purified by using anion-exchange, hydrophobic, and size-exclusive column chromatographies. The purified enzyme was characterized for its temperature and pH optima and stability.

Conclusions

The 16S rDNA analysis of selected isolate revealed that it belong to Bacillus megaterium. The optimum temperature and pH of purified FAE were 60°C and pH 7.0. The enzyme was stable in a wide pH ranges (pH 1–8), more than 80 % of initial activity remained after 16 h of incubation at 4 °C and stable up to 80°C for 1 h. The specific activity was 298.01 U/mg. This ferulic acid esterase have potential applications for several industries that high temperature and wide pH range are needed.
EFFECT OF PHOSPHOGYPSUM AND SEWAGE SLUDGE FERTILIZATION ON HEMP (CANNABIS SATIVA) YIELD AND MYCORRHIZA AND SOIL RESPIRATION, MICROBIAL BIOMASS AND DEHYDROGENASE ACTIVITY

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Background
Phosphogypsum and sewage sludge are formed as a by-product during production of fertilizers from phosphate rock and industrial or municipal waste-water treatment, respectively.

Objectives
The aim of present study was to investigate the hemp yield and arbuscular mycorrhizal fungi (AMF) root colonization and spore formation in response to phosphogypsum and municipal sewage sludge fertilization. Additionally the effect of phosphogypsum and sewage sludge on soil respiration, microbial biomass and dehydrogenase activity were also examined.

Methods
The field spots with three hemp varieties Beniko, Białobrzeskie and Tygra were studied. An equivalent of 170 kg per ha pure nitrogen was used as fertilization in treatments with sewage sludge. The phosphogypsum at the level 100, 500 and 1000 kg per ha was applied.
AMF root colonization by microscopic and molecular techniques were studied. For determination of soil microbial biomass fumigation-extraction method (PN-ISO 14240-2) was used. Soil respiration by determination of $\text{CO}_2$ release by titration (PN-ISO 16072) and dehydrogenase activity by TTC method (PN-ISO 23753-1) were measured.

Conclusions
Application in agriculture seems to be one of the most reasonable possibility for utilization of phosphogypsum and sewage sludge, especially as fertilizer in energetic plant production. Additionally, phosphogypsum in combination with sewage sludge significantly increased soil biological activity.

Financial support was provided by EU Project at Warsaw Life Sciences University ‘A program to improve the level of the didactic approach to the question of how to obtain raw plant materials for the purposes of energy production in the context of the Europe 2020 Strategy objectives’.
PALLERONIA ABYSSALIS SP. NOV., ISOLATED FROM THE DEEP MEDITERRANEAN SEA AND THE EMENDED DESCRIPTION OF THE GENUS PALLERONIA AND OF THE SPECIES PALLERONIA MARISMINORIS

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Background

Three strains designated 221-F1\textsuperscript{T}, 221-F2 and 3030-F1 were isolated from the Matapan Vavilov Deep canyon, also known as Calypso Deep in the Eastern Mediterranean Sea, at a depth of 4,908 meters.

Objectives

To describe a new species based on 16S rRNA gene sequence analysis and phenotypic characteristics. These strains were found to be most closely related to \textit{Palleronia marisminoris} and \textit{Hwanghaeicola aestuarii}.

Methods

The description is based on a polyphasic approach with extensive phenotypic, quimiotaxonomic, genotypic and phylogenetic analysis.

Conclusions

The strains were observed to be red-pigmented and to form non-motile cocci or pleomorphic cells. The cells were found to stain Gram-negative, to be strictly aerobic, oxidase and catalase positive. Strains 221-F1\textsuperscript{T}, 221-F2 and 3030-F1 were found to be mesophilic and to grow in medium containing up to 13 % NaCl. The major polar lipids of the three strains were identified as diphosphatidylglycerol, phosphatidylglycerol, phosphatidylcholine, an unidentified glycolipid and an unidentified aminolipid. Ubiquinone 10 (U-10) was found to be the major respiratory quinone. The DNA G+C content of strain 221-F1\textsuperscript{T} was determined to be 64.7 mol %. The new isolates were unambiguously more closely related to the type strain of \textit{Palleronia marisminoris} (95.3\% of similarity) than to \textit{Hwanghaeicola aestuarii} (94.7\% of similarity). Based on phylogenetic, physiological and biochemical characteristics we describe a new species represented by strain 221-F1\textsuperscript{T} (=CECT 8504\textsuperscript{T} =LMG 27977\textsuperscript{T}) for which we propose the name \textit{Palleronia abyssalis} sp. nov. We also propose to emend the description of the genus \textit{Palleronia} and the species \textit{Palleronia marisminoris} to reflect new results obtained in this study.
ANAEROBIC SULFATE-REDUCING BACTERIA IN THE OXIC WATER COLUMN OF THE BLACK SEA

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Background

The Black Sea is a world’s largest meromictic basin, where deep sulfide-rich waters do not mix with the upper oxic waters. Interestingly, sulfate-reducing bacteria (SRB) were detected there not only in the anoxic zone but also in the oxic water layer. A pure culture of a new SRB species, possessing different systems of antioxidative defense, was isolated for the first time from the Black Sea oxygenated water column.

Objectives

We analyzed the phylogenetic composition of SRB community in the Black Sea oxic waters where representatives of \textit{Desulfovibrio-Desulfomicrobium}, \textit{Desulfococcus-Desulfonema-Desulfosarcina} and \textit{Desulfotomaculum} subgroups were revealed. The pure culture of psychrophilic Gram-negative SRB was isolated from the subsurface oxic waters and described as \textit{Desulfofrigus euxinos}. This new species possesses different systems of antioxidative defense including \textit{O}_2 reduction and ROS detoxification.

Methods

FISH, nested and qRT-PCR with primers to 16S rRNA and \textit{dsrB} genes, DGGE and nucleotide sequencing were used to investigate the SRB community. DNA-DNA hybridization, fatty acids analysis and growth tests with different electron donors/acceptors were performed to characterize the SRB species. Oxygen consumption rates were measured with a Clark-type electrode. Key genes of antioxidative defense were identified by DOP-PCR, and inverse PCR was carried out for the sequence analysis of the entire genes.

Conclusions

While some SRB are resistant to aerobic conditions because of multi-component antioxidative mechanisms, their presence in the sea oxygenated waters is likely to depend also on inhabiting anoxic microniches within suspended organic particles. The work was supported by RFBR projects nos. 10-04-00220-a and 12-04-91052-CNRS-a (PICS #6041).
Background

Bacteria play an essential role in food webs and biogeochemical cycles in aquatic ecosystems. Investigating ecological relationships among bacterial taxa and between abiotic and biotic ecosystem components is critical to environmental control and functional traits of individual bacterial group. However, such information is scarce. Here we present the results of a study on the spatio-temporal dynamics of bacterial communities in a stratified lake.

Objectives

The main objectives of the present study were to: (a) investigate the bacterial diversity and dynamics through time and space; (b) identify relations between specific bacterial groups and environmental parameters.

Methods

We collected over a year monthly samples throughout the water column and from the sediment of a holomictic lake in the Netherlands. Subsequently, we sequenced the 16S rRNA genes of bacteria on an Illumina MiSeq platform to determine the diversity and relative abundance. Simultaneously different physical and chemical parameters were measured. Statistical analysis was performed to detect possible interactions among different populations and to determine relations between specific bacterial groups and environmental parameters.

Conclusions

All sequences were affiliated to members of phyla commonly found in freshwater systems. Bacterial community changed dramatically over time and space in the water column, but clear seasonal successions could be observed. Contrastingly, the bacterial community in the sediment was quite stable, and mainly composed of Proteobacteria and Bacteroidetes. Statistical analysis showed Actinobacteria were positively correlated with temperature, while Cyanobacteria and Planctomycetes co-varied with nitrate. Betaproteobacteria, Deltaproteobacteria, Gammaproteobacteria,
*Epsilonproteobacteria* and *Firmicutes* were positively correlated with ammonium and negatively correlated with dissolved oxygen and pH.
EXAMINATION OF THE RARE BIOSPHERE IN MICROBIAL MATS FROM SALAR DE HUASCO, AN ANDEAN HIGH ALTITUDE (3820 M) SALINE WETLAND

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Background
The high-altitude wetlands of the Chilean Altiplano (Atacama Desert) are unique and remote aquatic ecosystems, and are considered part of the cold biosphere. These ecosystems characteristically contain organisms with a high level of endemism that can prosper under extreme environmental conditions. Furthermore, microbial diversity is typically high, and includes a large contribution of previously undescribed groups of Bacteria and Archaea. Microbial mats are common, but are typically small in area and depth relative to those described from other environments.

Objectives
To describe members of the rare biosphere from a micro-profile taken from Salar de Huasco microbial mat.

Methods
Bacterial diversity was described by pyrosequencing of 16S rRNA genes of each layer (5) of a microbial mat (<5 mm depth) as well as metagenomic analysis (454 Roche) of microbial mat samples. Oxygen microprofiles were performed in situ using a microprofiling system (Unisense). Microbial diversity was characterized by the presence of groups of low relative abundance (rare biosphere) and also low sequence identity with sequences of available databases. At the phyla level, rare bacteria from the microbial mats (1.6-0.05% of relative abundance) were affiliated with Planctomycetes, Verrucomicrobia, Chlorobi, Acidobacteria, SR1, Armatimonadetes (ex-OP10), OD1, BRC1, Saccharibacteria (ex-TM7), OP11, Lentisphaerae, Spirochaetes, Synergistetes, and others.

Conclusions
Different members of the rare biosphere were present across the microbial mat micro-profile. Particular rare groups could be associated with the specific conditions in each layer of the microbial mat, providing clues to their likely metabolic function.

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MICROBIAL COMMUNITIES IN OPERATIONAL DRINKING WATER DISTRIBUTION SYSTEMS: IMPLICATIONS FOR DRINKING WATER QUALITY

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Background

Microorganisms inhabiting drinking water distribution systems (DWDS), particularly those attached to pipes forming biofilms, can affect the safety and quality of the water. The presence of unfavourable microorganisms in DWDS is influenced by a number of factors ranging from physico-chemical properties of the source water to the characteristics of the pipe infrastructure itself. The difficulty of accessing the internal surface of pipes has limited the study of the microbial ecology of DWDS. Most previous research in this area has been based on laboratory models and/or assessing only few selected microorganisms under controlled conditions which do not represent real systems.

Objectives

To better understand the microbial ecology of DWDS, microbial communities were monitored in two chlorinated networks supplied with different source waters.

Methods

Biofilm sampling devices fitted with coupons designed to maintain boundary layer conditions at the pipe water interface were installed in the networks. Over a one year-period, water physico-chemical parameters were measured and the dynamics of plankton and biofilm communities were monitored using Illumina MySeq sequencing.

Conclusions

Clear differences in the composition of the microbial communities supplied with different source waters were detected. Microbial diversity in the groundwater supplied network was higher than previously thought in chlorinated systems. Source water and the hydraulic regime in the system were critical factors shifting microbial communities and influencing the quality of the water. This new understanding of the microbial ecology of DWDS is vital to improve control and management strategies to help safeguard drinking water quality and ultimately public health.
Background

Cancer and infectious diseases are still the main two health problems facing the humankind. Actinomycetes from the coastal habitats of Saudi Arabia can represent a good source for drug discovery.

Objectives

The main objectives of the present study were screening the marine actinomycete strains isolated from the coastal habitats of Saudi Arabia for production of anticancer and antimicrobial compounds and identifying them to the species level by the polyphasic approach.

Methods

Eight samples were collected from the Arabian Gulf habitats of Saudi Arabia. The selective isolation of actinomycetes was carried out on three different recommended media. Twenty six strains were selected to study their diversity by examining their morphology and determining their cell wall diaminopimelic acid type and whole-cell sugar pattern. Sixteen representative strains were screened for their cytotoxic and antimicrobial activities and their taxonomic positions were confirmed by the phylogenetic analysis of the 16S rRNA gene sequence.

Conclusions

Based on dissimilarities in their appearance on two media, twenty six strains were selected and primarily assigned to 8 actinomycete genera; *Micromonospora*, *Streptomyces*, *Nocardiopsis*, *Amycolatopsis*, *Microtetraspora*, *Nonomuraea*, *Actinopolyspora* and *Saccharomonospora*. These results showed that there is good actinomycete genus diversity in the Saudi coastal habitats. Out of the selected sixteen strains, seven strains showed promising activity against the tested Gram-positive, Gram-negative bacteria and yeast strains. In addition, the actinomycete extracts showed potent anticancer activities against human T cell leukemia (Jurkat) and human laryngeal carcinoma (Hep-2) cell lines. The results are encouraging as novel strains have been isolated and identified and the active secondary metabolites are promising.
Background

Iron is an abundant redox-active element in many environments and iron(III)-reducing and iron(II)-oxidizing bacteria are largely responsible for microbial iron redox cycling. Three metabolic types of neutrophilic Fe(II)-oxidizing bacteria are known, i.e. microaerophilic, nitrate-reducing, and phototrophic Fe(II)-oxidizers. Based on the geochemical conditions in redox-stratified marine sediments, all three can potentially co-exist with overlapping niches.

Objectives

Consequently, there is potential for competition for their electron donor, Fe(II), that is formed by Fe(III)-reducing microorganisms which can interact with the iron-oxidizing microbial community. However, most studies of iron cycling have so far focused only on a single metabolic type of iron(II) oxidation coupled to iron(III) reduction.

Methods

In our study, we therefore determined the spatial distribution of the different metabolic types of iron-oxidizing and iron-reducing bacteria along the steep redox gradients within a coastal marine sediment from Aarhus Bay, Denmark, by MPN studies, isolations and qPCR. Furthermore, the iron(II)-oxidizing and iron(III)-reducing activity of the different metabolic types and their potential contribution to iron mineral formation within the sediment were quantified by microcosm studies.

Conclusions

From our results we conclude that microbial iron cycling occurs in these sediments and involves microaerophilic, phototrophic and nitrate-reducing Fe(II)-oxidizers as well as Fe(III)-reducers. The quantification of rates of iron oxidation and reduction under dynamic environmental conditions, including day-night cycles, input of nitrate or O₂, reveals the competition and ecological network of iron-oxidizing and -reducing bacteria and is an essential step in order to evaluate the importance of microbial iron cycling in coastal marine sediments.
Background

Algal surfaces are an untapped source of microbial diversity and could represent a promising source of novel therapeutic agents. Very little is known about the diversity and bioactive potential of Gram-positive bacteria associated with the surface of Antarctic macroalgae. Furthermore, it remains unclear the natural distribution, diversity and ecological contributions of Antarctic pigmented heterotrophic bacteria. To our best knowledge, this study is the first research carried out on the diversity and biosynthetic potential of pigmented bacteria associated with Antarctic macroalgae.

Objectives

This study aims to explore the phylogenetic diversity and potential to produce secondary metabolites of Gram-positive, yellow-pigmented bacteria isolated from the surface of intertidal and subtidal Antarctic macroalgae.

Methods

Representative species of green, red and brown algae were collected from the intertidal and subtidal zone (5 and 30 m) of King George Island, Antarctica, in January 2014. Surface-associated bacteria were investigated by cultivation-based methods and 16S rRNA gene sequencing. Yellow-pigmented isolates were screened for the presence of genes encoding polyketide synthases (PKS) by PCR amplification with degenerate primers.

Conclusions

The phylogenetic analysis showed that the yellow-pigmented epibionts belonged to the genera Arthrobacter, Citricoccus, Kocuria, Labedella, Microbacterium, Salinibacterium and Staphylococcus. PKS sequences were detected in Kocuria and Staphylococcus isolates. It highlights that Antarctic macroalgae are a unique source of microbial diversity for natural product research.

Acknowledgments
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STUDYING THE GREAT POTENTIAL OF CULTIVABLE BACTERIAL COMMUNITIES ASSOCIATED WITH THE BROWN ALGA ASCOPHYLLUM NODOSUM

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Background
Bacteria associated with algae differ markedly from those living freely in seawater and represent great potential for the production of diverse bioactive compounds as they interact in multiple complex ways with their host.

Objectives
This work aims to study cultivable bacteria associated with the brown alga Ascophyllum nodosum, showing polysaccharolytic activities.

Methods
To isolate cultivable microorganisms, algal thalli of Ascophyllum nodosum were swabbed with sterile cotton tips and marine agar plates were inoculated. Three-hundred isolated bacteria were screened for agarase, kappa- or iota-carrageenase activities on specific marine media. Thirty-two bacteria with polysaccharolytic activities were isolated and a part of their 16S rDNA (8F-1492R) were amplified and sequenced. Twenty-seven were classified as Flavobacteria and five as Gammaproteobacteria. Putative new strains and species of Zobellia, Maribacter, Cellulophaga, Shewanella, Glaciecola, Pseudoalteromonas and Colwellia were identified by phylogenetic analysis. Genomics libraries with their DNA were constructed in Escherichia coli and Bacillus subtilis and are currently screened for diverse enzymatic activities (agarases, iota- and kappa-carrageenases, cellulases, beta-glucosidases, sulfatases and amylases).

Conclusions
In an era where high throughput sequencing is mostly used to study bacterial communities, cultivation methods are underestimated. Here, we revealed that only ten percent of the cultivable bacteria on this brown alga could degrade algal polysaccharides, which lead to asking us; who and what are the 90 other percents doing there? Furthermore, by this cultivation method we could also identify putative new bacterial strains/species, which are screened for polysaccharidases. Novel glycoside hydrolases from unknown marine bacteria represent great biotechnological potential as they should have original industrial properties.
Background

*Microcystis aeruginosa* is a bloom forming cyanobacteria impacting freshwater environments worldwide. Blooms are formed by a mix of strains, some able to produce a hepatotoxic peptide called microcystin, others not.

Objectives

Our objective was to compare two *M. aeruginosa* strains, a microcystin producing and a non producing one, isolated from reservoirs in São Paulo, Brazil.

Methods

The strains were cultured (23±2 °C, 40-50 µmol photons m⁻².s⁻¹, 14-10h light–dark cycle, ASM-1 medium), cells collected at exponential growth phase and morphological and physiological aspects were investigated. Proteomic shotgun analysis was performed using iTRAQ and 2DLC-MS/MS.

Conclusions

The strains were similar in chlorophyll-a and phycocyanin content, the non toxic strain had higher carotenoid content. The toxic strain was richer in antioxidants. It formed floating colonies with numerous cells embedded in mucilage (diameter 150-800µm). In contrast, the non toxic strain exhibited small colonies (100µm) with few cells dispersed in thick mucilage, not floating. Electronic microscopy revealed thylakoids and polyphosphate granules equally present in both strains, aerotopes more abundant in the toxic one, cyanophycin granules and carboxysomes more evident in the non toxic strain. Sixty nine differentially expressed proteins were found. Both strains expressed proteins related to photosynthesis, energy metabolism and translation. Gas vesicle and stress response proteins were more abundant in the toxic strain while transport and protein folding functions were more expressed in the non toxic one. These results are discussed in relation to the other physiological aspects investigated. These differences probably represent distinct ecological strategies adopted by each strain.
Background
The release of effluents into aquatic systems leads to an increase in nutrients, a process known as eutrophication. This process promotes large proliferations of cyanobacteria (blooms) that can be harmful to the environment and to humans due to the production of cyanotoxins. The cyclic peptide microcystin (MC) is the prevalent and most studied cyanotoxin in blooms, with more than 80 variants. It has been reported consistently since 1996 in Jacarepagua Lagoon (RJ). MC is rarely found dissolved in the water column, but adsorbs to particles in suspension which settle at the bottom sediment or undergo biodegradation by microorganisms.

Objectives
The objective of this study was to evaluate the potential for biodegradation of MC-LR by bacteria found in the lagoon sediment, testing both recently isolated strains and microbial assemblages.

Methods
MC-LR was incubated with isolated strains (15) or microbial assemblages for 1 and 7 days, and quantified by LCMS.

Conclusions
After 7 days only one strain showed degradation of MC. The microbial assemblage extracted directly from the sediment was very efficient in degradation, decreasing (by ~50%) the added MC. Part of the observed decrease in MC amount was recognized as adsorption to sediment particles. The 15 isolated strains were identified as Ralstonia or Bacillus by 16SrDNA sequencing. This indicated that the culture medium led to selection of microorganisms, probably eliminating those able to degrade MC. On the other hand, MC degradation by microbial assemblages reflects the metabolic diversity of this sample and points to a possible cooperative activity to complete the process of biodegradation.
MOLECULAR CHARACTERIZATION OF PSEUDOMONAS AERUGINOSA POPULATION IN AQUATIC ENVIRONMENTS OF GREECE

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Background

Recent isolation of Ps. aeruginosa strains from Greek aquatic environments and subsequent PFGE analysis demonstrated that these strains have a unique genotype. Environmental isolates may also have a unique phylogenetic position. Literature has revealed that the behavior of Ps. aeruginosa during typing is affected by its habitat and by stress factors such as antibiotics, biofilm formation e.t.c.

Objectives

Considering the advantages of MLST, as a typing method that focuses strictly on 7 conserved housekeeping genes, it was chosen for typing and discrimination of Ps. aeruginosa strains isolated from water samples from all over Greece. Additional information regarding the resistant phenotypes circulating in these environments is cited.

Methods

The activity of 14 antibiotics was tested against all isolates by the disk diffusion method, where additional phenotypical tests were performed according to published protocols. MLST was performed as described by Curran et al, 2004 with some alterations concerning the annealing temperatures of the seven housekeeping genes. The control strains Pseudomonas aeruginosa ATCC 27853, clinical control from HPA/NEQAS (External Quality Control) and PAO1 were used as standards of reference during the MLST and for reproducibility experiments.

Conclusions

The majority of the isolates exhibited the intrinsic antimicrobial resistance, while a significant proportion presented additional resistant mechanisms. The amended protocol of MLST seemed to produce allelic profiles for all the isolates so far tested. Thus this typing method is going to be used in a larger sampling program to obtain more data and to conclude in a reliable phylogenetic analysis for the first time in aquatic environments of Greece.
PHOTOCHEMICAL AND MICROBIAL ALTERATIONS OF DOM SPECTROSCOPIC PROPERTIES IN THE ESTUARINE SYSTEM RIA DE AVEIRO

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Background
Photochemical transformations of chromophoric dissolved organic matter (CDOM) influence microbial communities in aquatic systems.

Objectives
This work aimed to evaluate this influence in the estuary Ria de Aveiro. In order to achieve that, two sites representative of the marine (MZ) and brackish water (BZ) zones of the estuary were regularly surveyed.

Methods
Optical parameters of CDOM indicative of aromaticity and molecular weight were used to establish CDOM sources, and microbial abundance and activity was characterized. Additionally, microcosm experiments were performed in order to simulate photochemical reactions of CDOM and to evaluate microbial responses to changes in CDOM composition.

Conclusions
The CDOM of the two zones showed different spectral characteristics, with significantly higher values of the specific ultra-violet absorbance at 254 nm and of the absorption coefficient at 350 nm and lower SR ratio at BZ than at the MZ, reflecting the different amounts and prevailing sources of organic matter, as well as distinct riverine and oceanic influences. At the MZ, the abundance of bacteria and the aminopeptidase activity correlated with absorbance, suggesting a microbial contribution to the HMW CDOM pool. The irradiation of DOM resulted in loss of color and increase of its bioavailability. However, the extent of photoinduced transformations and microbial responses was dependent on the initial characteristics of CDOM. In Ria de Aveiro both photochemical and microbial processes yielded optical changes in CDOM and the overall results of these combined processes determine the fate of CDOM in the estuary and have an influence on local productivity and in adjacent coastal areas.
IN SITU DETECTION AND LOCALIZATION OF THE PROBIOTIC PEDIOCOCCUS SP. AB1 IN THE GUT OF ABALONE HALIOTIS GIGANTEA

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Background
We previously reported that host-derived Pediococcus sp. Ab1 can colonize the gut of abalone Haliotis gigantea and that the addition of Ab1 to commercial feed can alter gut microflora and increase alginate lyase and VSCFAs in the gut. From those viewpoints, addition of Ab1 can provide valuable effects in abalone aquaculture as well as in other aquaculture animals.

Objectives
In this study, to revealing the dynamics of probiotic Pediococcus Ab1, we tried to establish an optimum FISH protocol, and tried to in situ detection of this bacterium.

Methods
TAMRA-labeled Rpt probe was constructed and samples were fixed by 4% paraformaldehyde and hybridized at 40 °C for 3 h. In the case of described condition, we could discriminate clearly Ab1 from other control bacterial strains (Enterococcus, Lactobacillus). Fluorescence signals of Rpt probe-positive bacteria were easily distinguished from any detectable background autofluorescence.

Conclusions
Rpt probe-positive bacteria were detected in both probiotics supplemented animal and non-probiotics supplemented animal. Rpt probe-positive bacteria formed microbial colonies only in the probiotics supplemented animal. The mean numbers of Rpt probe-positive bacteria in the gut-attached samples of probiotic supplemented animal was 1.94×10⁷ cells/g-gut, and those of non-probiotic supplemented animal were 6.85×10⁵ cells/g-gut, respectively. From the thin section FISH analysis, Rpt positive cells were observed from gut samples of probiotics supplemented animal. Interestingly, Rpt positive cells were also observed from gills samples of them. From these results indicated that probiotic Pediococcus sp. ab1 is colonized and make a micro-colony in their host gut.
REDDUCTION OF BIOFILM FORMATION IN COOLING TOWER WATER SYSTEMS USING NANO-HYDROPHOBIC COATING

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Background
The proposed study is to reduce biofilm formation using nano-hydrophobic coating on cooling tower fill materials. Consequently, enhancing cooling performance, lengthen the material life, reducing clogging and biofilm associated pathogen bacteria.

Objectives
Primary aim is to reduce the slimy biofilm formation on fill material, which will lead to better cooling of water, longer material life and less clogging of the fills. It is widely known that cooling towers’ maintenance is heavily neglected. This end ups in shorter material life, expensive maintenance charges, inefficient cooling, excessive energy consumption and public health risk due to Legionella bacteria dissemination. While novel anti-biofilm approaches are still in nascent phases of development, our efforts are devoted for sustainable management of cooling towers and public health safety, where cooling towers are often associated with deadly Legionnaires’ disease.

Methods
Test surfaces were placed into biofilm reactor along with the untreated control coupons up to 6-months period for biofilm maturation. Natural bacterial communities were monitored to analyze the impact to mimic the real-life conditions. Surfaces were monthly analyzed in situ for their microbial load using epifluorescence microscopy.

Conclusions
Wettability is known to play a key role in biofilm formation on surfaces, because characteristics of surface properties affect the bacterial adhesion. Results showed that surface-conditioning with nano-silica significantly reduce (up to 90%) biofilm formation. Easy coating process is a facile and low cost method to prepare hydrophobic surface without any kinds of expensive compounds or methods.
THE REGULATION OF DIMETHYLSULFONIOPROPIONATE METABOLISM IN RUEGERIA POMEROYI DSS-3

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Background

Dimethylsulfoniopropionate (DMSP) plays a key role in the global sulfur cycle and marine microbial sulfur and carbon metabolism. The marine roseobacter Ruegeria pomeroyi DSS-3 is capable of metabolizing DMSP by either the cleavage or demethylation pathways. The cleavage pathway forms the volatile gas dimethylsulfide (DMS), the largest natural source of sulfur to the atmosphere. The demethylation pathway produces methanethiol (MeSH), which is readily assimilated or oxidized.

Objectives

A model for the regulation of DMSP metabolism has been developed for R. pomeroyi where the availability and turnover of methyl-tetrahydrofolate (THF) is a major regulatory point.

Methods

A combination of RNA-Seq, enzyme assays and growth experiments were performed to test predictions of the model.

Conclusions

According to RNA-Seq, the genes involved in THF biosynthesis and methyl-THF metabolism were significantly up-regulated during growth on DMSP as compared to acetate. Similarly, the specific activities of the methylene-THF reductase and formate dehydrogenase were two-fold higher in cell-free extracts of DMSP-grown cells. During growth on DMSP, treatment of cultures with low concentrations of the dihydrofolate reductase inhibitor trimethoprim increased production of DMS without inhibiting growth. This result was consistent with a model where high levels of methyl-THF metabolism allow DmdA to function at its maximal rate, leading to sufficient MMPA accumulation to overcome the DMSP-based inhibition of RPO_DmdB2, one of the two isozymes for the second step in the pathway. Based on the current
knowledge of DMSP metabolism, the regulation of the two pathways in *R. pomeroyi* appears to be complex and multifaceted.
EFFECT OF BOTANICAL EXTRACTS & ESSENTIAL OILS IN FEED ON LAYING HENS PERFORMANCE & SALMONELLA, E.COLI & TOTAL BACTERIAL COUNT IN FECES

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Background

Ph.d Research idea was to explore plant extracts effects

Objectives
The aim of the present study was to investigate the effect of Botanical Extracts and Essential oils on performance of laying hens

Methods
One hundred and twelve, 40 weeks old, white Novagin laying hens were randomly assigned to seven groups equally (n = 16). Each treatment (group) was replicated four times of four birds per replicate. Experimental diets were prepared by adding A: positive control, antibiotic lincomysin 4.4%, 120mg/kg of feed, antioxidant seldox (BHA, BHT, ethoxiquine and citric acid) 120mg/kg of feed, acetic acid 99.5% pure 0.15ml/kg of feed), B: negative control (N.C) , no antibiotic, antioxidant and acetic acid, C: N.C + black tea poly phenolic extract 1ml/kg of feed, D: N.C + black cumin seed poly phenolic extract 1ml/kg of feed, E: N.C + fenugreek seed poly phenolic extract 1ml/kg of feed, F: N.C + black cumin seed oil 1ml/kg of feed, and G: N.C + fenugreek seed oil 1ml/kg of feed.

Conclusions
Weekly egg production percentage of positive control is significantly higher than negative control (P<0.05) but plant extracts and oils has no significant difference between each other (P>0.05). On the other hand Total bacterial count significantly reduced with increasing age in all treatments except negative control but there is no significant difference between treatments when compared for presence of salmonella and E. coli (P>0.05) in feces.
EFFECT OF ROOT EXUDATES AND DIFFERENT CARBON SOURCES ON BACTERIAL GROWTH OF BACILLUS AMYLOLIQUEFACIENS FZB42 AND BACILLUS SUBTILIS BBG131 AND THEIR LIPOPEPTIDE PRODUCTION

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Background

Rhizosphere is the region that is closed to the roots of plant. Plant root exudates consist of a complex mixture of organic acids, sugars and amino acids, which are used as substrates for the growth of bacteria colonizing this rhizosphere.

Objectives

we highlighted on the effect of root exudates and some of carbon sources which found in the root exudates, on the bacterial growth and lipopeptides production

Methods

Ten of carbon sources (glucose, fructose, sucrose, maltose, xylose, glutamic acid, citric acid, succinic acid, oxalic acid, fumaric acid), and root exudates were used to study bacterial growth and lipopeptides production

Conclusions

Different growth kinetics were observed depending of the substrates. For both strains, growth on oxalic acid is very slow and glucose is the best substrate. As a conclusion, the different behavior of the two strains to consume the root exudates and the different carbon sources, which reflex to the growth and lipopeptides production, may be causing the different ability to colonize the rhizosphere.
ABUNDANCE AND DIVERSITY OF FUNGAL ENDOPHYES FROM THE GOTAJWAL FORESTS ON JEJU ISLAND IN KOREA

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Background

Endophytes are microorganisms that inhabit plant tissues by colonizing internal plant tissues without causing apparent harm to host. Particularly, some fungal endophytes have beneficial roles on plant growth and plant tolerances to environmental stresses such as abiotic and biotic stresses. Accordingly, identifying their abundance and diversity from unique forest ecosystem is an interested subject to discovery the beneficial fungal endophytes.

Objectives

The objective of this study is to construct endophytic fungal resources for discovering their beneficial functions, by collecting diverse fungal endophytes from the Gotjawal forest (Jeju, Korea), a unique forest ecosystem, and identifying their abundance and diversity.

Methods

Various plant tissues were collected from Gotjawal area on Jeju Island in Korea. They were sterilized and then, subjected to the endophytic fungal isolation. Subsequently, they were further pure-cultured and then subjected to the long-term storage in liquid nitrogen and the DNA isolation for the molecular identification, mediated by various DNA barcode markers.

Conclusions

Forty seven plant samples were collected from five Gotjawal forests on Jeju Island and composed of 34 species. Through the molecular identification, 196 fungal isolates were obtained from the plant tissues and classified as 3 classes, 11 orders, 17 families and 29 genera. Especially, fifty fungal MOTUs (molecular operational taxonomic units) were identified and classified further into four genera such as Colletitrichum, Xylaria, Pestalothiopsis, Fusarium. The fungal collection, obtained from Gotjawal forests, was deposited with classification information in National
Institute of Biological Resources as stocks and can become a useful resource to
discovery the beneficial functions of fungal endophytes.
DOES INTERACTION OF THE INVASIVE CORDGRASS SPARTINA DENSIFLORA WITH PLANT GROWTH-PROMOTING RHIZOBACTERIA EXPLAIN ITS TOLERANCE TO PHYSICOCHEMICAL PROPERTIES OF MARSHES SOILS?

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Background
Microbiota in the niches of the rhizosphere zones can affect plant growth and responses to environmental stress conditions via mutualistic interactions with host plants.

Objectives
A glasshouse experiment was designed to investigate the role of a bacterial consortium (Pseudomonas composti SDT3, Aeromonas aquariorum SDT 13 and Bacillus sp. SDT14) isolated from the rizhosphere of Spartina densiflora in its growth and physiological tolerance to the physicochemical properties of marshes soils.

Methods
Plants of S. densiflora were randomly assigned to two soil types with different physicochemical characteristics with and without inoculum for 50 days. Plant responses were examined using growth analysis, combined with measurements of gas exchange, efficiency of PSII biochemistry, total content of photosynthetic pigments and leaf water content. In addition the accumulation of nutrients in roots and leaves were determined.

Conclusions
The inoculation improved growth of S. densiflora through a beneficial effect on its photochemical apparatus due to its impact on chlorophyll concentration. This growth enhancement happened under both soil conditions and was mainly reflected in a greater length and diameter of roots. Modifications of pigment concentrations were linked to an increase in leaf magnesium content. Also, inoculation favoured LWC through the decline in gs and increment in root-to-shoot ratio. Moreover, this consortium was able to stimulate ions uptake in roots and leaves. Plant growth-promoting rhizobacteria of S. densiflora appears to play a significant role in its growth response and tolerance to the physicochemical properties of soils, through diverse protective effects on the photosynthetic apparatus, WUE and mineral nutrient balance.
Background

The plants growing under salinity stress conditions increase ethylene production, which induces root elongation inhibition. The bacterial enzyme 1-aminocyclopropane-1-carboxylate deaminase (ACCD) cleave ethylene precursor, the 1-aminocyclopropane-1-carboxylate, thus mitigating ethylene effects. Some strains also produce phytohormone indol acetic acid (IAA), which increases cell elongation.

Objectives

To determine the effects of inoculation with consortia formulated with IAA- and ACCD-producing endophytic (EB) and rhizospheric bacteria (RB) on salinity stress mitigation of wheat plants.

To compare the effects of EB and RB inoculation, as well as, different levels of bacterial IAA and ACCD production on salinity stress mitigation of wheat plants.

Methods

Twelve bacterial strains were isolated from avocado plants, and four consortia were formulated: 1) EB with higher ACCD and IAA production; 2) EB with lower ACCD and IAA production and 3) RB with higher ACCD and IAA production; 4) RB with lower ACCD and IAA production. Wheat seeds were inoculated with the bacterial consortia, and then seeds were grown under salt stress conditions. Length, dry weight and superoxide dismutase (SOD) activity of wheat shoot and roots were determined.

Conclusions

At lower levels of bacterial IAA and ACCD production, the EB were more efficient than RB consortia mitigating salt stress effects. Between RB consortia, only those with higher production are able to promote the growth of stressed plants. Both EB and RB consortia with higher production are able to increase SOD activity. Finally,
bacterial strains isolated from avocado plants mitigate plant stress and therefore have the potential to be used as commercial inoculum of avocado plants.
Background
Huanglongbing (HLB) is a destructive disease of citrus caused by phloem-limited bacteria, Candidatus Liberibacter spp. Although no known HLB-resistant citrus species or varieties have been identified, several studies report the group of Poncirus trifoliata and some of its hybrids, as more tolerant to the disease. One of the early manifestations of HLB is excessive starch accumulation in leaf chloroplasts.

Objectives
We hypothesize that callose deposition in the phloem may intervene photoassimilate exportation, causing the starch to over-accumulate. Thus, we examined citrus leaf phloem by microscopy to characterize plant responses to infection.

Methods
We have studied 3 genotypes (C. sinensis, C. sunki and P. trifoliata) and 7 hybrids obtained from crosses between C. sunki and P. trifoliata. All genotypes (3 replicates) were graft inoculated with budwood from HLB-infected plants and infection was confirmed by qPCR (TaqMan®) 8 months after plants inoculation.

Conclusions
We observed callose deposition in leaf petioles and accumulation of starch in leaves in all genotypes and hybrids, when compared with mock-inoculated controls. However, accumulation of starch and callose deposition were more abundant in C. sinensis, C. sunki and three of the hybrids. P. trifoliata and four of the hybrids showed the lowest starch accumulation and lower deposition of callose, when compared with others genotypes. Our results indicate that Liberibacter infection is accompanied by callose deposition in different level among the genotypes and the deposition of callose is higher in susceptible genotypes suggesting that the phloem plugging by callose inhibits phloem transport, contributing to the development of HLB symptoms.
FEMS-1816
Plant/microbes interactions

PYROSEQUENCING ANALYSIS OF SOIL FUNGAL COMMUNITIES OF DIFFERENT TRANSFORMATION SYSTEMS IN THE JAMBI PROVINCE (SUMATRA, INDONESIA)
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Background

In many tropical areas rainforests are cleared in order to transform them into rubber and oil palm plantations, although massive transformations are a major threat for biodiversity. Most of the tree species in rainforests are associated with fungi that form mutualistic interactions with roots and function as key link for nutrient exchange between the above- and belowground compartment. Therefore, in addition to the loss in tree species, transformation of primary forests into intensely managed plantations may lead as well to changes in the fungal community structure of soils or even to a loss of species.

Objectives

The goal of this study is to identify potential differences in fungal community structures in soils of tropical lowland rainforests and plantations sites. Furthermore soil-borne plant stress will be evaluated by analyzing arbuscular mycorrhizal (AM) spore density and morphology.

Methods

The study is implemented in one of the largest regions of tropical lowland rainforest in Southeast Asia, the Jambi Province (Sumatra, Indonesia). Two landscapes, National Park Bukit Duabelas and Harapan Rainforest, were investigated. Soils of different transformation systems (lowland rainforest, jungle rubber (extensive), intensive rubber and oil palm plantations) were investigated using the 454 Pyrosequencing approach. Fungal soil communities were characterized using internal transcribed spacer regions ITS1 and ITS2.

Conclusions

A strong increase of AM spore density in plantations compared to rainforest sites was detected supporting that monoculture tree crops may suffer from soil-born stress.
Currently we are investigating if transformation of tree species rich tropical rainforests to species poor plantations correlates with a loss of fungal diversity in soil.
Background

MicroRNAs (miRNAs) are endogenous small-RNAs transcribed from non-coding DNA, matching a target messenger RNA to repress translation or induce cleavage. They act in almost every biological plant activity, e.g. development, abiotic stress tolerance, signal transduction, and in defense from pathogens or parasites.

Objectives

To elucidate miRNAs role in plant-endophyte interactions, we constructed libraries from roots of Solanum lycopersicum endophytically colonized (Pmi) or not (Pm) by the hyphomycete Pochonia chlamydosporia. This fungus shows endophytic behaviour with growth promotion or nematode biocontrol effects. No data are available on tomato miRNAs role and targets in the endophytic interaction.

Methods

Illumina™ NGS of small-RNAs yielded $9 \cdot 10^6$ (Pmi) and $12 \cdot 10^6$ (Pm) reads per library. CLC Genomics Workbench was used for trimming, counting, annotation and data analysis.

Conclusions

Non-redundant, unique small-RNAs (869178 in Pmi, 958026 in Pm), were produced. MiRNAs expression was affected by endophytism. Analyses of tomato miRNAs (miRBase, rel.21), revealed miR156 and miR168 (conserved across higher plants), as most abundant in roots. Four further miRNAs (miR169a, miR169c, miR9473 and miR9476), out of 75 known in tomato, were expressed only in Pmi, with seven further (miR169d, miR1917, miR169e, miR394, miR167a, miR5300 and miR9475) over-expressed and 27 down-regulated (fold change range: 1.2–4.8). 37 remaining miRNAs were equally expressed in both conditions. A Pmi comparative analysis showed that 1732 out of 5055 Pmi down-regulated genes were miRNA targets,
involved in structural protein, metabolism, transcription factor, growth and development, stress-related, signaling pathways, storage and other processes.
CHARACTERIZATION OF THREE ENSIFER MELILOTI BACTERIOPHAGES FROM SOILS SUBJECTED TO ABIOTIC STRESS.

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Background

The presence of bacteriophages in soils was demonstrated as early as 1935. The presence of rhizobiophages in soils suggests that through selection or elimination of certain types of rhizobia, the rhizobiophages influence the evolution of bacterial populations.

Objectives

In this work, we isolated 3 *Ensifer meliloti* bacteriophages (EMph1, EMph2 and EMph3) from the rhizosphere soil of *Medicago marina* collected in two areas of the South of Spain. This study is part of a project in which the diversity of rhizobia strains that nodulate *M. marina* was studied in the Odiel river marshes (Huelva) and in the dunes of San Fernando beach (Cádiz), soils subjected to pH and salt stress.

Methods

Bacteriophages were isolated from rhizosphere soil samples by the enrichment technique of Barnet (1972). Phages were purified by three successive isolation of single plaques. The basic DNA manipulations and molecular techniques used have been described elsewhere. To study their host range, phages were assayed on 22 different bacteria, 13 of them were strains isolated from *M. marina* nodules, along with additional bacteria, including strains of *E. meliloti*, *E. medicae*, *E. fredii*, *R. tropici*, *R. etli*, *R. leguminosarum* bv. *viciae*, *R. l. bv. trifolii* and *Agrobacterium tumefaciens*.

Conclusions

The 3 phages were resistant to restriction with many enzymes tested. EMph1 and EMph3 displayed a wide host range within the whole bacteria tested. On the contrary, phage EMph2 exhibited a narrow host range which was capable of killing only 5 out of 22 bacteria tested.

THE NOVEL ADVANCES IN UNDERSTANDING OF PECTOBACTERIUM ATROSEPTICUM SCR1043 STRATEGIES DURING PLANT-MICROBE INTERACTIONS.

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Background

Many phytopathogenic bacteria colonize heterogeneous interior of their host consisting of various tissues that form diverse microniches for microorganisms. However, the dynamics of bacterial population structure in such heterogeneous and changing system as plant organism is poorly understood.

Objectives

In our investigations we described the dissociation of Pectobacterium atrosepticum SCR1043 (Pba) populations and the formation of various subpopulations in tobacco plants.

Methods

Microbiological methods; RT PCR; light, transmission electron microscopy; immunocytochemistry; chromatography.

The proliferative potential and bacteria distribution within different plant tissues were analyzed from inoculation time till long-term preservation of microorganisms in dead plant debris. We have revealed new structures totally occluding xylem vessels of the host plant and termed them "bacterial emboli". These structures are composed of tightly packed bacterial cells having a predominant spatial orientation and a peculiar way of formation. Using immunocytochemistry and biochemical methods it was shown that high molecular weight products of pectic compounds degradation and Pba exopolysaccharides play an important role in the bacterial emboli formation. Such structures were formed in the plant vessels both in the pronounced and inapparent disease symptoms after Pba infection. Bacterial emboli are likely to form conditions for bacterial downward migration through the xylem vessels resulting in rhizosphere colonization. After plant death bacteria were transformed to viable but non-culturable state.
Conclusions
Our investigation demonstrates that pectobacteria realize different strategies in colonization of various plant compartments and are able to pass through the life cycle symptomatically and asymptatically. This study was supported by RFBR (14-04-01750_A), RSF (15-14-10022), MK-7359.2015.4.
THE NOVEL ADVANCES IN UNDERSTANDING OF PECTOBACTERIUM ATROSEPTICUM SCR1043 STRATEGIES DURING PLANT-MICROBE INTERACTIONS.

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Conclusions
Our investigation demonstrates that pectobacteria realize different strategies in colonization of various plant compartments and are able to pass through the life cycle
symptomatically and asymptotically.
COMPOSING PLANT-MICROBIAL ASSOCIATION RESISTANT TO SOIL SALINIZATION

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Background
Potash mining by Belaruskali concern poses a grave environmental threat in terms of soil pollution with sodium, potassium and calcium chlorides. Salt-induced stress affects growth and productivity of cultivars. A promising method for salinized soil remediation is field introduction of legume crops, like alfalfa (*Medicago sativa*) upgrading soil fertility due to atmospheric nitrogen fixation of selected rhizobial strains and PGRP bacteria. This work was aimed at isolation of beneficial microorganisms enhancing alfalfa resistance to elevated salt concentrations in soil.

Objectives
Halophilic phosphate-solubilizing and nodulating bacteria.

Methods
Recovery and selection of phosphate-solubilizing bacteria was conducted on TY medium containing chlorides of sodium, potassium, calcium in concentration range 3-15%. Taxonomic affiliation of isolates was based on physiological-biochemical characterization using VITEK2 identification system.

Conclusions
Four halophilic bacterial cultures able to grow on TY medium comprising 15% NaCl were isolated from specimens of solid saliferous wastes sampled at Starobin potash deposit. The examined isolates were capable to solubilize phosphates and exert favorable influence on seed germination, growth and development of alfalfa. Treatment of alfalfa seeds by strain FM-3 under saline conditions increased germination rate by 50% as compared to control. Nitrogen-fixing strain *Ensifer meliloti* Mst 3-2 isolated from alfalfa nodules displayed good growth on TY medium containing 2% NaCl. Investigation allowed to refer three phosphate-solubilizing isolates to genus *Bacillus*. The isolated strains of nitrogen-fixing and phosphate-solubilizing bacteria are especially attractive as components of plant-microbial association resistant to soil salinization.
CULTIVATING BIOFUEL CROPS ON MARGINAL LAND: PLANT-ASSOCIATED BACTERIA OF LIGNIN-REDUCED (GMO) ARABIDOPSIS THALIANA AND THEIR CAPACITY TO FACILITATE GROWTH.

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Background

Producing second generation biofuels out of lignocellulosic biomass can contribute to reduce the food-fuel competition on fertile agricultural soils. However, the recalcitrance of lignin polymers is a limitation during processing the biomass to pure cellulose fibers. The efficiency of this process can be increased by using genetically modified crops with lower lignin-contents. Though, the genetic modification can have effects on the phenolic compounds inside the xylem which can influence the endophytic bacterial population.

To even further reduce the food-fuel competition, biomass can be cultivated on marginal land (containing e.g. cadmium (Cd)). An additional advantage can be the reduction of contaminants in these soils by making use of phytoremediation, a process that can be optimised by inoculating bacteria with plant growth promoting capacity (e.g. siderophores, IAA, organic acid and ACC deaminase production).

Objectives

We want to identify the effects of the genetic modification and/or Cd-exposure on the endophytic bacterial population and determine the endophytic growth promoting capacity.

Methods

The endophytes of different tissues of Arabidopsis thaliana wildtype and reduced lignin mutants, grown in a hydroponic system with and without 3µM CdSO₄, were isolated and characterised using 16S rDNA amplification/sequencing. Their in vitro plant growth promoting capacity was tested using colorimetric tests.

Conclusions
Differences in bacterial populations due to the genetic modification and Cd-exposure were observed in stems and seeds but they seemed to be less pronounced in leaves. The root endophytic populations were more or less identical. However, among all tissues differences in growth promoting capacity were found.
Background

Plants lack an immune system in the sense that it exists in animals. However, they are able to recognize potential pathogens. Lipoxygenase signaling pathway plays an important role in regulation of plant growth and development, cell signaling and defense. Lipoxygenase initiates synthesis of group of bioactive compounds collectively called oxylipins (fatty acid hydroperoxides, hydroxy-, oxo-, or keto-fatty acids, divinyl ethers, volatile aldehydes, or jasmonates). Usually oxylipins are not preliminary formed but synthesized de novo in response to mechanical damage, infection and other stress factors. The main role in the biosynthesis of plant oxylipins belongs to the CYP74 enzymes that are non-classical cytochromes P450: allene oxide synthases (AOS), hydroperoxide lyases (HPL), divinyl ether synthases (DES). A number of AOSs and HPLs have been cloned and characterized from various plant species. DESs are less studied. However, divinyl ethers were found in a large number of organisms.

Objectives

We studied interaction between *Linum usitatissimum*, *Ranunculus acris* and *Selaginella moellendorfii* plants with pathogen bacteria *Pectobacterium atrosepticum*SCRI1043.

Methods

Methods of microbiology, molecular biology, biochemistry, mass spectrometry, NMR and UV spectroscopy.

Conclusions

We cloned and characterized DESs from *L.usitatissimum* (LuDES), *R.acris* (RaDES) and *S.moellendorfii* (SmDES1 and SmDES2). LuDES, RaDES and SmDES2 convert
substrates into divinyl ethers (omega5Z)-etherolenic and (omega5Z)-etheroleic acids, respectively. The main reaction product of SmDES1 is (11Z)-etherolenic acid. (Omega5Z)-etherolenic acid possesses bactericide properties, whereas other divinyl ethers – bacteriostatic properties. Expression of DESs genes after infection is 50 times higher than in control. Thus, DES branch of lipoxygenase cascade involved in plant defense against pathogens.
BIOFILM FORMATION OF RALSTONIA SOLANACEARUM IS REQUIRED FOR ITS VIRULENCE

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Background
Cells of a bacterial pathogen \textit{Ralstonia solanacearum} strain OE1-1 (OE1-1) form biofilms on surfaces of tomato cells adjacent to intercellular spaces after invading tomato plants, and cause bacterial wilt. OE1-1 cells in intercellular fluids but not xylem fluids from tomato plants form mushroom-type biofilms. The production of major extracellular polysaccharide, EPS I, positively regulated by a multi-transcription regulator PhcA is involved in the biofilm formation by OE1-1 cells.

Objectives
We analyzed involvement of biofilm formation by OE1-1 cells in its virulence.

Methods
Among PhcA-positively regulated genes, we created \textit{ralA} encoding furanone synthetase-deleted mutant. Biofilm formation of OE1-1 and the mutant incubated in intercellular fluids from tomato plants were observed under the scanning electron microscope. EPS I production and expression of EPS I production-related genes, \textit{xpsR} and \textit{epsB}, were analyzed. Furthermore, virulence of the strains on tomato plants was analyzed.

Conclusions
The mushroom-type biofilm formation of the \textit{ralA}-deleted mutant significantly reduced, compared to those of OE1-1. Furthermore, expression of \textit{xpsR} and \textit{epsB}, and EPS I production of the mutant significantly reduced. Inoculation with the mutant into tomato plants using root dipping resulted in a significant reduction of bacterial growth in intercellular spaces and loss of its virulence. On the contrary, tomato plants inoculated with the mutant through petiole wilted. The transformation with native \textit{ralA} resulted in recovery of not only EPS I production and biofilm formation but also virulence. These results suggest that biofilm formation of which regulation is positively influenced by \textit{ralA} may be involved in OE1-1 virulence.
EFFECTS OF RHIZOBACTERIAL ACC DEAMINASE ACTIVITY ON THE GROWTH OF SOYBEAN UNDER SALINE CONDITION

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Background
1-amino-cyclopropane-1-carboxylate (ACC) deaminase is an enzyme that degrades the precursor of plant hormone ethylene. Plant growth-promoting bacteria that ACC-deaminase facilitate plant growth and development under stress conditions by decreasing plant ethylene levels.

Objectives
This study was conducted to analyse the changes in root architecture and root hair length of soybean induced by three rhizobacteria Pseudomonas putida TSAU1, Pseudomonas aureantiaca TSAU22, Pseudomonas sp. NUU8 strains containing ACC-deaminase.

Methods
ACC-deaminase activity was determined by monitoring the amount of ammonia generated due to hydrolysis of ACC by the rhizobacterial isolates containing ACC-deaminase. Four salinity levels (5.0, 7.5, 10.0 and 12.5 dSm⁻¹) were maintained in the gnotobiotic system using NaCl salt.

Conclusions
Results showed that the salinity adversely affected on the root length of soybean decreasing by 79% the heighest NaCl concentration (12.0 dSm⁻¹). The bacterial strains could utilize ACC an N source indicating the presence of ACC-deaminase that play a role in reducing ethylene levels in plants. The inoculation of seeds with the P. putida TSAU1, P. aureantiaca TSAU22, Pseudomonas sp. NUU8, increased the root and shoot length of soybean at 5.0, 7.5 and 10.0 dSm⁻¹ up to 58% compared to uninoculated plants exposed to salt stress. In summary, based on the results of our work we recommend utilization of rhizobacteria having capacity to produce ACC-deaminase to alleviate salt stress of soybean grown in the saline soils.
CO-INOCULATION OF BRADYRHIZOBIUM WITH PSEUDOMONAS STRAINS TO IMPROVE GROWTH AND YIELD OF SOYBEAN UNDER SALINE SOIL CONDITION
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Background
Salinity stress is one of the most serious factors limiting the productivity of agriculture, especially the negative effects on yield and nitrogen fixation of Leguminous plants. However, there are several reports on the positive impacts of co-inoculation with Pseudomonas and Rhizobium spp., but less attention has been paid how co-inoculation of root-colonising and PGPR bacteria will affect Rhizobium-Legume interactions under salinity stress.

Objectives
The objectives of this research were to observe if the co-inoculation of soybean with Bradyrhizobium japonicum NU1 and Pseudomonas sp.NUU8 can enhance salt tolerance, nodulation, plant growth, pod yield and grain yield of soybean under saline soil conditions.

Methods
The effects of bacterial inoculation on growth, nodulation, pod yield and grain yield of soybean plant grown under saline soil conditions were studied in field experiments. Plants were grown in saline soil of Sherobod district, Surkhandarya province, Uzbekistan.

Conclusions
The results showed that co-inoculation of with B.Japonicum NU1 and Pseudomonas sp. NUU8 gave more benefits in nodulation, plant growth, pod yield and grain yield of soybean compared to plants inoculated with B.japonicum NU1 alone. Under field condition, co-inoculation of B. japonicum NU1 and Pseudomonas sp. NUU8 strains significantly improved shoot dry weight by 38% and root dry weight by 58% of soybean compared with the uninoculated control. The synergistic use of B.japonicum NU1 and Pseudomonas sp. NUU8 also improved the nodulation, plant growth, pod yield and grain yield under salt-stress. The results suggested that these strains could be used to formulate a biofertilizer for sustainable production of soybean under salt stressed field conditions.
Background
Plant growth promoting rhizobacteria, *Azospirillum brasilense* SM releases phytohormone-Indole-3-acetic acid (IAA) and other plant growth regulators into the rhizosphere which enhances plant development. A crosstalk between IAA and gasotransmitter, Nitric oxide (NO) is speculated which may further benefit the plants.

Objectives
To identify NO production by strain SM and unravel potential rhizospheric crosstalk between IAA and NO.

Methods
The study involved creating mutant strains which overexpress essential NO metabolism genes, NO Fluorescence assay, IAA quantification by HPLC, determining plant-bacterial association by SEM, and Real time PCR for gene expression.

Conclusions
Production of NO and presence of NO metabolism genes i.e. nitrous oxide reductase (*nosZ*), nitrous oxide reductase regulator (*nosR*) and nitric oxide reductase (*norB*) were identified by fluorescence assay and PCR sequencing. Improved PGP response of their overexpressing mutant strains was mediated by increased NO and IAA levels. Surface colonization of strain SM on sorghum roots was established by electron microscopy and improved plant development was observed with the mutants. Quantitative IAA estimation suggested that *nosR* and *norBC* influences regulation of IAA biosynthesis in *A. brasilense* SM. The NO quencher, inhibitor and donor reduced or blocked IAA biosynthesis in wild type and mutants, emphasizing a common regulatory role of these molecules in IAA biosynthesis. Expression studies by qPCR showed positive influence of Tryptophan and Arginine on NO genes. IAA biosynthesis gene, indole-3-pyruvate decarboxylase (*ipdC*) was influenced by Tryptophan but not significantly by Arginine. These results impress some shared signalling mechanism or potential crosstalk involving IAA and NO in strain SM.
Background

Pristine Sphagnum-dominated bogs are N-limited and depend on N\(_2\) fixation by diazotrophs or atmospheric N-deposition for their N supply. Sphagnum mosses harbor a very diverse microbiome including numerous methanotrophs that contribute to the carbon supply of the mosses. It has been hypothesized that methane might also stimulate Sphagnum-associated N\(_2\) fixation during early stages of peat development, thereby increasing Sphagnum growth and N content, although other studies did not see this effect.

Objectives

The objectives are to elucidate whether methane indeed does stimulate N\(_2\) fixation in Sphagnum mosses from ombro-, oligo- and mesotrophic field sites and to test whether this is affected by oxygen level.

Methods

N\(_2\) fixation and CH\(_4\) oxidation activity were studied by incubating mosses with \(^{15}\)N\(_2\), \(^{15}\)N\(_2\) + \(^{13}\)CH\(_4\) or no additions. Furthermore, different oxygen regimes (aerobic, microaerobic (either N\(_2\)-He atmosphere or submerged) were applied. DNA and RNA are to be extracted and 16S rRNA and nitrogenase (nifH) genes quantified and analyzed by qPCR and high-throughput sequencing of amplicons.

Conclusions

The isotope tracer-studies showed that only in oligotrophic sites the diazotrophic communities associated with Sphagnum reacted positively to methane addition and microaerobic conditions. For mosses from ombrotrophic and mesotrophic sites, diazotrophic activity was neither affected by methane addition, nor by reduced oxygen concentration. Combined, these results indicate that methane dependent nitrogen fixation may only be important under certain environmental conditions in Sphagnum-dominated peatlands. Conditions proven to be important for
methanotrophy (pH, water level) and diazotrophic community composition are likely to affect diazotrophy as well, and will be subject of future studies.
INTERACTION BETWEEN MICROALGAE AND BACTERIA - ANALYSIS OF COMMON MECHANISMS TO COMMUNICATE

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Background
Algae live in close association with microorganisms. However, our knowledge on the physiology and metabolism of complex microbial communities associated with algae is very limited.

Objectives
A main goal of our research is to understand the underlying molecular mechanism of species divergence and adaptive processes in microalgae and bacterial systems. According to the fundamental questions “Who’s there?” and “What are they doing?”, bacterial communities associated with plants and their functions should be analyzed in more detail. We have analyzed the bacterial populations of different microalga and we speculate that these bacteria have evolved a common mechanism to communicate and manipulate their hosts.

Methods
The diversity and population dynamic of the bacteria were examined using scanning electron microscope, 16S rRNA gene analysis, MS and metagenome analyses. Moreover, the current work is focused on transcriptome analysis.

Conclusions
A comprehensive understanding of microalgae and bacteria interactions requires knowledge of the associated gene expression changes in both the microalgae and the bacteria. So, we will analyze the role of common signaling pathways and signals from eukaryotic and prokaryotic sites. The results will help us to understand overall principles of microbial adaptation and signal exchange in more detail.
Background
Bacteria are able to colonize a wide spectrum of habitats including the rhizosphere of plants. In such association bacteria may utilize plant root exudates components as carbon and energy source, but also other processes may take place, like degradation of toxics, competition with pathogenic bacteria, phytohormones synthesis, nitrogen fixation, quorum sensing control, and several others. Despite the relevance for agronomical applications, the molecular mechanisms underlying microbe-plant root exudates interactions are poorly understood.

Objectives
To improve understanding of plant-microbe interactions through transcriptomica and metabolomics profiling of selected bacteria exposed to plant root exudates when plants have been grown under different nutritional and stress conditions.

Methods
Cell cultures of two different Burkholderiales species able to associate with Arabidopsis (one plant growth promoting rhizobacterium –PGPR- and the other one a non-PGPR) were exposed to plant root exudates obtained from plants grown on control, N-limitation, and saline or chemical stress conditions, and gene expression profiling: Real Time PCR analysis for specific genes, and complete transcriptomic profile analysis by High-Seq Illumina platform, were obtained. Metabolomic profiling of plant root exudates was done by HPLC-MS.

Conclusions
Plant root exudates exhibit different composition profile according to the growth conditions of the plant. This may explain differential gene expression profiles observed with the PGPR and the non-PGPR species.
ENDOPHYTIC BACTERIA THE INTROVERT ENGINEERS OF RHIZOSPHERE: A CASE STUDY EMPLOYING THEIR USE FOR REJUVENATION OF APPLE TREES IN SIX DECADE OLD APPLE ORCHARDS

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Background

Apple cultivation a prominent industry in Himachal Pradesh, India occupies an area of 1,01,485 hectares and production reaching 2,11,295 metric tonnes. Today, the six decade old apple orchards are being replaced by new plantations. However, their survival is a major challenge which results in indiscriminate use of chemical inputs. This poses potential threat to environment/human health and therefore managing plant rhizosphere is critical.

Objectives

Work presented in here, highlights novelty of Bacillus sp. strain CKA1 (apple roots endophyte) in being used as rhizosphere engineer.

Methods

Microbial formulation was evaluated under field conditions for its effect on plant establishment; early growth; yield and rejuvenation of diseased apple trees. Whole genome sequencing of the bacterium helped in deciphering molecular mechanisms underlying the multifunctional plant growth promoting potential. Application of microbial formulation resulted in: increased plant biomass (75.40-80.11%); increased Nitrogen (2.42%), Phosphorus (0.58%) and Potassium (1.33%); increased fruit yields (35 to 45%); rejuvenation of diseased apple trees infected with Dematophora necatrix in farmer’s field over a three year trial. Comparative genomics paved the way for understanding molecular mechanisms involved in the strains ability to function as multifunctional plant growth promoter.

Conclusions
Apple orchard rejuvenation projects are being ambitiously launched by State agencies for restoring six decade old planted apple orchards. Integrated nutrient management systems comprising biological systems especially, managing rhizosphere using endophytes such as strain CKA1 opens up new avenues not only for improving crop yield but also in sustaining soil health.
POTENTIAL FIBRINOLYTIC ACTIVITY OF AN ENDOPHYTIC XYLARIA CURTA SPECIES FROM WESTERN GHATS OF INDIA

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Background

Endophytic fungi are prominent producers of putative phytochemicals and their analogues which can be exploited as therapeutic interventions/or pharmaceutical agents.

Objectives

The current investigation deals with isolation, purification and characterization of fibrinolytic enzymes produced by endophytic fungal isolates derived from different medicinal plants of Western Ghats of India.

Methods

In preliminary screening assays, 17% of the endophytic fungal isolates expressed fibrinolytic activity whereas 26% of the isolates exhibited proteolytic activity. Maximum fibrinolytic and proteolytic activity was shown by endophytic fungal isolate #37 CRSTBRT which exhibited a halo formation of 113.04 mm² on fibrin clot assay. The fibrinolytic enzyme (VM 22) from fungus was purified to electrophoretic homogeneity with the methods including ammonium sulphate precipitation, anion exchange and gel filtration chromatography. The molecular weight of the purified enzyme was estimated to be 35 KDa by SDS-PAGE, fibrin zymography and gel filtration chromatography. The enzyme was stable between 5.5 – 8.5 pH and below 50°C. The optimal pH for the enzymatic activity was 7.8 at 35°C. The purified enzyme also exhibited an 18 fold increase in the enzyme activity. In vitro studies also revealed that the enzyme could effectively catalyses the blood clot lysis. Further the endophytic fungal isolate #37 CRSTBRT was identified as a Xylaria curta species based on morphological and molecular taxonomic tools.

Conclusions

The fibrinolytic enzyme VM 22 directly lyse the fibrin clot and not by plasminogen activators indicating that this enzyme could be useful in thrombolytic therapy.
Background
There are evidence of the participation of salicylic acid (SA) pathway in the plant/virus interaction. Thioredoxins (Trxs) participate as a regulator in this mechanism, and in plants, the h-type play a fundamental role in tolerance of oxidative stress and defense responses against pathogens.

Objectives
The objective of this work was to analyze the effect of silenced CaTRXh1 in plants of Capsicum annuum during the compatible interaction with the begomovirus Euphorbia mosaic virus-Yucatán Peninsula (EuMV-YP).

Methods
CaTrxh1 was silenced in plants of Capsicum annuum using a VIGs vector based on the TRV. 15 days post-silencing, plants were infected with the EuMV-YP. A time course experiments was done. Plant samples were collected at different times post-inoculation. Gene expression of thioredoxin h, NPR1, and PR10, and EuMV-YP replication were evaluated on the time course by Real Time PCR.

Conclusions
NPR1 and PR10 genes increase their expression at 14 and 7 days post inoculation (DPI), respectively. In the silenced and infected plants, the expression of NPR1 decreases in approximately 41% at 28 DPI, while PR10 gene maintains the same expression at 7 DPI. At 4 hour post inoculation (HPI) it was observed an increase in the SA content in the plant infected with the EuMV-YP and at 1 HPI in the pTRV2:CaTRX/EuMV-YP, compared with the control plants. Also, the viral DNA accumulation was higher in the plants silenced and infected with the begomovirus. These results suggest the participation of CaTRXh1 on plants of pepper in the mechanism of defense during the compatible interaction to EuMV-YP.
Background
Ralstonia solanacearum is a soilborne bacterial pathogen and is the causal agent of bacterial wilt of more than 200 plant species, limiting the crop production around the world. The colonization of R. solanacearum strain OE1-1 (OE1-1) in intercellular spaces after invasion through wounds is required for its virulence.

Objectives
We analyzed colonization mechanism of OE1-1 in intercellular spaces.

Methods
The behavior of GFP-labeled OE1-1 cells in intercellular spaces of tomato plants was observed under the fluorescence microscope. OE1-1 cells in intercellular spaces were then observed under the scanning electron microscope (SEM). OE1-1 cells in intercellular fluids and xylem fluids from tomato plants were observed under the SEM. Furthermore, biofilm formation of the major extracellular polysaccharide, EPS I, productivity-deficient mutants was also analyzed.

Conclusions
The observation under the fluorescence microscope showed aggregation of fluorescence from GFP-labeled OE1-1 cells in intercellular spaces. The SEM observation showed attachment of OE1-1 cells on surfaces of tomato cells adjacent to intercellular spaces and biofilm-like structures by OE1-1 cells surrounded by an extracellular matrix. Interestingly, the SEM observation showed that OE1-1 cells incubated in intercellular fluids from tomato plants formed mushroom-type biofilms. On the contrary, incubation in xylem fluids led to a reduction of biofilm formation by OE1-1 cells. Furthermore, EPS I productivity-deficient mutants incubated in intercellular fluids significantly reduced their biofilm formation. Together, colonization of OE1-1 cells in intercellular spaces of tomato plants may consist of their attachment and biofilm formation on surfaces of tomato cells adjacent to intercellular spaces, in which EPS I productivity is involved.
LEAF SPRAYING WITH RHIZOBACTERIA FOR FOR BIOLOGICAL CONTROL OF XANTHOMONAS AXONOPODIS PV. PHASEOLI

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**Background**

The bean common blight (BCB) is the most important bacterial disease in the tropics. The chemical control of the BCB is neither efficient nor economic. Therefore, is important to develop new approaches to control CBB. Previous studies using seed microbiolization showed that DFs strains and combinations were effective in control leaf, root and vascular diseases (CORRÊA et al., 2014) and changed enzymatic activities (SILVA et al., 2009).

**Objectives**

So, here these rhizobacteria were evaluated by spraying them in leaves inoculated with *Xanthomonas axonopodis* pv. *phaseoli*.

**Methods**

The treatments were *Bacillus* (DFs093, DFs348 and DFs769), *Pseudomonas* (DFs513 and DFs831) and their combinations (DFs93+DFs769+DFs831 and DFs348+DFs769+DFs831) or water (test) sprayed 48 or 24 h before, 24 or 48 h after inoculation of the pathogen. The experiment was conducted in a greenhouse, in a design completely randomized, with six repetitions. The BCB severity was assessed after 96 hours each two days using a diagrammatic scale (0 to 6).

**Conclusions**

The test plants were 100% symptomatic after 120 hours and the final severity was between 5.35 and 5.48. Only the spraying 48 h before the pathogen inoculation result in a significant difference between the bacterial treatments. The treatments DFs513, DFs769, and the combination DFs348+DFs769+DFs831 showed under disease progress curve (AUDPC) (20.6, 26.8 and 22.8% respectively). This result suggests the occurrence of induced resistance, since it requires previous spraying, as occurs when these treatments were used to treat seeds.

CORRÊA et al., 2014. Biological Control, 72: 71-75.

Background

Seagrass meadows grow in coastal, biogeochemically active sediments. These productive ecosystems are distributed from tropical to temperate areas and are dominated by seagrasses, a group of marine angiosperms that is often threatened by the presence of hydrogen sulfide, a phytotoxic gas produced by belowground bacterial communities. The seagrass rhizosphere is still poorly described, however the occurrence of die-off events caused by high levels of sulfide stresses the need to understand these plants and bacterial communities present in their rhizosphere – rhizobiome.

Objectives

The aim of this research was to obtain insight into the seagrass rhizobiome and to answer the following questions: a) Is the rhizobiome plant-specific?; b) Does the rhizobiome co-evolve with its host?; and c) Who are the core species of the seagrass rhizobiome?

Methods

Rhizosphere samples of Zostera marina, Z. noltii and Cymodocea nodosa, bulk sediments and seawater were collected from Portugal, and the former two rhizospheres were additionally collected from France. Next Generation Sequencing of 16S rDNA amplicons was performed using MiSeq Illumina platform.

Conclusions

The rhizobiome of seagrasses differs significantly from their surrounding environment. Although they don’t vary at a local scale, the rhizobiomes were significantly different between plants from different geographical locations. The core rhizobiome of seagrasses is mainly composed of OTUs involved in the sulfur cycle (sulfate reduction and sulfur oxidation), and known for their ability to fix nitrogen.
Moreover, our results point to a niche differentiation of sulfur bacteria, in which sulfide oxidation is performed by different taxa.
EFFECT OF GFP-LABELED PAENIBACILLUS POLYMIXA ON GROWTH OF AGRICULTURAL CROPS

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Background
Paenibacillus polymixa strain P2b-2R isolated from internal stem tissue of a naturally regenerating pine seedling fixes nitrogen in association with lodgepole pine seedlings and promotes their growth. It has been shown to colonize lodgepole pine seedling tissues endophytically using a green fluorescent protein (GFP)-labelled derivative of P2b-2R.

Objectives
We wanted to see if the GFP labelled derivative of P2b-2R would fix N and promote growth of corn and canola in ways similar to the wild type strain.

Methods
We inoculated corn and canola seeds with wild type P2b-2R or the GFP labelled derivative of P2b-2R and seedlings were grown for 40 days in a N-limited soil mix. Seedlings were harvested 20, 30 and 40 days after inoculation and evaluated for biological nitrogen fixation and growth promotion.

Conclusions
Seedlings inoculated with the GFP labelled P2b-2R strain derived small amounts of N from the atmosphere (upto 17%) but grew significantly larger than seedlings inoculated with wild type P2b-2R. Seedlings growth was promoted by inoculation with GFP labeled P2b-2R with an increase of upto 40% in height and 70% in biomass as compared to wild type P2b-2R. Thus, we concluded that GFP modification of strain P2b-2R resulted in a significant enhancement of its growth promotion efficacy of corn and canola.
CAN AN ENDOPHYTE ISOLATED FROM LODGEPOLE PINE TREES RESIDE INSIDE AGRICULTURAL CROPS AND FIX N?

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Background
Several bacterial strains of *Paenibacillus* that possessed N-fixing ability were isolated from extracts of surface-sterilized lodgepole pine seedling and tree tissues. One strain, *Paenibacillus Polymyxa* P2b-2R, was found to fix high amounts of nitrogen when reintroduced to lodgepole pine and western red cedar (gymnosperms). But can this bacterial strain fix nitrogen if introduced into agricultural crops?

Objectives
We wanted to determine if this bacterial strain could fix N and promote plant growth while living inside or in rhizosphere of agricultural crops (Corn and Canola).

Methods
We inoculated corn and canola seeds with P2b-2R and grew seedlings for 40-60 days. Corn seedlings were harvested 10, 20 and 30 days after inoculation and canola seedlings were harvested after 20, 40 and 60 days after inoculation. Seedlings were evaluated for biological nitrogen fixation and growth promotion.

Conclusions
Seedling growth was promoted significantly by inoculation with P2b-2R with an increase of upto 35% in height and 30% in biomass from control. P2b-2R also fixed more than 20% of atmospheric nitrogen while living both inside the plant and in the rhizosphere. These results suggest that this bacterial strain has a broad range of hosts and is successful in N-fixation and growth promotion in agricultural crops.
EVALUATION OF TEMPERATURE, PH AND CARBON SOURCE IN INDOLE ACETIC ACID PRODUCTION BY YEASTS ISOLATED FROM RIZOSPHERE AND PHYLOPLANE

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Background
It is possible to find naturally in rhizosphere and phylloplane some yeasts species capable of producing phytohormones, solubilize phosphates and other minerals, as well as acting as biological control agent. Despite that, scarce information is known about this microorganism group as plant growth promoters.

Objectives
Considering these, the aim of this study was evaluate the production of phytohormone auxin (indole acetic acid (IAA)) by yeasts, with variations of culture conditions.

Methods
The yeasts evaluated were Rhodotorula mucilaginosa (2F32) and Trichosporon asahii (3S44), isolated from phylloplane and rhizosphere of sugar cane, respectively, in Brazil, São Paulo State. Yeasts were cultivated in Potato Broth medium, with 20 g/L of glucose or sucrose, as carbon source; the cultivation of yeasts was carried out at temperatures 22°C, 25°C or 30°C; pH was modified to 3.0, 4.5 or 6.0. All cultures were analyzed by colorimetric technique with reaction of broth and Salkowsky reagent; the cells count was realized to evaluate the development of yeasts.

Conclusions
The results for T. asahii showed that IAA production with sucrose was superior at pH 6.0, and glucose was superior at pH 4.5; these results indicate direct relation between carbon source and pH medium. Rh. mucilaginosa presented a high IAA production if compared with T. asahii or even with microorganisms related in literature; the higher production (655 μg/ml) was obtained at pH 6.0 and glucose as carbon source. Temperature was not able to influence IAA production, significantly. Variations of culture conditions did not affect the cell growth of yeasts.
ANALYSIS OF METABOLIC CHANGES IN MESORHIZOBIUM LOTI DURING NODULE MATURATION BY QUANTITATIVE PROTEOMICS

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Background

Rhizobia are nitrogen-fixing bacteria that establish a symbiotic relationship with leguminous plants. After entering the root cells, rhizobia differentiate into bacteroids within the root nodule of the host plant, and provide the plant with fixed nitrogen. Inside nodule, it have been believed that bacteroids construct optimal environment for nitrogen fixation, but the details are still unknown. We have previously evaluated the intercellular proteome of rhizobia1.

Objectives

To understand the mechanism by which rhizobia alter their metabolism to establish nitrogen-fixing symbiotic relationship with hosts, time-course quantitative proteome analysis of Mesorhizobium loti, one of the model rhizobia, was performed during the nodule maturation.

Methods

Lotus japonicus was inoculated with M. loti. Nodules were harvested at 2, 3, and 4 weeks post-inoculation. Bacteroids were extracted from nodules, and their proteins were extracted, digested by trypsin, and labeled by Tandem Mass Tag. The prepared samples were analyzed by LC-MS/MS equipped with a 500 cm-long monolithic silica capillary column2.

Conclusions

Using a quantitative proteomics approach, we identified and quantified 537 proteins in M. loti bacteroids. The results revealed significant changes in the carbon and amino acid metabolisms upon differentiating into bacteroids. Furthermore, our findings suggest that M. loti enters a nitrogen-deficient condition during the early stages of nodule development, and changes to a nitrogen-rich condition in the intermediate stages3.

Reference

1) Tatsukami Y. et al., BMC Microbiol., 13, 180 (2013)
3) Nambu M. et al., submitted
FUNCTIONAL GENOMICS ANALYSIS OF SALMONELLA-TOMATO INTERACTIONS REVEALS MICROBIAL ADAPTATIONS AND PLANT RESPONSES TO THE INFECTION BY A HUMAN PATHOGEN
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Background
As the number and severity of produce-associated gastroenteritis outbreaks increase, we are coming to recognize that we know little about the ecology of Salmonella and enterovirulent E. coli outside of their animal hosts.

Objectives
The objective of this work was to delineate mechanisms behind interactions of a model human pathogen, Salmonella enterica, and its alternate plant host.

Methods
TnSeq and RNAseq analyses of Salmonella behavior in tomatoes and tomato responses to the colonization by the pathogen were used to reveal key mechanisms of bacterial adaptation to persistence within tomato and plant responses to the colonization by the human pathogen. Expression of individual genes was confirmed by qPCR and Recombinase-based in vivo Technology (RIVET).

Conclusions
Salmonella virulence genes located on the Pathogenicity Islands were not involved in persistence within tomato fruit. We identified important functions for the regulators of capsule synthesis (yihT, rcsB) in the phenotype. Salmonella genes involved in de novo synthesis of amino acids were critical to persistence within tomato fruits. Salmonella gene expression depended on the host maturity and genotype, for example, RIVET assays confirmed that cysB was expressed strongly in a tomato cultivar known to be resistant to a tomato pathogen Ralstonia solanacearum, while fadH was responsive to the maturity-dependent accumulation of specific fatty acids. Proliferation of the pathogen was restricted in the plant mutants defective in ethylene synthesis and response (rin, Nr, nor). Tomato genes involved in responses to plant pathogens were activated in response to Salmonella infection, however, typical responses to infection were only observed in ethylene mutants.
Background

Bacterial endophytes colonize the internal tissues of plants without causing harm. An important subgroup are seed endophytes that are transferred from one generation to the next and that are already present during the early plant growth stages. Several studies suggest that seeds can serve as a vector for beneficial bacteria and that these bacteria could improve plant growth.

Objectives

We want to determine the effect of different cultivation substrates, which are frequently used during bulking of Arabidopsis thaliana seeds, on the seed endophytic population of this model plant.

Methods

Arabidopsis thaliana was cultivated on sand (bacteria-poor) and on a mixture of sand with potting soil (bacteria-rich). Seeds harvested on both substrates were sown again on the same substrate until the radicle emerged or until there were 3 week-old leaves. Cultivable endophytes were isolated from soil, seeds, radicles and leaves, and identified using 16S rDNA amplification and sequencing. Total bacterial populations were identified by a direct DNA extraction on soil, seeds, radicles and leaves, followed by amplification and 454 pyrosequencing of the V5-V7 hypervariable region of the 16S rDNA.

Conclusions

Despite large differences in the soil bacterial populations, the seed endophytic populations were very similar, which indicates that plants are able to select which bacteria can become seed endophytes. This can impact on the establishment of the endophyte community of the next generation of plants.
BACKGROUND
The use of inorganic fertilizers has been established to cause environmental problems.

OBJECTIVES
The study investigates the effect of a biofertilizer, *Trichoderma longibrachiatum* NGJ167 and NPK fertilizer on the growth of two varieties of lettuce (Greatlake and Eden).

METHODS
The study was carried out in the screenhouse by inoculating soils with mycelial plugs of *T. longibrachiatum* NGJ167 and compared with NPK-fertilizer. Thereafter, *in-vitro* and conventionally propagated lettuce were planted on the treated soils and plant growth parameters observed.

CONCLUSIONS
Results showed that the micro-propagated Greatlake lettuce treated with *T. longibrachiatum* had higher plant height (22.23 cm) while the *in-vitro* propagated Greatlake lettuce treated with NPK-fertilizer had mean plant height value of 14.17 cm. The conventional propagated Greatlake and Eden treated with *T. longibrachiatum* had mean plant heights of 17.23 cm and 15.07 cm respectively while the controls had average values of 2.67 cm and 12.4 cm respectively. The average number of leaves of the *in-vitro* propagated Greatlake and Eden lettuce treated with *T. longibrachiatum* (7 and 4 respectively) were higher than those of the NPK-fertilized Greatlake and Eden lettuce (6 and 5 respectively). The result of the fresh biomass revealed that the conventional propagated Eden lettuce treated with *T. longibrachiatum* had an average value of 19.77 g while the NPK-fertilized plants had a mean of 15 g. The implication of these results showed that *T. longibrachiatum* NGJ167 was better than NPK fertilizer with respect to the growth of lettuce. The study concluded that the use of *T. longibrachiatum* improved the growth of lettuce.
MOLECULAR CHAPERONES OF THE TYPE III SECRETION SYSTEM ARE NOT REQUIRED FOR SECRETION OF THEIR PARTNER SUBSTRATES, BUT CAN BLOCK EFFECTOR TRANSPORT IN PECTOBACTERIUM CAROTOVORUM

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Background

Pectobacterium carotovorum (Pca) is a plant pathogenic soft rot bacterium that can use the type III secretion system (T3SS) to suppress host defence. T3SS is utilized by many Gram negative bacterial pathogens to inject bacterial virulence proteins into eukaryotic cells, where they can manipulate host cell processes to pathogen advantage. T3SS chaperones are involved in protein transport, but their exact function is still unclear in many cases. In Pca two potential T3SS chaperones are coded for by the divergent transcriptional units, dspEF and hrpWshcW. Chaperone function was suggested for the DspF protein in related bacterium Erwinia amylovora, but not in Pca, while the ShcW protein was not studied before.

Objectives

Determine the role of DspF and ShcW in T3SS-dependent protein transport in Pca.

Methods

PCR, molecular cloning, directed mutagenesis, DNA sequencing, far-western blotting, two-hybrid screening.

Conclusions

We have found no evidence for chaperone requirement for stability or secretion of the DspE and HrpW effector proteins. On the other hand, introduction of additional copies of the dspF or shcW genes into Pca cells blocks their ability to induce hypersensitive response on nonhost plants. This phenotype is caused by a severe defect in translocation of effectors into plant cells. Translocation of the DspE-Cya fusion appeared to be completely blocked while translocation of AvrPto-Cya – reduced from Pca cells carrying dspF on a plasmid. Thus, additional molecules of T3SS chaperones can block effector translocation, while their native amounts are not required for secretion/translocation. This suggests that the main function of the DspF and ShcW chaperones may be to repress premature effector transport.
Background

Plants are constantly exposed to microbes, such as bacteria, fungi or viruses and most plants are resistant to most microbes. To be successful, a pathogen must overcome constitutive defenses or suppress induced defenses. Our group previously demonstrated that cyclic $\beta$-(1,2)-glucan, a polysaccharide synthesized by Xanthomonas campestris pv. campestris, suppresses the accumulation of callose deposition, as well as the local and systemic expression of PR1, a defense-related gene associated with salicylic acid responses (The Plant Cell. 2007: 2077-2089).

Objectives

To gain knowledge about the mechanism of cyclic glucan in the modulation of plant defense

Methods

We performed Xanthomonas campestris pv. campestris infection assays on A. thaliana leaves pretreated with flagellin (flg22) and purified cyclic glucan or both. Measurement of WRKY22, WRKY33 and MPK3 expression was done by RT-PCR and binding assays was made using Arabidopsis membranes and labeled $^{14}$C-cyclic glucans.

Conclusions

We observed that cyclic glucans of X. campestris suppresses partially the immunity elicited by flg22 in Arabidopsis thaliana. In addition, this compound downregulates WRKY22, WRKY33 and MPK3 expression, early defense genes induced by this elicitor. In order to get further insights into how CG modulates flg22 responses, we performed binding assays with Arabidopsis membranes and labeled $^{14}$C-CG. The radiolabeled glucan, $^{14}$C-CG, bound specifically to wild-type plant extracts, suggesting the existence of direct interaction between CG and an still unknown Arabidopsis receptor. The present results shed light on the mechanism by which Xcc CG hijacks, at least partially, plan immune response.
Background

Citrus canker is one of the most important and aggressive bacterial diseases of citrus trees. The causal agent of this disease is *Xanthomonas citri* subsp. *citri* (Xcc). Biofilm formation on citrus leaves plays an important role in epiphytic survival of Xcc. Previous work from our laboratory described a genetic screen for biofilm formation-defective mutants in Xcc, this work identified that a mutant with a transposon insertion in XAC3733 (xbmR) had significantly reduced attachment to a polystyrene surface (Microbiology. 2013: 159, 1911–1919).

Objectives

To gain knowledge about the regulatory mechanisms of Xcc infection

Methods

For in vitro analysis of biofilm formation, we used confocal laser scanning microscopy. For pathogenicity assays grapefruits leaves was used as the host plant for Xcc. Gene expression levels was obtained by qRT-PCR. Intracellular alteration in c-di-GMP was achieved by exogenous expression of either the diguanylate cyclase WspR19 from *P. fluorescens* or the c-di-GMP phosphodiesterase PA2567 from *P. aeruginosa*.

Conclusions

A knock-out of xbmR led to a substantial downregulation of *fliA*, which encodes a σ28 transcription factor, as well as *fliC* and XAC0350 which are potential member of the σ28 regulon. XAC0350 encodes an HD-GYP domain cyclic di-GMP phosphodiesterase. These findings suggest that XbmR is a key regulator of flagellar-
dependent motility and chemotaxis exerting its action through a regulatory pathway that involves FliA and c-di-GMP.
MICROBIAL RESOURCES TO BIODEGRADE COMPLEX TAR AFTER STEAM GASIFICATION PROCESSES

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Background

Biomasses gasification produces toxic compounds (tar), mainly composed by polycyclic aromatic hydrocarbons (PAH). Up to now the problem of the tar disposal is not solved. Bioremediation of tar with a mixed microbial consortium could be a highly promising and innovative approach for the handling of this by-product.

Objectives

The aim of this study was to assess the tar-degradation process in a laboratory-scale microcosm.

Methods

Triplicate tar-containing microcosms (PAH 13,904 mg/L) were incubated for 20 days with a microbial consortium selected from pinewood and polluted soils. After 20 days HPLC and GC/MS showed a significant reduction of all the PAH compounds to about 150 mg/L whereas PAH concentration in the negative controls was about 8,400 mg/L. Reliability of distinct replicated microcosms was very high since microbial communities showed 97% of similarity. Automated fingerprints and ribosomal gene pyrosequencing showed, at first, a noticeable lag-phase (0-2 days), where fungi were the leading part of the community. A second phase was evident between 3 and 10 days of degradation with specific bacterial genera. After 15 days, a stable phase with 24 bacterial genera and only one fungal genus was detected. Overall, a total of 59 bacterial and 22 fungal genera participated in the degradation.

Conclusions

Bacterial species grew according to a “cross-feeding” behavior. Natural and polluted soils proved to successfully provide valuable microbial taxa for innovative services such as tar degradation.
COMPARISON OF BACTERIOPLANKTON COMMUNITIES AND DIESEL BIOREMEDIATION ABILITY OF NATIVE BACTERIA FROM DIFFERENT MEDITERRANEAN TOURIST PORTS

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Background

Tourist ports are subjected to seasonal impacts determining high pollution by petroleum hydrocarbons. Bioremediation is an emerging clean-up technique based on the addition of amendments or microorganisms able to stimulate the degradation activity of the autochthonous bacteria.

Objectives

This study was carried out within MAPMED, a multidisciplinary project aimed to improve the environmental sustainability of tourist ports in the Mediterranean Sea with regard to hydrocarbon pollution. Cagliari (Italy), El Kantaoui (Tunisia), and Heraklion (Greece) were selected as case study sites. The present study aims to compare bacterioplankton, hydrocarbon degraders, and the effects of different bioremediation additives on the autochthonous communities among the selected sites.

Methods

The structures of bacterioplankton were compared by 16S rRNA T-RFLP among sites, sections with different usage and seasons (winter, beginning and end of the tourist season). Cultivable hydrocarbon degraders were characterized regarding phylogenetic position and catabolic abilities. Different bioremediation treatments (biostimulation with nutrients and bioaugmentation with selected strains) were tested in seawater microcosms supplemented with diesel. The monitoring parameters are the titles of heterotrophs and diesel degraders, and the hydrocarbon removal efficiency. At the end of treatments, the bacterioplankton communities are compared by 16S rRNA gene analysis.

Conclusions

The structure analysis highlights a clear seasonal variation in bacterioplankton and differentiates Heraklion communities from those found in Cagliari and El Kantaoui. Degraders exhibiting different ecological strategies dominate the three communities, copiotrophics in Cagliari and hydrocarbonoclastic bacteria in El Kantaoui and
Heraklion. Comparison of the effects of different bioremediation treatments is currently in progress.
EXTRACELLULAR DNA AS BIOSCAFFOLD FOR THE SYNTHESIS OF SUPERIOR PALLADIUM(0) NANOCATALYSTS HIGHLY ACTIVE IN THE DETOXIFICATION OF PERSISTENT ORGANOHALOGEN POLLUTANTS

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Background
In a search for new aqueous-phase systems for catalyzing reactions of environmental and industrial importance, we prepared novel palladium(0) nanocatalysts in the presence of Shewanella oneidensis strains. Especially a nuclease-deficient mutant of Shewanella oneidensis MR-1 showed an increased tolerance against elevated heavy-metal stress, obviously as a result of high levels of extracellular DNA (eDNA) and additional adaptive response mechanisms.

Objectives
The main objective of our study was to assess whether eDNA is able to protect S. oneidensis cells from toxic Pd(II) ions and can simultaneously serve as a scaffold for nanocatalyst synthesis.

Methods
With eDNA as nucleation site, it was possible to retrieve superior nanocatalysts, bearing small sizes, a monodisperse size distribution and low aggregation behaviour which are important prerequisites for high catalytic activity. Furthermore, in comparison to catalysts produced by the Shewanella oneidensis wildtype, the nanocatalysts synthesized by the mutant were more resistant against catalyst transformation, deactivation and poisoning. The catalytic properties of the palladium(0) nanocatalysts were tested using a newly developed real-time PTR-ToF-MS analytical technique, which is able to follow the nanocatalyst-mediated detoxification of highly-persistent anthropogenic pollutants at environmentally relevant concentrations. The palladium(0) nanocatalysts synthesized in the presence of the nuclease-deficient mutant of Shewanella oneidensis MR-1 showed a high catalytic activity for the dehalogenation of hexachlorobenzene and triclosan.

Conclusions
Further tests with "old" and emerging halogenated persistent organic pollutants are under way and will demonstrate the high potential of Pd(0) nanocatalysts for clean-up of drinking water, aquifers and specific industrial effluents, as well as the suitability of PTR-ToF-MS to assess the respective transformation and detoxification reactions.
COMPARATIVE STUDY OF THE CAPACITY OF CARRIERS: ANOXKALDNES K1, POLYPROPYLENE PADS AND GRANULAR CORKSORB TO FORM HYDROCARBON-DEGRADING BIOFILMS AND REMOVE HYDROCARBONS FROM DIESEL POLLUTED WATER

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Background

Hydrocarbons are a widely distributed pollutant that reaches the environment from multiple sources and involve significant risks to the environment that must be minimized.

Objectives

To design a new treatment system for hydrocarbon contaminated water based on biofilm technology, we studied three different carriers 1) AnoxKaldnes™ (a plastic carrier). 2) Polypropylene pads, these are oil absorbent pads composed of 100% meltblown polypropylene that absorb oil only and not water. 3) Granular thermal treated hydrophobic cork which absorbs oil and solvents without absorbing water.

Methods

Using 1L bioreactor with 400 ml of water and 200 ml of carrier, kinetics of growth and adherence to carriers were determined by enumeration of viable cells. Development of biofilm was visualized by scanning electron microscopy. TPH, n-alkanes, branched-alkanes and naphthalene removal were determined by GC/MS at the beginning and at the end of the assays. Pseudoalteromonas elyakovii strain W18 was inoculated to study biofilm formation. Waters with and without 1% (v/v) diesel, were used to analyze the characteristics of formed biofilm and the efficiency of hydrocarbon removal by indigenous microorganisms.

Conclusions

No inhibitory effect of carriers was observed, even in bioreactors with high hydrocarbon contamination. SEM analyses showed that Polypropylene pads and Granular CorkSorb allow microbial biofilm formation. GC/SM analyses showed that Polypropylene pads not only enhance microbial growth but also retain high amount of hydrocarbons.

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NOVEL ANAEROBE OBTAINED FROM A HEXADECANE-DEGRADING CONSORTIUM
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Background

Aliphatic hydrocarbons (AHC) are abundant in crude oil and fuels, and are frequent contaminants of water, soil and sediments. There is potential for AHC bioremediation using sulfate as electron acceptor, due to its abundance in marine environments and natural presence in soils and groundwater.

Objectives

In this work sulfate-reducing anaerobic microorganisms involved in AHC biodegradation were studied.

Methods

Anaerobic sludge was incubated at 37°C with hexadecane (1mM) and sulfate (20mM) in serum vials. Cultures were successively transferred to fresh medium until a stable enrichment was obtained (monitored by microscopy and PCR-DGGE of 16S rRNA gene). For isolation of AHC-degrading bacteria, serial dilutions and successive transfers are now running using palmitate (1mM) as an easier substrate.

Conclusions

Cultures growing on palmitate show two main bacterial cell types: a rod-shaped bacterium closely related to Desulfomonile limimaris (94% identity) was predominant in the first 30 days of incubation, when 83% of the added palmitate was degraded coupled to 4 mM sulfate reduction (suggesting stoichiometric palmitate conversion to acetate); and an oval-shaped bacterium related to Desulforhabdus amnigena (99% identity) that mainly developed when incubations where extended and a total of 11.5 mM sulfate was reduced. Growth of Desulforhabdus was stimulated when incubated with acetate. The role of the Desulfomonile in AHC degradation will be further discussed in the presentation, as well as its halorespiring ability, a characteristic of
the Desulfomonile genera. Further characterization of this novel bacterium is important due to its high potential for bioremediation of hydrocarbons, fats and halogenated pollutants.
APPLICATION OF HIGHLY EFFECTIVE MICROBIAL DEGRADERS OF OILS AND FATS IN WASTEWATERS DECONTAMINATION

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Background

Disposal of municipal sewage and industrial effluents remains a top priority challenge. Annual discharge of wastewaters containing lipid substances in Belarus reaches 10-12 mln ton. It is recommended to utilize high-efficient microbial strains decomposing oils and fats for their remediation.

Objectives

Searching and examination of microorganisms-degraders of lipid compounds.

Methods

Isolation of microbial strains was carried out by enrichment culture technique. Lipase activity was evaluated by Ota-Yamada method. Dynamics of lipid dissimilation was monitored on model water comprising fats (lard, dairy) and oils (sunflower and olive) in concentration 1000 mg/l as the sole carbon sources.

Conclusions

Four superactive bacterial strains *Rhodococcus ruber* 2B, *Rhodococcus sp.* R1-3FN, *Bacillus subtilis* 6/2-APF1, *Pseudomonas putida* 10AP – degraders of fatty substrates showing lipase activity levels in the range 0.65-0.70 u/mg protein were derived by adaptive selection method. It was found that the selected strains were capable to consume 83.4-94.2 % of animal fats and 82.3-90.4 % of vegetable oils in 7 days. Nutrient medium composition was defined and optimized in terms of nitrogen, potassium, phosphorus and carbon sources. It allowed to attain maximum biomass concentrations (3.0-4.8 g/l by 48 h of fermentation) with elevated lipase activity.

Based on the obtained data technology of manufacturing microbial preparation to accelerate disposal of lipid pollutants in wastewaters was developed.
Introduction of new microbial product into biodecontamination scheme promoted 15.9 % decline in chemical oxygen demand, increased degree of lipid removal by 59 % and improved quality of activated sludge.
APPLICATION OF CARBON STABLE ISOTOPE ANALYSIS (CSIA) TO INVESTIGATE BIODEGRADATION AND DIRECT PHOTOLYSIS OF ANTIBIOTIC SULFAMETHOXAZOLE

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Background

Sulfamethoxazole (SMX) represents sulfonamide antibiotics (SAs) widely used in veterinary and human medicine. SMX is incompletely metabolized and enters the environment with wastewater discharge, animal manure and aquaculture. SMX is frequently detected in wastewater treatment plants, surface and ground water. In the environment it undergoes different degradation processes, such as biodegradation and photodegradation.

Understanding of SA’s removal routes is crucial for an assessment of their environmental impact. Compound specific stable isotope analysis (CSIA) can provide additional information on the organic contaminants’ transformation processes in complex environments. In CSIA, the changes in isotope composition of the parent compound are monitored during (bio)transformation processes and the isotope enrichment of the investigated contaminant provides an evidence for its (bio)degradation without a need of metabolite analysis.

Objectives

This research aims to evaluate the applicability of CSIA for the assessment of SMX transformation pathways. Therefore, the isotope fractionation of SMX during biotic degradation by Microbacterium sp. strain BR1 and abiotic transformation via direct photolysis was determined.

Methods

Bio- and photodegradation assays; HPLC; LC-IRMS; quantification of enrichment factors

Conclusions
A significant difference in isotope fractionation during biotic and abiotic SMX decomposition was observed, showing that CSIA has a potential for distinguishing these two degradation processes. Isotope fractionation during direct photolysis was variable and depended on pH of the solution.

This work shows a new application for CSIA - monitoring of sinks, sources and environmental behaviour of water-soluble contaminants, among which pharmaceuticals are of particular concern.
FEMS-2550
Pollutant degradation

BIOREMEDIATION OF DRILL CUTTINGS BY HALOPHILIC BACTERIA
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Background

Exploration and drilling of oil requires the use of drilling fluid to lubricate and cool the drill bit. Oil based muds (OBMs) containing diesel, are preferred drilling fluids. After mixing with drill cuttings, OBMs form complex industrial wastes containing hydrocarbons and brine. Bioremediation is a promising alternative to physicochemical methods for cleaning up drilling wastes.

Objectives

The goal of this study was to isolate halophilic bacterial consortium capable of degrading diesel oil in drilling muds and evaluating their applicability in the lab scale drill cuttings microcosm.

Methods

Between 10 microbial consortia isolated from different saline environments, the best consortium was isolated from Qom sample which grew very well on Bushnell Haas medium containing diesel with the bacterial count about $1.9 \times 10^{15}$ CFU/ml after 96h of incubation. For evaluating the applicability of the consortium in bioremediation of drill cuttings, it was mixed by 50% of fine sands or biological active soil in the lab scale microcosms and the effect of consortium inoculation was compared with corresponding un-inoculated control for three month.

Conclusions

The Measurement of the amount of remaining pollutant (TPH) after 3 month showed that the most degradation rate (40%) belonged to the microcosm containing 50% fine sand and consortia inoculation. Sequencing of 16S rRNA gene revealed that the dominant strain of the consortium belonged to the Dietzia genus. It could be concluded that bioaugmentation is useful for remediation of drill cuttings when the cuttings mixed with fine sand which has a weak microbial flora and allowing the better growth of inoculated consortium.
FEMS-1548
Pollutant degradation

ISOLATION AND CHARACTERIZATION OF CONFLUENTIMICROBIUM SP. NS6, A NEW NAPHTHALENE-DEGRADING BACTERIUM, ISOLATED FROM AN OIL-CONTAMINATED TIDAL FLAT

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Background
Polycyclic aromatic hydrocarbons (PAHs) are great environmental concerns because of their persistence, toxicity, and carcinogenicity. Many studies for bacteria capable of PAH degradation have been reported in seawater, but PAH-biodegrading bacteria in tidal flats have been rarely explored.

Objectives
The purpose of this study was to isolate strains NS6 as a novel PAH degrader capable of PAH degradation in tidal flats and characterize its physiological and PAH-biodegrading properties.

Methods
We established a slurry type enrichment system based on tidal flat sediment and seawater using naphthalene as a sole carbon and energy source. The enrichment culture was spread on marine agar (MA) and incubated at 25°C for 2 days. Colonies grown on MA were characterized phylogenetically based on their 16S rRNA gene sequences and the naphthalene degradation abilities were evaluated in seawater, tidal flat slurry, and ONR7a. Finally, a strain showing a good naphthalene degradation property was isolated and its PAH degradation properties and genes were characterized.

Conclusions
A naphthalene degrading bacterium, designated strain NS6, was isolated from the slurry type enrichment and classified into a member of Confluentimicrobium based on its 16S rRNA gene sequence. Confluentimicrobium sp. NS6 degraded PAH compounds quickly in all tested media. Naphthalene dioxygenase gene of Confluentimicrobium sp. NS6 was successfully amplified by only NDO 201 primer set, which suggested that strain NS6 may metabolize naphthalene via a nah catabolic pathway alike Rhodococcus. In addition, the PAH degradation properties, physiologies, and PAH degradation gene structure of strain NS6 will be investigated and discussed in detail in the poster section.
Background
The use of white rot fungi such as *Pleurotus ostreatus* and their lignin-modifying enzymes has become an effective treatment for various organic soil and water pollutants. The manganese peroxidase gene family (*mnps*) is a major part of the ligninolytic system of *P. ostreatus*. This gene family is comprised of nine genes encoding short manganese peroxidases (short-MnPs) or versatile peroxidases (VPs). The VPs contain unique active sites which are responsible of direct oxidation (in the absence of Mn^{2+}) of various aromatic compounds.

Objectives
Here we study the oxidation mechanisms of aromatic compounds by VP1 of *P. ostreatus*.

Methods
We show that in Mn^{2+}-deficient GP medium *vp1* (encoding VP1) has a key and non-redundant function. We used the azo-dye Orange II (OII) as a model contaminant, its decolorization occurs only during the idiophase where the abundance transcripts of *mnps* indicate that *vp1* is the predominantly expressed and a Δ*vp1* strain showed a drastic reduction in the decolorization. Three degradation metabolites were identified by LC-MS indicating both asymmetric and symmetric enzymatic cleavage of the azo bond. The presence of asymmetric cleavage diminishes the toxicity level of the degradation products. To better understand mechanisms of degradation and detoxification we purified and characterized VP1. The purified enzyme degraded 60% of OII (50µM) within the first min and continued up 90% within 6 min.

Conclusions
The non-specific oxidation properties of VP1 and its unique ability to degrade organic compound in the absence of Mn^{2+} suggest that it may have potential applications for treatment of contaminated water.
Background

Accumulation of synthetic polymers in the environment is cause for concern, as they have been associated with various negative effects on ecosystem health, especially marine systems. Knowledge on biodegradation pathways of conventional polymers is very limited, but it is known that these compounds are hardly biodegradable.

Objectives

Here, we investigated whether fungi causing white and brown rot of wood are capable of degrading the widespread polymer polystyrene (PS) as well as its water-soluble analogue polystyrene sulfonate (PSS). Their well-known extracellular oxidation capabilities made them promising candidates for the degradation of these recalcitrant substrates.

Methods

PSS analysis by size exclusion chromatography revealed that it was practically inert to white-rot fungi, but vulnerable to various brown-rot strains, which caused strong depolymerisation leading to molecular mass reductions of more than 90 %. Detailed investigations pointed to Fenton reactions driven by fungal hydroquinones as agents of depolymerisation. Solid PS films were significantly more recalcitrant, as no changes in their bulk properties were observable after long periods of fungal treatment. Indications of biodegradation only came from surface property measurements, which revealed increased hydrophilicity through slightly decreased water contact angles. Investigations by X-ray photoelectron spectroscopy revealed that surface oxygen content had increased significantly in fungi-treated samples.

Conclusions
These results demonstrate that biodegradation of recalcitrant synthetic polymers is, in principle, possible. However, the different observations between PS and PSS indicate that biological attack on the solid polymer proceeds extremely slowly, raising questions about the feasibility of PS waste biodegradation.
FUNCTIONAL ANALYSIS OF A PLASMID-BORNE ALKYL SULFATE DEGRADATION MODULE OF PSYCHROBACTER SP.

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Background

Plasmid pP62BP1 (34.5 kb), harboured by strain Psychrobacter sp. DAB_AL62B isolated from ornithogenic deposits in Spitsbergen (Svalbard Archipelago, Norway), was found to carry a phenotypic module, named SLF, which consists of four open reading frames (slfCHSL) encoding putative catabolic enzymes and a gene (slfR) for transcriptional regulator of AraC/XylS family. Based on the comparative in silico analyses, the module was predicted to take part in the metabolism of alkyl sulfates, e.g. sodium dodecyl sulfate (SDS), a popular anionic surfactant.

Objectives

The aim of our work was to perform a functional characterization of the SLF module both in native host and in other bacterial strains as well as to elucidate the regulatory mechanism of the slfRCHSL genes expression.

Methods

Qualitative and quantitative SDS degradation assays were designed based on the properties of Stains-All, a carbocyanine dye. Promoter strength was examined in β-galactosidase activity assays. Transcriptional organization of the module was characterized with the use of reverse transcription PCR.

Conclusions

Growth experiments revealed that 0.17 mM concentration of SDS in the medium is sufficient to exert bacteriostatic effect on several tested Psychrobacter spp. strains. Nonetheless, the pP62BP1 containing strain was capable of the complete SDS degradation in these conditions. The cloning of the whole SLF module and its slfR-deficient derivative into Escherichia coli TG1 resulted in obtaining transformants able to degrade SDS. The molecular studies showed that the activity of the slfCHSL operon is dependent on SifR protein which acts as a negative transcriptional regulator.
MOLINATE BIOREMEDIATION STRATEGY USING A MICROENCAPSULATION APPROACH

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Background

Molinate is a thiocarbamate herbicide used in rice cultivation to control grass weeds, however its use had led to environmental contamination, being critical the establishment of remediation procedures. A mixed culture has been described as being able to mineralize molinate as the sole source of carbon, nitrogen and energy [1, 2], primarily, by the activity of molinate hydrolase (MolA) [3], encoded by Gulosibacter molinativorax ON4T. Previous studies, using soil microcosms, have demonstrated that a bioaugmentation strategy using the mixed culture was a feasible approach to bioremediate paddy soils contaminated with molinate [4].

Objectives

To improve the bioremediation strategy, we aimed at microencapsulating ON4T cells or MolA, using a spray-drying process that uses biopolymers as encapsulating agent [5].

Methods

Several encapsulating agents (sodium and calcium alginate, arabic gum, chitosan and modified chitosan) were used. The spray-drying product yield was about 50% for all the agents. The metabolic activity of the microencapsulated cells was proven by a qualitative colorimetric method using a redox dye (tetrazolium violet).

Conclusions

There was a linear correlation between the amount of encapsulated cells and the colour development intensity. The feasibility of using optimized microencapsulated biomaterial will be tested in contaminated soil and water using a microcosm approach.


INACTIVATION OF PSEUDOMONAS AERUGINOSA BY ZINC OXIDE NANO PARTICLES IN AQUEOUS SOLUTION

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Background

With the emergence of microorganisms resistant to multiple antimicrobial agents there is increased request for promotion of disinfection methods.

Objectives

Since ZnO nanoparticles (ZnO-NPs) exhibit strong antibacterial activities on a broad spectrum of bacteria the aim of this study was to evaluate the antimicrobial activity of ZnO-NPs against pseudomonas aeruginosa as a model for gram-negative bacteria.

Methods

Muller Hinton broth was used as a growing medium for pseudomonas aeruginosa. Photocatalytic experiment was carried out in a laboratory-scale batch reactor with low pressure ultraviolet irradiation (380 nm). Different experimental parameters such as amount of ZnO-NPs, contact time, inorganic and organic substances and pH on photocatalytic inactivation of pseudomonas aeruginosa cells have been studied. An initial pseudomonas aeruginosa concentration of $10^8$ CFU/mL was used for all experiments.

Conclusions

Result showed that, almost all the initial pseudomonas aeruginosa cell ($10^8$ CFU/ml) was inactivated in 60 min in the presence of 2 g/l ZnO-NPs. Photocatalytic inactivation of bacteria was found to follow first order kinetics. The initial pH of the water did not play an important role on the inactivation rate within a range of 6–8 pH units. The amount of photocatalyst also plays an important role in photocatalytic inactivation rate. As the result showed increasing the photocatalyst amount provided more rapid inactivation. Addition of some inorganic ions to the suspension affects the sensitivity of pseudomonas aeruginosa and caused to retard the inactivation rates.
Since the sensitivity of *pseudomonas aeruginosa* to photocatalytic treatment was fairly good, it is therefore, recommended to use this nano-particle for water treatment.
ADAPTATION TO AROMATIC HYDROCARBONS INVOLVES A NEW HYBRID TWO-COMPONENT REGULATORY SYSTEM IN AZOARCUS SP. CIB

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Background

The catabolic processes performed in anaerobiosis are becoming of great environmental interest because many anoxic ecosystems are frequently contaminated with aromatic compounds. The facultative anaerobe β-proteobacterium Azoarcus sp.CIB has been used as model system to study basic aspects on the biochemistry and genetics underlying the anaerobic metabolism of several aromatic compounds. Within the operon responsible for the anaerobic degradation of toluene, we identified an orphan gene, tolR, without homologues. The TolR protein shows an atypical architecture which corresponds to a new hybrid two-component system (HTCS), where the sensor histidine-kinase and the c-di-GMP phosphodiesterase-containing response regulator (RR) components are fused in a single polypeptide.

Objectives

The main objective was to demonstrate that TolR constitutes a new HTCS involved in anaerobic adaptation to aromatic hydrocarbons in Azoarcus sp.CIB

Methods

An in vivo approach for confirming the c-di-GMP phosphodiesterase activity of the tolR gene was done using a lacZ-based P. aeruginosa reporter strain. In vitro approaches by using purified TolR and truncated forms were used to demonstrate toluene-induced autokinase and transphosphorylation activity. Transcriptomic studies were done to identify genes involved in the TolR-mediated response to aromatic hydrocarbons.

Conclusions

TolR encodes a unique HTCS whose RR is not a transcriptional regulator but a c-di-GMP phosphodiesterase. We demonstrate for the first time the effector-dependent induction of the intramolecular phosphorelay in HTCS. Transcriptomic studies confirmed that TolR is involved in a signalling network that controls the morphological, metabolic and stress response programs. The toluene-dependent control of c-di-GMP levels represents an unprecedented mechanism of bacterial adaptation to aromatic hydrocarbons.
BIODEGRADATION POTENTIAL OF PENICILLIUM SP. ISOLATED FROM SOUTH CHINA SEA

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Background
Some halophiles fungal genera were found from sea sediment samples are potentially play an important role in nutrient regeneration cycles as decomposers.

Objectives
In this study, three genera of microfungi derived from South China Sea were isolated as Penicillium, Aspergillus and Talaromyces in order to identify the fungal saprotrophic properties.

Methods
For the first stage of investigation by using dermatophyte test medium (DTM), the results showed that only the ascomycete *Penicillium* sp. highly potential as saprophytic microfungi. This fungus was used to observe their early thallus development *in vitro* and to determine a degradation potential of the textile dyes Trypan blue (TB), Methyl orange (MO), Sudan III (S3), Cango red (CR) and Bromocresolgreen (BG) in broth culture.

Conclusions
*Penicillium* sp. displays a distinctive morphological growth, produced an orange pigment in all different media as an additional features of the strain. In contrast, there were significant differences in physiological tests of *Penicillium* sp. such as thermophilic osmomophilic, pH, salinity, xerotolerant and nitrogen stress. In the presence of textile dyes in liquid media was able decolorized by *Penicillium* sp. Thus, this study shows that halophilic *Penicillium* sp. from sediment of South China Sea is hereby reported for the first time and will be used as bio-indicator of water pollution and also in biological wastewater treatment. In additional this investigation also demonstrated the need of extra and intracellular enzymes associated to decolorization and biodegradation assays.
Background

Biogas resultant from the anaerobic treatment of sludge is composed mainly of CH$_4$ and CO$_2$. H$_2$S, a minor component, is of particular concern due to its corrosive and environmentally hazardous properties. Some microorganisms are able to perform enzymatic oxidation of H$_2$S to sulfur and/or sulfate. Biological treatment using these microorganisms can be an alternative to physical/chemical processes, avoiding the production of secondary waste.

Objectives

This work aims at the characterization of enzyme(s) responsible for H$_2$S oxidation in Pseudomonas sp.

Methods

Isolates A9, B9 and C1, identified as Pseudomonas spp., were isolated through enrichment from wastewater treatment plant deodorant bioreactor supplied with H$_2$S streams. These strains were screened for the presence of genes responsible for H$_2$S oxidation by PCR using degenerated primers for group II sulfide quinone:oxidoreductase. Fragments with the expected size were purified from the gel and subjected to sequencing. Translated sequences were compared using the BLAST software. Total protein profiles were screened by SDS-PAGE, before and after the addition of H$_2$S. Activity tests were conducted using crude cell extract as catalyst and measuring the formation of sulfate as the reaction product.

Conclusions

The amplified fragment from isolate A9 showed 51% identity with an oxidoreductase from Pseudomonas sp. M1, indicating that this bacterium has the machinery required to the desired activity. The protein profile of the strains when grown in the presence H$_2$S differ from the profile of the same strain grown without sulfide.
Background

The biodegradability of PBAT (poly(butylene adipate-co-butylene terephthalate)) under aerobic conditions has been examined in several studies. However, there is considerably less known about PBAT biodegradation in anaerobic environments.

Objectives

Amongst others Clostridium species are known to be typically present during naturally occurring anaerobic degradation processes of polymers (e.g. in biogas plants). Hence, various Clostridium species were in-silico screened for esterases that are possibly active on synthetic polyesters like PBAT.

Methods

First, a selection of identified hydrolases from Clostridium species were successfully expressed in E. coli BL21-Gold(DE3). The activity of these esterases was confirmed on the soluble standard substrate p-nitrophenyl butyrate as well as on the PBAT polymer. All esterases were characterized in detail including the determination of their crystal structure or modeling of the proteins. The crystal structure of one promising esterase revealed the presence of a metal ion that lies deep beneath the protein surface. Furthermore, Clostridium cultures were tested for their ability to express these active and naturally occurring enzymes in situ and degrade the PBAT polymer.
Conclusions

We were able to prove that the novel hydrolases from *C. botulinum* and *C. hathewayi* hydrolyze the aliphatic-aromatic polyester PBAT. These enzymes are promising candidates for industrial applications or pollutant degradation. Furthermore, the study provides information about these anaerobic strains and their naturally occurring enzymes.
USE AND IMPACT OF MNO2 MICROPARTICLES FOR SULFIDE SCAVENGING IN MICROBIAL BROTHS

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Background
Hydrogen sulfide (HS⁻) is ubiquitously present in many waste streams originating from industrial activities as well as in sewage. It needs to be removed, as it can be toxic to bacteria, induces concrete corrosion and forms an odour nuisance. MnO₂ has been proposed for HS⁻ adsorption and oxidation, the addition of such particles to microbial broths may enable selective sulfide scavenging. However potentially negative effects of addition of MnO₂ microparticles (MPs) on microbial metabolism may occur.

Objectives
Therefore, we evaluated the efficiency of this MnO₂ MPs based HS⁻ removal strategy and test the effect of the particles and the sulfide scavenging on bacterial growth and microbial community activity.

Methods
Abiotic batch tests were set up to test impact of pH on HS⁻ removal. HS⁻ removal rate after 24h at pH 7.2 was twice that at pH 10.5. Scanning electron microscopy with energy dispersive X-ray spectroscopy showed the presence of sulfur as well as MnS on the particle surface. The effect of HS⁻, MnO₂ MPs and mixtures of both on the growth of single bacterial strains, and mixed microbial community activity were assessed using an optical density microplate reader placed into an anaerobic chamber and biochemical methane potential assays respectively.

Conclusions
The presence of MnO₂ MPs did not affect bacterial growth under any of the conditions tested. Inhibitory effects due to presence of HS⁻ on bacterial growth were observed, upon addition of MnO₂ MPs normal growth could be restored. We finally observed that a mixture of HS⁻ and MnO₂ MPs positively impacted methanogenic activity.
IN STREPTOMYCES, STRONG ANTIBIOTIC PRODUCTION CORRELATES WITH EXTENSIVE STORAGE LIPIDS DEGRADATION AND HIGH ATP GENERATION.  

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Background  

Some Streptomyces strains produce antibiotics and others don't. Why? For instance S. lividans and S. coelicolor are phylogenetically closely related strains possessing the same functional antibiotic biosynthetic pathways but S. lividans produces antibiotics at low level whereas S. coelicolor produces them at high levels. Interestingly, a dramatic overproduction of antibiotics occurred in S. lividans upon interruption of the ppk gene. This mutant strain experiences energetic stress (ATP deficit) since Ppk regenerates ATP from ADP and polyphosphate. However, the role of the Ppk in relation to the regulation of antibiotic biosynthesis remains unclear.

Objectives  

To better understand it we assess the global impact of strong antibiotic production on the cellular metabolism.

Methods  

To do so proteomic and lipidomic analysis were conducted and the cellular content in polyphosphate, ATP/ADP and free Pi was assayed throughout growth of the three strains of interest grown in phosphate limitation or proficiency.

Conclusions  

Our results revealed that an active degradation of storage lipids (TriAcylGlycerol, TAG) was taking place in the antibiotic producing strains. TAG degradation generates precursors used directly or indirectly for antibiotics biosynthesis as well as numerous reduced co-factors (FADH2) whose re-oxidation by the respiratory chain yields high levels of ATP. Whereas the ppk mutant likely mobilizes its storage lipids to re-
establish its energetic balance, the reason of the extensive TAG degradation taking place in *S. coelicolor* is still not understood.

References

Le Maréchal P *et al.* (2013) Comparative proteomic analysis of the wild-type and the *ppk* mutant of *S. lividans* revealed the importance of storage lipids for antibiotic biosynthesis. AEM 79-19:5907-5917.
Background

Chlorendic acid (1,4,5,6,7,7-hexachlorobicyclo[2.2.1]hept-5-ene-2,3-dicarboxylic acid, known as HET acid, or Hetron 92) has been identified in the past as the dead-end product of the biodegradation of the cyclodiene pesticides Aldrin, Endrin, Dieldrin, Endosulfan, Chlordane, and Heptachlor. On the other hand, this compound is still being produced as a flame retardant, and used in respective materials and corrosion-resistant equipment. Chlorendic acid has been considered as non-biodegradable until today.

Objectives

Although the hexachlorocyclopentadiene moiety of the compound is highly halogenated and does not carry any hydrogen atom, its dichloroethene substructure may provide the side of bacterial attack by the mechanism of anoxic dehalogenation as known from tetrachloroethylene or polyhalogenated aromatics (halo-respiration), and thus represent the initial side of a biochemical reaction. Further steps of dehalogenation then may proceed aerobically as, for instance, in the catabolic pathway of hexachlorocyclohexane (HCH, Lindane).

Methods

We started enrichment experiments with sediments from the River Elbe (Hamburg, Germany) the former contamination of which by pesticides from production plants of chemicals has been well documented, and established a gradient bioreactor system with an anaerobic zone at the bottom towards an aerobic zone at the top. From the latter we could already isolate the dominant bacterium responsible for the mineralization of the (almost) dehalogenated carbon backbone. 16S rDNA analysis revealed this species as 98% identical to Pseudomonas aeruginosa.

Conclusions

Analyses of the 16S rRNA gene sequences in the anoxic and transition zones will determine the bacterial community structure involved in the catabolic sequence. Identification of intermediates of the combined anaerobic-aerobic pathway is underway.
MICROBIAL DEGRADATION OF EMERGING TRACE ORGANIC CONTAMINANTS: SULFAMETHOXAZOLE, PIRACETAM AND RITALINIC ACID

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Background

Contamination with pharmaceuticals is an emerging environmental problem. Most pharmaceuticals are only partially eliminated in biological wastewater treatment processes, leading to contamination of ground- and surface water. Among them, antibiotics are of particular concern due to potential spread of antibiotic resistance. Sulfamethoxazole (SMX) represents sulfonamides, which belong to most prescribed and consumed antibiotics worldwide. Only a few microbial strains using SMX as a sole carbon source have been isolated and metabolic pathways for most of them have not been described.

The nootropic drugs are a new group of emerging contaminants, very popular these days because of their increasing use as pharmaceuticals and illicit drugs. Some of them, e.g. piracetam and ritalinic acid, have been recently detected in drinking water but so far their potential biodegradation has received very little attention.

Objectives

This research aims at isolation and characterization of novel microbial strains capable to degrade SMX, piracetam and ritalinic acid.

Methods

Enrichment and isolation of strains was carried out under aerobic conditions using various environmental samples as inocula: wastewater, activated sludge, soil, birds feces etc. Degradation is monitored by HPLC-DAD; new method for ritalinic acid and piracetam have been established; 16S rRNA analysis was performed; radio assays with 14C labeled ritalinic acid will be carried out.

Conclusions
Two novel strains capable of SMX degradation have been isolated. Enrichment cultures with piracetam and ritalinic acid are active and the isolation of strains is ongoing. It is a first attempt to obtain the strains growing on nootropic drugs.
DEGRADATION OF TRICLOSAN AND TRICLOCARBAN BY A WASTEWATER MICROORGANISM
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Background
Since last several decades, biocides like triclosan and triclocarban have been used as a common ingredient in many synthetic products especially in industrial and personal care products. Hence, the widespread use of triclosan and triclocarban has raised major concern over their impact on eco-geological system.

Objectives
We have isolated and identified a triclosan-degrading bacterium from wastewater treatment plant (WWTP) named as Sphingomonas sp. strain YL-JM2C.

Methods
Response surface methodology was used to understand the suitable conditions for the degradation of triclosan in strain YL-JM2C. The bacterium can efficiently degrade triclosan (5 mg L\(^{-1}\)) with biomass between 0.1-0.3 g L\(^{-1}\) at 30-35 °C with pH 7-8. By GC-MS analysis, the intermediates of triclosan biodegradation were identified as 2,4-dichlorophenol, 2-chlorohydroquinone and catechol. Tracking experiment using \(^{13}\)C labeled triclosan confirmed that the \(^{13}\)C labeled triclosan was completely mineralized into carbon dioxide by strain YL-JM2C, and the phospholipid fatty acids (PLFAs) extracted from bacterial cells showed that part of labeled carbon from \(^{13}\)C labeled triclosan was incorporated into the strain’s fatty acids.

Conclusions
These results indicated that the bacterium was able to utilize triclosan as a carbon source. We also observed that the bacterium was also able to degrade 30 to 35% of triclocarban (4 mg L\(^{-1}\)). In Sphingomonas sp. strain YL-JM2C, triclocarban was initially hydrolyzed into 3,4-dichloroaniline and 4-chloroaniline. 3,4-Dichloroaniline was further transformed to 4-chlorocatechol via 4-chloroaniline. As per our knowledge, this is the first report of bacterial degradation of triclocarban by a pure isolated bacterial strain.
COMPARATIVE AND TRANSCRIPTIONAL ANALYSIS OF THE PREDICTED SECRETOME IN THE LIGNOCELLULOSE DEGRADING BASIDIOMYCETE PLEUROTUS OSTREATUS

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Background

Fungi interact with their environment by means of secreted proteins to obtain nutrients, elicit responses and modify their surroundings. Consequently, lifestyle influences the set of fungal secreted proteins.

Objectives

To perform a combined bioinformatics-transcriptomics study of the secretome of the basidiomycete Pleurotus ostreatus and to explore the conservation of these secreted proteins across the Basidiomycota phylum

Methods

We identified bioinformatically the set of secretable proteins in two monokaryotic strains (haplotypes) of the white-rot basidiomycete P. ostreatus (PC9 and PC15) using the web pipeline SECRETOOL [1]. Then, we performed two RNA-seq analyses to study the relationship between the functional profile of the predicted secretome and the expression level of each group. Finally, we used the set of proteins secreted by P. ostreatus as a query to search for similar proteins in the fungal genomes released in JGI Mycocosm[2]

Conclusions
538 and 554 protein models were predicted to be secreted (4.41% and 4.77% of PC9 and PC15 gene models, respectively). The functional annotation of these proteins revealed the unknown (37.2%), glycosyl hydrolases (26.5%) and red-ox enzymes (11.54%) as the main functional groups, in a similar distribution for the two strains. The expression level of these groups further enhances the relevance of the unknown group and was significantly different in the two strains (revealing different responses to the same environment). Furthermore, the presence of similar proteins to *P. ostreatus* secreted proteins in other basidiomycetes was used to cluster them into

![Figure 1: Dendrogram showing several basidiomycetes arranged by presence of similar proteins (E-value<e-20) to PC15 predicted to be secreted proteins.](image-url)
groups coherent with their particular lifestyles rather than with their corresponding phylogenetic positions.
NEW INSIGHTS INTO THE T6SS OF PSEUDOMONAS AERUGINOSA

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Background

*Pseudomonas aeruginosa* is a gram-negative opportunistic pathogen capable of living in a wide variety of environmental niches and is the third most common cause of all hospital-acquired infections. *P. aeruginosa* utilises a range of secretion systems and virulence factors to enable it to prosper. The Type VI Secretion System (T6SS) is one such factor. This nanomachine can modulate invasion of epithelial cells and is widely known for its ability to facilitate the delivery of effector proteins into both eukaryotic cells and bacterial competitors.

Objectives

The majority of *P. aeruginosa* strains have three complete T6SS clusters [H1-, H2-, and H3-T6SS] however additional ‘orphan’ genes or short operons are present in the *P. aeruginosa* genomes, which have the characteristics of T6SS components. In this study we have employed a range of molecular microbial techniques to further investigate components of the T6SS and related systems.

Methods

Through the construction of *lacZ* fusions, transposon mutagenesis, specific gene deletions, complementation and labelling with V5 tags we have identified new secreted products and mechanistic insights that contribute to the function of the *P. aeruginosa* T6SS. We also characterized global regulatory networks that are instrumental to the expression of genes encoding either core components of the T6SS machine or cognate T6SS effectors.

Conclusions

Overall secretion systems are of vital importance for Bacteria to interact with their environment and cause infection. A detailed knowledge of the mechanisms of secretion combined with an understanding of the role of the secreted factors should provide the basis for the development of new therapies to combat bacterial infections.
EXPRESSION OF THE MENINGOCOCCAL AUTOTRANSPORTER AUTB AND ITS IMPACT ON BIOFILM FORMATION

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Background
Autotransporters are secreted proteins of Gram-negative bacteria often involved in virulence. They consist of a translocator domain that inserts as a β-barrel in the outer membrane and a fused passenger domain that is translocated across the outer membrane. Genomes of the human pathogen Neisseria meningitidis contain eight autotransporter genes. Seven have been characterized to a certain extent, but the autotransporter AutB remains poorly studied. Previous studies suggested that the autB gene is a pseudogene.

Objectives
Our objective was to analyze whether AutB can be expressed and to elucidate its possible function.

Methods
In this study we used a combination of genomic and proteomic assays to determine the expression of the autotransporter. Included were also diverse functional assays to analyse its function.

Conclusions
Bioinformatics analysis of available genome sequences indicated that AutB expression is prone to phase variation, and can be inhibited by the presence of premature stop codons or other genetic disruptions. However, several genome sequences contain an intact autB gene. Western blotting and RT-PCR assays demonstrated the expression of AutB in strains with an intact gene. Proteinase K-accessibility assays evidenced that AutB is secreted but its passenger remains attached to the cell surface. Functional assays revealed the involvement of AutB in biofilm formation in a strain-dependent mode. We conclude that autB is a pseudogene only in some strains and that AutB may have a relevant function in the formation of bacterial communities.
THE OLOGOMERIZATION STATE OF THE TYPE III EFFECTOR/CHAPERONE COMPLEX CAN PROMOTE ITS RECOGNITION BY THE TYPE III SECRETION SYSTEM IN SALMONELLA.

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Background

Many bacterial pathogens use Type Three Secretion System (T3SS) to inject virulence factors, named effectors, directly into the cytoplasm of target eukaryotic cells. Before being translocated, effectors are maintained in a secretion-competent state by interacting with specific chaperones. The molecular mechanism of the recognition process between the effector/chaperone complex and the T3SS is still unclear but a highly conserved hexameric-ring-shaped ATPase localized at the basis of the needle complex has a critical role in the specific recognition process of chaperone-effector complex.

Objectives

As previous works showed that chaperones and effectors share structural motifs that could be at the origin of the recognition by TTSS, we investigated the structure of the complex between SopB, and its cognate chaperone SigE that is the form recognized by TTSS.

Methods

In this work we performed a biochemical and structural characterization of Salmonella SopB/SigE chaperone/effector complex by SAXS.

Conclusions

Our results showed that the SopB/SigE complex is assembled in dynamic homo-hexameric-ring-shaped structures with an internal tunnel. In this ring, the chaperone maintains a disordered N-terminal end of SopB molecules, in good position to be reached and processed by the T3SS. This ring dimensionally fits the ring-organized molecules of the injectisome including ATPase hexameric ring; this organization suggests that this structural feature is important for the ATPase recognition by T3SS. As effectors share neither sequence nor structural identity, the quaternary oligomeric structure could constitute a strategy evolved to promote the specificity and efficiency of T3SS recognition.
Background
VirB4-like proteins are associated with all bacterial Type 4 Secretion Systems (T4SS) described to date. These signature ATPases function in assembly of the T4SS channel and biogenesis of extracellular pili. They are also required for translocation of secretion substrates and nucleoprotein uptake during pilus-mediated phage infection.

Objectives
Very little is known about the regulation of VirB4-like ATPases in protein trafficking. Our paradigm to study control of VirB4-like ATPases is TraC of the F-like conjugative plasmid R1. Here we test the hypotheses that ATPase activity is controlled by (i) oligomeric state and (ii) intra-molecularly, by auto-inhibition.

Methods
Full length and truncated alleles of traC were modified with a strep-tag and evaluated functionally by complementation of conjugative transfer of a ΔtraC derivative of R1. Multimer formation, nucleotide binding and hydrolysis by purified TraC proteins were measured under various conditions in the presence and absence of protein interaction partners using NMR and biochemical techniques.

Conclusions
Full length TraC behaves as a monomer in solution with and without a 20-fold molar excess of non-hydrolysable ATP. This form binds but does not hydrolyze nucleotide. Preliminary data indicates that proteolytic processing of the protein stimulates both the formation of apparent hexamers and enzyme activity.

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Background

The type VI secretion system (T6SS) involves ~13 core proteins and shares similarities with the tail of the bacteriophage T4. The current model of the structural organization of the T6SS suggests that its core components are assembled in several subcomplexes that are associated to form a dynamic phage-like machinery. A first set of T6SS components forms a trans-envelope complex which consists of the two inner membrane proteins TssM and TssL, and the outer membrane lipoprotein TssJ. A second group of T6SS proteins forms the bacteriophage tail-like structure. It includes the components Hcp and VgrG, and the sheath-like structure made of the two proteins TssB and TssC. Hcp proteins are hexameric rings which are able to form a rigid tube capped by the puncturing device made by VgrG proteins. Contraction of the TssB/TssC sheath propels the Hcp-VgrG structure, which ultimately results in the injection of toxins into target cells.

Objectives

The remaining T6SS core components are likely to form what is known as the baseplate structure in the phage. In the T6SS, it connects the tail-like structure to the membrane subcomplex. Functional and structural information about these T6SS proteins as well as their assembly within the machinery remain poorly described.

Methods

Here, we used biochemical and electron microscopy techniques to determine the structure and function of individual components of the T6SS in Pseudomonas aeruginosa. Additionally, using a systematic analysis of protein-protein interaction, we determined the interaction network between T6SS proteins.
Conclusions

Overall these data led to propose a comprehensive working model of the T6SS.
Background
Mastitis, infection and inflammation of the mammary gland, is a well-known problem in the dairy industry, affecting cows worldwide and causing considerable financial losses. Multiple bacterial species can cause mastitis and *E. coli* is often involved. Despite many years of mastitis research, no efficient measures exist to prevent or treat the disease, and only little is known about specific virulence factors of the bacteria.

Objectives
Our goal is to understand the molecular mechanisms of host-pathogen interactions in the mammary gland and relate them to disease processes, in hope that understanding these mechanisms will lead to development of novel tools to combat *E. coli* mastitis.

Methods
Using genome sequencing and analysis of six clinical isolates, we found that type VI secretion system (T6SS) gene clusters are present in all. Furthermore, using unbiased screening of strains for reduced colonization, fitness and virulence in our murine mastitis model, we have identified in P4-NR strain a new pathogenicity island encoding the core components of T6SS and its hallmark effectors Hcp, VgrG and Rhs. Next, we have shown that specific deletions of T6SS genes reduced *in vivo* pathogenicity.

Conclusions
Based on our results we hypothesize that T6SS is an important virulence mechanism of MPEC. To our knowledge, we are the first to describe relevance of T6SS in the pathogenesis of mastitis. We intend to validate our findings in dairy cows and field strains and study the molecular mechanisms of T6SS associated with MPEC virulence. The identified mechanism may provide new targets for diagnostic, preventive and therapeutic intervention.
BACTERIAL KILLING VIA A TYPE IV SECRETION SYSTEM
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Background
Type IV secretion systems (T4SSs) are multiprotein complexes that transport effector proteins and protein-DNA complexes through bacterial membranes to the extracellular milieu or directly into the cytoplasm of other cells. They are essential for host colonization by many medically important microbes as well as for horizontal transfer of genetic material between bacteria and from bacteria to plants. Many bacteria of the family Xanthomonadaceae, that occupy diverse environmental niches, carry a T4SS with unknown function but with several characteristics that distinguishes it from other T4SSs.

Objectives
The aim of this study is to determine the function of the Xanthomonadaceae T4SS.

Methods
We have used structural biology, spectroscopy, enzymology and novel bacterial competition and secretion experiments to address this question.

Conclusions
The Xanthomonas citri (Xac) T4SS provides these cells the capacity to kill other gram-negative bacterial species in a contact-dependent manner. The secretion of one Type IV bacterial effector protein is shown to require a conserved C-terminal domain and its bacteriolytic activity is neutralized by a cognate immunity protein whose 3D structure is similar to peptidoglycan hydrolase inhibitors. This is the first demonstration of the involvement of a T4SS in bacterial killing and points to this special class of T4SS as a mediator of both antagonistic and cooperative interbacterial interactions and therefore an important driving force in the evolution of bacterial species.
STUDIES ON THE REGULATION OF TYPE VI SECRETION SYSTEMS IN ESCHERICHIA COLI

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Background

Type VI secretion systems (T6SS) are complex protein transport machineries and widespread among pathogenic Gram-negative bacteria. Accordingly, one could assume that they may contribute directly or indirectly to pathogenesis; for example, by targeting eukaryotic host cells or by competing with other bacteria in microbial populations.

Objectives

The genomes of the uropathogenic *Escherichia coli* (UPEC) strains often harbor one or more genomic islands encoding putative T6SSs. As the expression of T6SSs seems to be tightly regulated, most of the secretion systems are inactive under standard laboratory conditions. The screening for potential regulators directing gene expression of T6SSs is a first step to get a better understanding under which conditions these secretion systems may be active and contribute to the fitness and/or pathogenicity of *E. coli*.

Methods

To elucidate the transcriptional regulation of the T6SSs in UPEC strains, chromosomal and plasmid-based promoter-reporter gene fusions to core genes of both genomic islands have been constructed. The promoter activity has been tested in different *E. coli* K-12 deletion backgrounds as well as in various wild type *E. coli* strains.

Conclusions

The screening using the promoter-reporter gene fusion demonstrated that the availability of nucleoid-associated proteins and the growth phase, i.e. the stationary phase, affect transcription of genes coding for core elements of T6SSs.
TOWARDS A BETTER UNDERSTANDING OF THE BACTERIAL TYPE II SECRETION PATHWAY
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Background
The bacterial type II secretion system (T2SS) is unique in its ability to promote the transport of large folded and sometimes multimeric proteins. In this secretion process, exoproteins are first translocated into the periplasm. The final release into the medium requires a multiprotein complex called the secreton.

Objectives
Although the 12 individual components of the secreton have been identified, its mode of action remains obscure.

Methods
We set up various dedicated in vitro and in vivo protein-protein interaction experiments to identify the Pseudomonas aeruginosa Xcp T2SS periplasmic interactome.

Conclusions
BIAcore experiments revealed that three Xcp components, XcpP, the secretin XcpQ, and the pseudopilus tip, directly and specifically interact with secreted exoproteins. Affinity chromatography co-purification indicated that the XcpY periplasmic domain interacts with the secreted substrate and a component of the pseudopilus tip XcpW. Interestingly, the periplasmic domain of another member of the Xcp inner membrane platform, XcpZ co-elutes with the XcpY/substrate and the XcpY/XcpW complexes during affinity chromatography. Finally the direct interaction between the secreted substrate and XcpY was confirmed by in situ photo-crosslinking. All together, our results allowed us to propose the most advanced integrative model of Xcp T2SS assembly and function.
Background
Bacterial type III secretion systems are cell-envelope spanning effector protein-delivery machines employed by Gram negative pathogens and symbionts. The membrane-embedded core unit of these secretion systems is termed needle complex. It can be divided into a base that anchors the complex to the inner and outer membranes, a hollow filament composed of inner rod and needle subunits that serves as conduit for substrate proteins, and an export apparatus facilitating substrate translocation located at the center of the base.

Objectives
While the low resolution structural analysis of the base has revealed the stoichiometry of this large ring-forming complex composed of three different proteins, the stoichiometries of the five hydrophobic export apparatus components and of the inner rod are largely unknown.

Methods

We employed a mass-spectrometry based peptide-concatenated standard strategy to evaluate the stoichiometry of the entire needle complex.

Conclusions
Using this strategy, we could validate the hitherto suggested stoichiometries of the base components and major export apparatus protein. We further propose a stoichiometry of 5:1:1:1:1 for the minor export apparatus components SpaP, SpaQ, and SpaR and for the switch-protein SpaS of the type III secretion system located on pathogenicity island 1 of Salmonella Typhimurium. The herein presented data also suggest that the inner rod is merely a rod but rather a disc because of the low stoichiometry observed.

Even though the complexity and hydrophobic character of the evaluated components allowed only the obtainment of approximate stoichiometries in this study, the gained results provide the first complete picture and a valuable framework for further investigations.
CHARACTERIZATION OF A NEWLY DISCOVERED TOXIN-ANTITOXIN FAMILY PREDICTED TO HAVE AN ADP-RIbosYLATING TOXIN

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Background

Toxin-antitoxin (TA) systems are highly abundant in bacteria and are typically composed of adjacent genes that encode both stable toxin and its unstable ‘antidote’. TA systems were shown to have a diverse set of physiological roles that range from phage defense to bacterial persistence. We recently reported the discovery of six new types of TA systems (Sberro \textit{et al.} Discovery of functional toxin/antitoxin systems in bacteria by shotgun cloning. Mol Cell. 2013 50(1):136-48).

Objectives

Here, we aimed in characterizing one of these newly discovered TA systems, Pmen TA. We propose that this toxin is involved in ADP-ribosylation, a post-transcriptional modification of crucial components within the bacterial cell. This mechanism is observed in a variety of bacterial exotoxins.

Methods

We use a combination of bioinformatics tools, molecular biology and biochemistry in order to attain initial characterization of this TA system after expressing it in \textit{E.coli}.

Conclusions

The Pmen Toxin is predicted to have an NAD-binding domain that allows ADP-ribosylation activity. The anti-toxin is unstable and physically binds the toxin. It has an N-terminal domain, known to bind ADP-ribose, and its removal eliminates the rescue effect against the toxin. The conserved residues of the toxin resemble the core active site residues in other ADP-ribosylating toxins, such as Cholera toxin. Substitution of the most conserved active site residue leads to abolishment of toxicity. Possible common evolutionary origin between toxin-antitoxin systems and lethal exotoxins suggests that TA systems may be a genetic source for the emergence of pathogen-related toxins.
THE CELLULAR LOCALIZATION AND RIBOSOME ASSOCIATION OF ACINETOBACTER BAUMANNII TYPE II TOXIN-ANTITOXIN SYSTEM COMPONENTS
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Background
Most prokaryotic genomes, including human pathogens, code for toxin-antitoxin (TA) systems, usually consisting of a toxin and its cognate antitoxin, which counteracts toxin activity. TA systems have been associated with a variety of biological functions, including general stress response, biofilm formation and pathogenicity. \textit{Acinetobacter baumannii} is multidrug-resistant pathogen causing serious hospital-acquired infections worldwide. The internally encoded TA systems can be a useful tool in understanding and preventing the spread of multi-drug resistant \textit{A. baumannii}.

Objectives
Recently, at least five functional \textit{A. baumannii} TA systems (RelBE, HicAB, HigBA, SplTA, CheTA) were discovered. Their toxins were shown to inhibit translation in \textit{Escherichia coli} cells. In order to characterize the \textit{A. baumannii} toxins in more detail, we have investigated their cellular localization and ribosome-association.

Methods
Series of inducible \textit{E. coli} expression vectors encoding \textit{A. baumannii} TA components, fused with fluorescent proteins have been constructed. The functionality of fusion proteins in \textit{E. coli} was confirmed using the kill-rescue assay. The ability of toxins to interact with ribosome was tested using ribosome fractionation, cellular localization was accessed by fluorescence microscopy.

Conclusions
Microscopy of \textit{E. coli} where the fusion toxins, alone or with their cognate antitoxins, were expressed, has shown either polar or whole cell localization pattern. The analysis of fluorescence profiles of \textit{E. coli} ribosomal fractions has demonstrated only RelE associated with ribosome, while association with the ribosome is unlikely for HicA, HigB and SplT toxins of \textit{A. baumannii}. The latter toxins might cleave RNA independently from ribosome, or inhibit translation by other mechanisms.
Background
Microcystins, produced primarily by cyanobacterium *Microcystis aeruginosa*, are the most commonly studied cyanotoxins (1). Most studies have focused only on the toxic effects of microcystins on humans and animals while their physiological or ecological roles have not been elucidated. Therefore, there is a considerable interest in understanding the importance of cyanotoxins in natural environments (2).

Objectives
The aim of the present study was to clarify the ecological significance of microcystins in algal succession.

Methods

*Desmodesmus subspicatus* was cultured in a medium together with dialysis tubing containing *M. aeruginosa* 7806. Cultures were incubated under 30 μEm⁻²s⁻¹ continuous irradiation at 23°C for 4 weeks. Microcystin-LR was quantified using mass spectrophotometer and algal density was obtained by measuring the OD₇₅₀.

Conclusions

Microcystin-LR was determined in the media after two weeks but there was no change in *D. subspicatus* growth rate. After 4 weeks incubation, *D. subspicatus* growth decreased. Using co-cultivation method gave the opportunity to investigate the algal interactions at continuous toxin production. By continuing the growth of *M. aeruginosa*, toxin concentration increased and caused a decrease in *D. subspicatus* growth. Results provided evidences to suppose the role of toxin as an allelochemical in algal interactions.

References:


Background

*Campylobacter fetus* cause human infection and are important pathogens. Recent comparative genomics of *C. fetus* subspecies revealed *Fic* loci encoding Fido superfamily proteins, which may contribute to niche adaptation and pathogenicity.

Objectives

We analysed function of 4 *Fic* loci organizes as toxin-antitoxin (TA) modules on the chromosome and ICE of *C. fetus* subsp. *Venerealis* 84-112.

Methods

The *C. fetus* proteins and mutant variants were expressed ectopically in *E. coli*, yeast and in transiently transfected HeLa cells. Prevalence of *fic* genes in 102 *C. fetus* isolates was surveyed with PCR.

Conclusions

We show that Fic proteins are cytotoxic to human cells but not *S. cerevisiae*. The *fic* loci are organized as TA modules on the chromosome and ICE of *C. fetus* subsp. *Venerealis* 84-112. Expression in *E. coli* validated the cytotoxic and neutralizing activities of the proteins, providing the first functional evidence for TA systems in *Campylobacter*. Reversal of *fic*-mediated filamentation and growth inhibition in *E. coli* also revealed antitoxin crossreactivity between loci. Key active site residues involved in adenylylation by Fic proteins are conserved in Fic1, Fic3 and Fic4, but degenerated in Fic2. We show the non-canonical Fic2 disrupts assembly of *E. coli* ribosomes. *fic* genes are prevalent in *C. fetus* subsp. *Venerealis* but not generally conserved among *Campylobacters*. Strikingly, homologous genes are found in some *Campylobacters* and unrelated pathogens adapted to the human and animal
urogenital tract. *C. fetus* fic genes thus appear to be important to adaptation and virulence in this niche.
TOXIN-ANTITOXIN SYSTEMS ON ANTIBIOTIC RESISTANCE PLASMIDS: JUST PLASMID MAINTENANCE?

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Background

Toxin-Antitoxin (TA) systems are small genetic elements composed of a toxin gene and its cognate antitoxin. They were originally discovered as plasmid-borne loci that promote plasmid maintenance by killing daughter cells that have lost their TA encoding plasmid, a phenomenon known as post-segregational killing. Nowadays it is known that a great diversity of plasmids, including the antibiotic resistance (AbR) plasmids, encodes one or even more TA systems. The widespread dissemination of antibiotic resistance genes is ascribed to the spread of bacterial plasmids encoding these genes and this successful dissemination it is attributed in part to TA systems and its participation in plasmid maintenance. However, there are some lights indicating that plasmidial TA systems could participate in functions other than plasmid maintenance.

Objectives

Our main aim was to elucidate the diversity and function of TA systems encoded on AbR plasmids. Specifically to characterize a new putative TA system encoded on an IncX4 plasmid.

Methods

By bioinformatic tools a novel putative TA system was identified in pJIE143, an IncX4 plasmid carrying blaCTX-M-15 from an Escherichia coli ST131 isolate. We analyzed the effect of the expression of this system on E. coli growth and its participation on plasmid maintenance.

Conclusions

pJIE143 encodes a novel TA system that appears to be related exclusively to IncX plasmids. The characteristics and putative functions of this novel TA system will be discussed in this work. These results reveal that AbR plasmids could contain an underappreciated diversity of TA systems, which could have functions further of plasmid maintenance.
Background
The German cockroach, Blattella germanica (L.) has been recognized as a serious health problem throughout the world. Control failures due to insecticide resistance and chemical contamination of environment have led some researchers focus on the other alternative strategy controls. Microbial insecticides such as those containing entomopathogenic fungi could be of high significance. Lecanicillium muscarium and Beauveria bassiana grow naturally in soils throughout the world and act as a parasite on various arthropod species, causing white muscardine disease. Thus, these two species could be considered as entomopathogenic fungi.

Objectives
The current study conducted to evaluate the toxicity of Beauveria bassiana and Lecanicillium muscarium against German cockroach, Blattella germanica.

Methods
Conidial formulations of L. muscarium (PTCC 5184) and B. bassiana (PTCC5197) were prepared in aqueous suspensions with Tween 20. Bioassays were performed using two methods including submersion of cockroaches in conidial suspension and baiting. Data were analyzed by Probit program and LC50 and LC90 were estimated.

Conclusions
The obtained results indicated that both fungi species were toxic against German cockroach however; Beauveria bassiana was significantly 4.8 fold more toxic than L. muscarium against German cockroach using submersion method.
REGULATION OF TOXIN-ANTITOXIN SYSTEMS WITHIN THE SOS REGULON IN E. COLI

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Background

TA systems (TAs) encode a stable toxic protein and its cognate unstable antitoxin that is either an antisense RNA (type I) or a protein (type II). Under steady-state conditions, both components are produced and cells are growing normally. Under conditions in which the antitoxin:toxin ratio is perturbed in favor of toxin, cell growth can be impaired. Therefore, maintaining an appropriate ratio is essential for viability. While negative autoregulation is observed for type II systems, few information is available for type I. Some of these TAs (symER, tisAB-istR and dinQ-agr) are part of the SOS regulon, as well as two type II TAs (dinJ-yafQ and yafNO).

Objectives

Our objective is to investigate the transcriptional regulation of these TAs at the population and single cell levels under steady-state growth and SOS conditions.

Methods

Approaches based on fluorescent reporters fused to TA promoters combined with flow cytometry and microscopy analysis are currently being developed.

Conclusions

Our data confirmed that the dinJ-yafQ and symE promoters are activated during SOS response, although at different timing and levels. Interestingly, the dinJ-yafQ promoter is active both in wild-type and dinJ-yafQ mutant strains, indicating that SOS regulation prevails on autoregulation. Furthermore, heterogeneity in promoter activity was observed at the single cell level, suggesting that phenotypic heterogeneity might play a role in the SOS response. This will be further investigated.
Background

Type II toxin-antitoxin systems (TAs) are small bicistronic modules encoding a toxic protein and its cognate antitoxin protein. TAs are widespread in bacterial and archeal genomes.

Objectives

Our hypothesis is that TAs might constitute a reservoir of toxic domains that are recruited by polymorphic toxin systems.

Methods

Polymorphic toxin systems, such as CDI and Rhs, are composed of large conserved proteins containing variable carboxy-terminal (CdiA-CT or RhsA-CT) domains showing toxic activity and small immunity proteins. Genetic organization of these systems is analogous to that of TAs: a toxic gene or domain followed by an antitoxin or immunity gene. A bioinformatics approach is currently developed to evaluate whether CdiA-CT and RhsA-CT toxic domains are found in TAs. Domains are collected and HMM profiles are built. Bacterial genomes are scrutinized for the presence of these domains under TA organization. Functionality of these systems is tested in *E. coli*. Toxin activity and targets are then investigated.

Conclusions

We identified and experimentally validated novel toxic domains that are found in TAs organization and in CDI and/or Rhs systems. Upon overexpression, toxins from TAs and CdiA-CT/Rhs-CT domains are toxic and antagonized by the expression of cognate antitoxin and/or immunity proteins. These domains were shown to inhibit translation. These data indicate that toxic domains are indeed shared by TAs and polymorphic toxin systems. Recruitment of toxic domains encoded by TAs by more
sophisticated systems might constitute a strong selective pressure on TAs and might explain why these small modules are evolutionary successful.
NOVEL FAMILY OF TOXINS WITH ACETYLTRANSFERASE DOMAIN CONSTITUTE TYPE II TOXIN-ANTITOXIN SYSTEMS

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Background
Type II toxin-antitoxin (TA) systems are small genetic loci coding for a toxic protein and its antidote. Such systems are widespread in bacterial and archaeal genomes. These systems stabilize mobile genetic elements and may act as stress response elements. Toxins of type II TA systems are most commonly mRNAses, with the exception of gyrase inhibitors and several families that have more elaborate enzymatic functions. We have found novel class of toxins that possess acetyltransferase domain and constitute typical type II TA systems.

Objectives
To functionally describe TA modules of this novel family and explore mechanism of toxicity of acetyltransferase domain toxin.

Methods
TA systems from \textit{Escherichia coli} and \textit{Vibrio cholerae} have been functionally validated by killing/rescue assay. Activity of toxin has been tested by measuring rates of replication, transcription and translation, mRNAse activity by Northern blots analysis. Mutational analysis on acetyltransferase active site has been performed. TA complex has been purified and separated by affinity chromatography. Bioinformatics analysis has been performed to describe the prevalence and diversity of this novel family.

Conclusions
Small acetyltransferase domain proteins found in pair with RHH-domain antitoxins constitute functional TA pairs. When expressed together they form a heteromultimeric complex of 2 toxins and 2 antitoxins. When expressed alone toxins possessing acetyltransferase domain inhibit translation without degrading mRNAs and therefore exhibit novel toxicity mechanism. Mutations in the acetyltransferase active site abolish toxicity, strongly indicating that toxicity relies on target(s) acetylation. Detailed analysis of translation inhibition \textit{in vitro} is currently under progress and should reveal the target(s) of the toxin.
CLAMP ALLOSTERY AND STEROSELECTIVITY GOVERN THE MOLECULAR MECHANISM OF PROTEIN TRANSLOCATION THROUGH THE ANTHRAX TOXIN PROTECTIVE ANTIGEN CHANNEL

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Background

Anthrax toxin, composed of protective antigen (PA) and lethal (LF) and edema (EF) factors, is a model system for transmembrane protein translocation. The PA component self-assembles into a proton motive force (PMF)-driven channel, which translocates LF and EF into target cells. The channel contains multiple nonspecific polypeptide-clamp sites, which catalyze unfolding and translocation. It is unknown how clamps facilitate transport while avoiding tightly bound kinetic traps.

Objectives

As the α clamp, specifically recognizes α helix, we asked how the conformational dynamics of the translocating chain might avoid kinetic traps during translocation.

Methods

~50-residue peptides (based upon the sequence of LF) were synthesized with identical sequences but variable stereochemistries to control for side-chain chemistry while manipulating helical sense and backbone flexibility. Single-channel and ensemble planar lipid bilayer electrophysiology monitored peptide binding, dissociation, and translocation.

Conclusions

Cooperative binding and allosteric modulation of translocation were only observed with an isotactic peptide synthesized only with l-enantiomers (αl). Cooperativity depended on acidic pH and the channel's α-clamp and ϕ-clamp sites. Incorporation of δ-enantiomers disrupted cooperativity, allosteric control of translocation, and slowed the rate of translocation. The translocation of αl occurs via a subsecond series of consecutive irreversible transitions and is unlike the translocation of a syndiotactic
peptide, synthesized with alternating L- and D-enantiomers (αld), which translocates 100-times slower (as slow as 776-residue LF). Therefore, to avoid kinetic traps, which may otherwise impede translocation, the channel's clamps operate allosterically through efficient, repeated, stereospecific contact along the length of the translocating peptide chain.
Background

The symbiotic interaction between the bacterium *Sinorhizobium meliloti* and the legume *Medicago sp* results in the development of new root organs, the nodules, where the differentiated bacteria (the bacteroids) reduce atmospheric N\(_2\) to ammonia. A few weeks after symbiosis establishment, nodules present a premature senescence.

Objectives

Since VapBC-type Toxin-Antitoxin (TA) systems of pathogenic bacteria is involved in the survival of the bacteria, we examined the role of *S. meliloti* VapBC modules in bacteroid viability and nodule senescence. The *vapBC* genes form an operon negatively regulated by the TA complex. Free toxin (VapC) has an RNase activity that is neutralized by the antitoxin (VapB).

Methods

We showed that, during the symbiotic interaction, a mutation in the *vapC5* toxin of the VapBC5 module, leads to a higher nitrogen-fixing activity, plant yield increase and a delayed nodule senescent phenotype. Thus, inactivation of this toxin improves symbiotic efficiency (Lipuma et al., 2014). We obtained an opposite phenotype with another mutant of toxin deletion (*vapC7*): aberrant nodules, where nitrogen fixation is nearly abolished. Nodule ultrastructure analysis and flow cytometry demonstrated that bacteroid differentiation was followed by a rapid bacterial death leading to early senescence phenotype. These results demonstrate that this Toxin Antitoxin module is essential for bacteroid viability (in preparation).

Conclusions

We demonstrated the role of two Toxin Antitoxin systems in *S. meliloti* survival during *Medicago sp.* interaction. The two mutants analysed show different phenotypes. To a better understand the role of all VapBC systems; phylogenetic analyses were done to define the evolution origin of these genes.
VAPC FROM THE LEPTOSPIRAL VAPBC TOXIN-ANTITOXIN MODULE DISPLAYS SPECIFIC RIBONUCLEASE ACTIVITY ON THE INITIATOR TRNA.

VAPC HIGH PRESSURE REFOLDING AND IN SILICO STRUCTURAL 3D STUDIES.

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Background

The most accepted hypothesis of the physiological function of the ubiquitous prokaryotic Toxin-Antitoxin (TA) operons is the reversible cessation of cellular growth under stress conditions. vapBC operons, present in Leptospira interrogans, are classified based on the presence of a ribonucleasic PIN domain in the VapC toxin. Their mechanisms of action remain mostly unknown. The expression of the leptospiral VapC in E. coli promotes a strong bacterial growth arrestment, making it difficult to obtain the recombinant protein.

Objectives

Produce soluble and active VapC in order to find its target and characterize its activity. Study and rank alternative 3D models of VapC.

Methods

vapBC locus, vapB and vapC of L. interrogans were cloned and expressed in E. coli. Proteins were purified by IMAC after refolding by pressurization (VapC). VapB and VapC interaction was tested by pull-down assay and affinity blotting. Ribonuclease activity was tested towards E. coli rRNA and tRNAfMet. Models of leptospiral VapC were generated and ranked using SWISS-MODEL.

Conclusions

VapC was refolded by high hydrostatic pressure, providing a new method to obtain the active toxin. VapB neutralizes the activity of VapC in vivo and in vitro. The 3D model of the leptospiral VapC closely matches the Shigella’s VapC X-ray structure. In agreement, leptospiral VapC shows no activity towards E. coli rRNA and was found to cleave tRNAfMet. This finding suggests that the cleavage of the initiator tRNA may
represent a common mechanism to a larger group of bacteria and potentially configures a mechanism of post-transcriptional regulation leading to the inhibition of global translation.
Background
MazF family toxins, representative endoribonucleases that cleave single-stranded RNA (ssRNA) in a sequence-specific fashion, are distributed among bacterial and archaeal lineages. Since their sequence pattern of cleavage sites is highly diverse, they are of importance not only in terms of prokaryotic physiologies, but biotechnological applications.

Objectives
To elucidate physiological roles of MazF family toxins and/or to expand the biotechnological utilization of these endoribonucleases, we established a novel easy-to-use method to effectively determine the pattern of the sequences at their cleavage sites.

Methods
The sequences cleaved by MazF family toxins were determined by a modified RNA-Seq protocol. To evaluate our method, MazF, an ACA specific endoribonuclease derived from Escherichia coli, was selected as a model target. Besides, the sequence digested by an uncharacterized MazF homologue from Proteobacteria was also identified using this method. The sequences recognized by these endoribonucleases were further analyzed by using Fluorescence Resonance Energy Transfer (FRET) based assays.

Conclusions
The method developed in this study correctly inferred the sequence pattern at the cleavage sites of the model target, E. coli MazF, as ACA. Furthermore, the possible sequence pattern recognized by MazF homologue from Proteobacteria was consistent with the sequence confirmed by FRET based method, indicative that the present method is a useful and straightforward means to explore the potential sequence recognition pattern of these endoribonuclease. This method may be applicable to various toxins that catalyze ssRNA cleavage in a sequence-specific manner, facilitating subsequent physiological investigations and practical uses of these enzymes in the genetic engineering.
IN VIVO CHARACTERIZATION OF A BACTERIAL KILLER TYPE IV SECRETION SYSTEM EFFECOR/IMMUNITY PROTEIN PAIR

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Background

Type IV secretion Systems (T4SS) are bacterial nanomachines that transfer proteins across the cell envelope and into target cells. We recently provided the first demonstration that the Xanthomonas citri (Xac) T4SS mediates the contact-dependent killing of other bacterial cells [1]. One Xanthomonas T4SS effector (X-Tfe) X-Tfe{sub XAC2609} has in vitro lysozyme-like lytic activity. This enzymatic activity is inhibited by the soluble domain of the putative Xanthomonas T4SS immunity protein (X-Tfi) X-Tfi{sup XAC2610}.


Objectives

Here, we set out to investigate the in vivo function of X-Tfi{sup XAC2610}. Since X-Tfe{sup XAC2609} is predicted to act in the bacterial periplasm and X-Tfi{sup XAC2610} has a lipobox which directs it to the bacterial outer membrane, the hypothesis is that X-Tfi{sup XAC2610} acts as an immunity factor against the possibly detrimental autolytic effects of X-Tfe{sup XAC2609}.

Methods

Xac mutants strains carrying single gene deletions (ΔX-Tfi{sup XAC2610}, ΔX-Tfi{sup XAC2609} and ΔT4SS) and double mutants (ΔX-Tfi{sup XAC2610},ΔT4SS and ΔX-Tfi{sup XAC2610}, ΔX-Tfi{sup XAC2609}) were created. Colony phenotypes and cellular viabilities were compared for the above mutant and wild type Xac strains.

Conclusions

1) X-Tfi{sup XAC2610} provides immunity against the in vivo auto-lytic activity of X-Tfe{sup XAC2609}.

2) The in vivo autolytic activity of X-Tfe{sup XAC2609} does not depend on a functional T4SS.
3) $\text{X-Tri}^{\text{XAC2610}}$ inhibitory activity is also independent of the Xac T4SS.
PLASMID-BORNE MAZEF TOXIN-ANTITOXIN LOCI ARE WIDESPREAD IN ANTIBIOTIC-RESISTANT STAPHYLOCOCCUS AUREUS, AND TOXIN ACTIVATION PROVIDES SELECTIVE KILLING

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Background
Antibiotic-resistant bacteria have become a global concern and new strategies to control pathogenic bacteria are urgently needed. In theory, activation of the toxin or inhibition of the antitoxin within a bacterial toxin-antitoxin (TA) system could provide a potent new antibiotic therapy.

Objectives
We aimed to identify the location of TA systems, to evaluate the functionality of TA systems in clinical isolates of antibiotic-resistant S. aureus, to determine the functionality of TA system and to evaluate the antitoxin as an attractive antimicrobial target for the eradication of antibiotic-resistant S. aureus.

Methods
To evaluate potential TA loci as therapeutic targets, we screened the plasmid and chromosome sequences of 1000 clinical isolates of S. aureus for the presence of TA loci.
Plasmid transformation, ATPase assay, Turbidity and PNA assay were performed.

Conclusions
Plasmid-borne mazEF TA loci were present in all tested, antibiotic-resistant S. aureus strains in Iran and Malaysia. A subset of strains expressed the mazE and mazF transcripts, and ATPase and growth assays revealed that the mazEF TA loci were functional. In addition, the plasmid-borne copies were stable in the absence of antibiotic selection. To activate toxin expression, we targeted the mazE antitoxin mRNA using peptide nucleic acid (PNA) oligomers. The anti-mazE oligomers were bactericidal against drug-resistant S. aureus containing mazEF and did not inhibit strains lacking mazEF. Therefore, mazEF TA loci are widespread in drug-resistant strains of S. aureus and are plasmid-borne, and activation of toxin activity by silencing of the antitoxin gene provides a means to selectively kill drug-resistant strains.
CHARACTERIZATION OF TOXIN-ANTITOXIN SYSTEMS IN THE HUMAN PATHOGEN STREPTOCOCCUS PYOGENES

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Background

The Gram-positive bacterium Streptococcus pyogenes, also known as group A Streptococcus (GAS), is an important human pathogen responsible for a wide spectrum of diseases such as pharyngitis, necrotizing fasciitis, and toxic shock syndrome. In addition to encoding multiple factors for adaptation and pathogenicity, GAS contains several predicted type II toxin-antitoxin (TA) loci. TA systems are widespread in bacterial pathogens and enable bacteria to adapt to rapidly changing environmental conditions, thereby contributing to the pathogenicity of the organisms.

Objectives

To understand the role in physiology and virulence of these putative TAs in S. pyogenes, we aim to characterize hypothetical toxin-antitoxins (TAs) and unravel the molecular mechanisms involved in their regulation and mode of action.

Methods

Operon architecture and expression of the predicted loci was analyzed by RT-PCR, RNA sequencing (RNA-seq), and reporter-fusions. Toxin and antitoxin activity and function was assessed by growth arrest assays, activity assays, Western blotting and microscopy.

Conclusions

A series of transcriptional fusions was constructed that enabled the confirmation of TAs annotation. Three out of four predicted TAs cause growth arrest upon overexpression. We show that two of the predicted toxins cause a reduction in CFUs but not in optical density, which can be explained by a cell division defect. Expression of the cognate antitoxins was able to relieve the growth arrests. Understanding the roles of TAs will increase our knowledge on the regulation of bacterial pathogenicity, and may reveal key targets for potential novel antibacterial strategies.
EXPOSURE TO TICKS AND SEROPREVALENCE OF BORRELIA BURGDORFERI
LIVING IN THE PROVINCE OF ERZINCAN, TURKEY

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Background
Lyme borreliosis is a zoonosis caused by Borrelia burgdorferi transmitted by ticks, especially Ixodes species.

Objectives
The aim of this study was to determine the Borrelia Burgdorferi seroprevalence caused by ticks in the province of Erzincan, which has a high tick population due to its geographical position and climate conditions.

Methods
368 people, who were live in rural area in the province of Erzincan located in the east of Turkey were included in the study. In all the acquired serum samples, the B.burgdorferi IgG antibodies were screened by using the ELISA method (SERION ELISA classic Borrelia burgdorferi IgG). The positive and doubtful results obtained were confirmed using the Western blot method (The VIRO-BLOT Anti-Borrelia IgG test).

Conclusions
B.burgdorferi IgG antibodies were determined as positive (4.1%) in 15 of all cases and suspicious in 36 cases by using the ELISA method. B.burgdorferi IgG levels were examined by using the Western Blot method in totally 51 serum samples that were determined to be positive and doubtful according to the result of the SERION ELISA classic Borrelia burgdorferi IgG test. Evaluating all the results together; B.burgdorferi IgG positivity was found at the rate of 2.17 % in 8 of 368 cases. The story of tick contact was determined in 3 of cases that were determined to have positive B.burgdorferi IgG. This study proved the presence of Borrelia Burgdorferi in the province of Erzincan. It is thought to offer the preventive health services for those in the risk group in order to minimize the tick exposure.
FEMS-1905
Vector-borne pathogens

SEROPREVALENCE OF CRIMEAN-CONGO HEMORRHAGIC FEVER LIVING IN THE PROVINCE OF ERZINCAN, TURKEY
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Background

Crimean-Congo Haemorrhagic Fever (CCHF) is a zoonotic disease caused by a tick-borne CCHF of the genus Nairovirus of the family Bunyaviridae.

Objectives

The aim of this study was to determine seroprevalence of the CCHF in animal breeders in the province of Erzincan, for which there has been no seroprevalence study despite being accepted as an endemic region in terms of the CCHF by the Ministry of Health.

Methods

372 people, who were live in the province of Erzincan located in the east of Turkey were included in the study. In all the obtained 372 serum samples, the CCHF IgG antibodies were screened by using the ELISA method. The samples that were found to be CCHF IgG positive were examined with CCHF IgM. CCHF IgM-IgG ELISA kits were used in order to determine the antibody levels. IgM positive samples were processed for detection of viral RNA through RT-PCR.

Conclusions

In 13.9% (52/372) of all samples included in the study, the CCHF IgG was determined as positive. This ratio was found respectively as follows; 16.7% (29/174) in people who had a tick contact/bite, 12.4% (18/145) in people who animal breeders and had no tick contact and 9.4% (5/53) in people who were not animal breeders in the city center and had no tick contact. Seven samples were found positive for IgM whereas viral RNA was detected in 1 samples. In the province of Erzincan where many CCHF-related death cases are reported every year, the CCHF IgG antibody level was found to be high as expected.
Background
Cutaneous Leishmaniasis (CL) is a common endemic parasitological disease in Iran. This disease is always serious health problems in Esfahan province specially Badrood city and its prevalence has been doubled over the last decade.

Objectives
This study was designed to determine the epidemiology and clinical characteristics of CL in Badrood in 2013.

Methods
This descriptive cross sectional study was on all detected patients with cutaneous leishmaniasis in the city Badrood by active detection within one year in 2014. Patients were visited by physician and a questionnaire including information about demographic and characteristics of wounds were filled through interview and examination. Diagnosis was confirmed by revealing of leishman body in smear of wounds.

Conclusions
we found 63 definitive case of CL. 55.6% of patients were male. Age group 15-30 years had the highest rate (23.8%) among the patients. 66.7% of patients had history of stay in Agha Ali Abbas region. All wounds were rural form. Most patients with cutaneous leishmaniasis in Badrood had a wound with diameter less than 3 cm, secretion, painless, itch and extremities was the most common site for infection.

Based on the findings of this study, cutaneous leishmaniasis in Badrood city is rural form and most patient are men and older than 15 year. there was complete or partial healing after treatment in the most patients.
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Based on the findings of this study, cutaneous leishmaniasis in Badrood city is rural form and most patient are men and older than 15 year. there was complete or partial healing after treatment in the most patients.
PREVALENCE OF INTESTINAL PARASITES AMONG PRIMARY SCHOOL PUPILS IN ABAKALIKI METROPOLIS
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Background
Intestinal parasites are still major health challenges in tropical and sub-tropical areas. They are commonly found among people with low socio-economic status and in unhygienic environments. Children are usually more infected than adults due to easier fecal-oral transmission¹. Children with parasitic infections are usually undernourished, weak and may have complications like diarrhoea, nutritional disorders and potentially lethal systemic disease.

Objectives
To determine the prevalence of intestinal parasites among primary school children in Abakaliki metropolis.

Methods
A total of 180 stool samples were aseptically collected and examined macroscopically by wet preparation (saline and iodine) and then microscopically by formal ether concentration techniques. The study was approved by Anambra State University Teaching Hospital Review Board.

Conclusions
Out of the 180 stool samples examined, intestinal parasitic infection prevalence of 25.0% was recorded. Hookworm showed the highest of prevalence, followed by Ascaris lumbricoides and Gardia lamblia, while Entaemoba histolytica had the least prevalence. Prevalence in male was insignificantly higher than that of female (p>0.05). The highest prevalence was recorded amongst age group 5-7 years. There was no significant difference in the rate of parasitic infection in relation to age (p>0.05). Higher number of parasites was seen in the concentration technique as compared to other laboratory techniques (normal saline and Iodine method). Higher prevalence was seen among the children of farmers compared to other children. A well-structured control programme is needed to reduce mortality and morbidity from intestinal parasites and sustained commitment by the government and policy makers.
are needed to improve the quality of life of these children.
EVALUATION OF HAIR SAMPLES AS TARGET TO DIRECT DETECTION OF LEISHMANIA SPP. IN LEPORIDAE

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Background

Leishmaniasis is a parasitic zoonotic disease caused by protozoa of the genus Leishmania, responsible of high mortality in developing countries and considered one of the three major neglected parasitic diseases in developed countries. Wild reservoirs have a key role in certain epidemiological situations, i.e. Leporidae were significantly involved in the largest recorded European outbreak of leishmaniasis, in Madrid (Spain), still active. The possibility of monitoring Leishmania spp. using samples easy to obtain, with noninvasive techniques and stable for transport/storage is an essential key for surveillance programs in wild reservoirs. Previous studies have shown that hair samples are ideal for this purpose in dogs, mice and some wild species, but their suitability in wild Leporidae has not been studied before.

Objectives

Assessment of hair as a target for direct detection of Leishmania infantum in wild Leporidae.

Methods

Seventeen wild rabbits and nine hares from Madrid were included. Sera from all the animals were tested by immunofluorescence antibody test (IFAT). Moreover, hair, skin and spleen samples were subjected to DNA extraction and analyzed by real-time PCR for Leishmania infantum.

Conclusions
All animals sampled (n=26) were positive to at least one of the three classic samples (serum, skin, spleen): 23 individuals (88.5%, 95%CI: 77.3%-100%) were positive in IFAT and 25 (96.2%, 95%CI: 88.2%-100%) were positive in skin/spleen RT-PCR.

A total of 22 hair samples (84.6%, 95% CI: 69.7%-99.8%) were positive. Despite the limited sample size, our results demonstrate that hair could be considered an adequate sample for direct diagnosis of *Leishmania infantum* infection in wild *Leporidae*. 
A NEW SPECIES OF CHOLEOEIMERIA (APICOMPLEXA: EIMERIIDAE) PARASITIC IN THE ROUGH-TAILED GECKO CYRTOPODION SCABRUM (SAURIA: GEKKONIDAE) IN EGYPT

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Background

Cyrtopodion Fitzinger is a large genus of the Gekkonidae Gray (gecko) family with currently 28 described species. In contrast to the large diversity of this genus, only Eimeria lineri McAllister et al. 1988 has so far been described. In Egypt, the genus Cyrtopodion is represented only by Cyrtopodion scabrum Heyden 1827. Genus Choleoeimeria Paperna and Landsberg 1989 was proposed to incorporate eimeriid-like coccidia of elongate-ellipsoidal oocysts (L/W ratio 1.6–2.2), with sporocysts that lack a Stieda body and that develop in the epithelium of the gall bladder and bile duct of their hosts.

Objectives

The present work aims to establish the taxonomic status of an unknown Choleoeimeria from the gallbladder of the rough-tailed gecko Cyrtopodion scabrum using the characteristics of exogenous and endogenous stages of the parasite.

Methods

A total of 15 rough-tailed geckoes, Cyrtopodion scabrum were collected and examined for coccidian infection. The presence of oocysts was determined by direct microscopic examination of bile contents. To study the endogenous stages, infected gallbladders were fixed in 10% neutral buffered formalin and embedded in paraffin. Sections were cut and stained with haematoxylin and eosin (H&E).

Conclusions

Based on the combination of the oocyst morphology, site and characters of the endogenous development; we assigned the present species to the genus Choleoeimeria and to which the name Choleoeimeria scabrum n. sp. was suggested.
BACTERIAL CONTAMINATION OF TOYS AND PLAY AREAS IN RESTAURANTS AND SHOPPING MALLS IN RIYADH, SAUDI ARABIA

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Background

This study was conducted to determine the load of pathogenic bacterial contamination and explore the prevalence of pathogenic bacterial species in play areas of fast food restaurants and shopping malls in Riyadh, Saudi Arabia. This novel study aims to pilot other studies in the area to explore bacterial contamination for purposes preventing pediatric disease

Objectives

The study seeks to identify and determine the load and antibiotic sensitivity of bacterial organisms in the play areas present in fast food restaurants and shopping malls in Riyadh, Saudi Arabia.

Methods

The sample included three fast food restaurants’ play pits and two shopping mall play areas in Riyadh, Saudi Arabia during the month of March in 2012. Nineteen samples were isolated using sterile swabs from surfaces of balls in ball pits and different arcade games. Samples were inoculated onto blood agar and MacConkey agar and were incubated 24 hours at 37º C. Bacterial species were identified and tested for their antibiotic sensitivity. Bacteria isolated from a total of 19 samples included Klebsiella pneumoniae, Staphylococcus spp., Diphtheroids, Pantoea agglomerans, Enterococci spp., Streptococci viridans and two fungal growths that included Aspergillus spp. and Penicillium spp.

Conclusions

The results of this study confirm that toys in play areas and arcades could serve as media for the transmission of pathogenic microorganisms and fungal elements that pose a potential health risk for children.
EFFECTS OF ORAL ADMINISTRATION OF THE BUTYRIVIBRIO GROUP BACTERIUM ON THE INFLAMMATORY DRY SKIN CONDITION IN CHRONIC IRRITANT CONTACT DERMATITIS

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Background
The genus Butyrivibrio is one of the predominant residents in the rumen, and also inhibits the intestine of monogastric animals. We previously reported that oral administration of B. fibrisolvens to mice alleviated experimental enterocolitis and suppressed the formation of colorectal aberrant crypt foci. Although the preventive mechanisms of this bacterium have been still unknown, it might have the potential to improve immune responses. Thus, some strains of this bacterial group might be used for improving the dysfunction in inflammatory skin diseases such as irritant contact dermatitis (ICD).

Objectives
The newly strain of Butyrivibrio group to improve the skin barrier function was isolated and its effect on inflammatory dry skin was examined.

Methods
Several strains of the Butyrivibrio group bacteria were isolated from the goat rumen and confirmed by reading the nearly full-length 16S rRNA gene sequences. Some strains were examined as the candidates for probiotic bacteria that recover the skin barrier function. Chronic ICD was induced to mice with oxazolone at the ears by repeated application. The isolated strains were administrated as the freeze-dried cells. The beneficial effects on cutaneous inflammation were evaluated by the dermatitis severity score, lymph node weight, histological analysis, and mRNA expression of IL-1β and IL-6 in the lesional ear.

Conclusions
When the isolated strain was administrated, the development of skin roughness or exfoliation was alleviated and mRNA expression of the pro-inflammatory cytokines was suppressed. In conclusion, this newly isolated Butyrivibrio strain might be used as a probiotics for the restoring the skin barrier function.
EMULSION OF THE AMAZON FOR DENTAL CAVITY CLEANING: PHYSICO-CHEMICAL QUALITY CONTROL AND EVALUATION OF ANTIBACTERIAL ACTIVITY

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Background
The dental field has been developed with the purpose of seeking new bioactive principles for the formulation of drugs with different applicability.

Objectives
Analyze the stability of an emulsion of copaiba oil (Copaifera multijuga – CM) for dental cavity cleaning. In this emulsion was performed physico-chemical tests and antibacterial test in different environments and periods.
Methods
The results of the pH and density determination were analyzed by ANOVA and Bonferroni's test (p < 0.05). For other tests were given descriptive analysis. In the centrifugation, there was no phase separation at time 0, in 3 months (freezer, refrigerator and air conditioning) and in 6 months (in the freezer and refrigerator); in the pH, the incubator, the room temperature protected from light, the room temperature and the air conditioning had statistically significant difference from the control group (time 0); at density, the incubator had the lowest density values; in microbiological evaluation, there was no bacterial growth; for the evaluation of organoleptic characters, there was only change in color of the emulsion in 3 months (incubator) and 6 months (incubator, room temperature protected from light and room temperature). The CM showed bacteriostatic and bactericidal activity, respectively, in concentrations of 13.33μL/ml and 15 μL/ml for S. mutans; 20 μL/ml and 23.33 μL/mL for S. salivarius and 13.33 μL/ml, 15 μL/ml for L. paracasei; 8.33μL/ml and 8.33μL/ml for S. oralis.

Conclusions
The authors concluded that the CM demonstrated bactericidal activity; presented conditions of stability and quality without contaminants, the best storage environment is refrigerator.
A CASE OF CHRONIC OTITIS MEDIA DUE TO ACHROMOBACTER XYLOSOXIDANS IN AN IMMUNOCOMPETENT PATIENT

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Background

Achromobacter xylosoxidans is an aerobic, oxidase positive, non-fermentative, gram-negative bacillus and often misidentified as Pseudomonas aeruginosa by its oxidase positive feature. It may cause serious nosocomial and community-acquired infections especially in patients with immune deficiency.

Objectives

We present a case of chronic otitis media due to A. xylosoxidans which is rarely isolated from clinical materials in routine microbiology laboratory practice. In this report we emphasize clinical impact of A. xylosoxidans and its antimicrobial resistance profile.

Methods
A 20-year-old immunocompetent man was admitted to otorhinolaryngology service of our hospital with the complaints of left ear pain, difficulty in hearing, left ear discharge beginning from nearly three months his admission. His history revealed unsuccessful treatment attempts with various oral antimicrobials with the diagnosis of otitis media. Tympanic membrane perforation and mucopurulent yellow discharge was observed. He had no systemic symptoms. Microscopic examination of the Gram stained smear revealed PMNL and gram negative bacilli. *A. xylosoxidans* was isolated from discharge. Identification was performed by VITEK 2 automated system. Antimicrobial susceptibility was detected by disc diffusion method according to CLSI criteria for *Pseudomonas* spp. It was resistant to penicillins, most of the beta lactamases, fluoroquinolones, aminoglycosides and susceptible to piperacillin-tazobactam and imipenem.

**Conclusions**

*A. xylosoxidans* strains are often multidrug resistant. Empirical antibiotic therapy with piperacillin-tazobactam or a carbapenem is a reasonable choice until the results of susceptibility tests are available. It should be kept in mind that even an opportunistic and generally nosocomial pathogen like *A. xylosoxidans* can cause community-acquired infection like otitis media in a immunocompetent host.
EFFECT OF TRANS-CHALCONE ON TRANSCRIPTION OF GENES RELATED TO CELL WALL AND FATTY ACID SYNTHESIS IN THE DERMATOPHYTE TRICHOPHYTON RUBRUM

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Background
Trichophyton rubrum is the main causative agent of dermatophytosis worldwide. The infections caused by this dermatophyte are not lethal but are uncomfortable and difficult to treat. Moreover, invasive infections are rising due to modern medical interventions and immunosuppressive diseases. The development of fungal resistance limits the arsenal of antifungal drugs, which has a narrow spectrum of activity and often presents toxicity to the host. Therefore, novel drugs with more specific and effective mechanisms of action against dermatophytes are urgently needed. Chalcones drew attention by numerous pharmacological applications, including antifungal activity.

Objectives
The aim of this study was to evaluate the modulation of genes involved in cell wall and fatty acid synthesis of T. rubrum after the exposure to trans-chalcone.

Methods
The modulation of fatty acid synthesis genes (FAS 1p and FAS 2p subunits, acetyl-CoA carboxylase 2p subunit) and genes of cell wall synthesis (beta-1,3-glucanosyltransferase and glycosylphosphatidylinositol anchored protein) was checked by qPCR after the exposure to MIC (7.8 µg/mL) and ¼ MIC (1.95 µg/mL) of trans-chalcone, respectively, for 16h for fatty acid genes and 7h for cell wall genes.

Conclusions
Trans-chalcone promoted a down-regulation of genes of fatty acid synthesis whereas cell wall genes were up-regulated. These results suggest that trans-chalcone may act on different fungal specific targets, which could be considered an advantage for an antifungal compound.
REP-PCR GENOMIC FINGERPRINTING AND NEW INSIGHTS INTO PHYSIOLOGICAL TRAITS OF ACIDIPLASMA STRAINS

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Background

Acidiplasma is the genus of acidophilic, ferrous iron-oxidizing, heterotrophic, cell wall-lacking, moderately thermophilic microorganisms belonging to the family Ferroplasmaceae. All representatives of the genus have high similarity of 16S rRNA gene (99-100%). The genus includes two species (A. aeolicum and A. cupricumulans) separated by DNA–DNA hybridization (DDH) values. However complexity of DDH suggests to necessity of application of alternative tools for distinguishing representatives within the genus. Also information about physiological traits of the genus may be incomplete. In our previous study the new physiological trait (ability to oxidize elemental sulfur) was shown by the example of strain Acidiplasma MBA-1.

Objectives

The aims of this study were to apply rep-PCR fingerprinting technique for distinguishing Acidiplasma representatives and to investigate sulfur oxidation by the strains A. aeolicum V¹, A. cupricumulans BH2¹, Acidiplasma MBA-1, Ferroplasma acidiphilum Y¹.

![Graph showing pH and sulfite oxidation over time](image)

**Figure 1.** Oxidation of sulfur by Acidiplasma strains and Ferroplasma acidiphilum Y¹
Methods
The genomic fingerprints were obtained using BOXA1, BOXS1, ERIC and REP primers. Oxidation of sulfur was estimated by pH decreasing and increasing of sulfates-ions concentrations.

Conclusions
Sulfur was oxidized by all *Acidiplasma* strains, but not by *F. acidiphilum YT* (Fig. 1). It suggests sulfur oxidation can be widely distributed among *Acidiplasma* representatives, but not among *Ferroplasma* representatives. Each of *Acidiplasma* strains possess its unique genomic fingerprint (Fig. 2), it suggests genomic fingerprinting could be rapid, highly discriminatory screening techniques for *Acidiplasma* representatives. The work was supported by the Russian Foundation for Basic Research, project no. 14-04-31210 mol_a and Core Research Facility ‘Bioengineering’ equipment.
FEMS-2995
Free subjects - 2

TREATMENT AGAINST THE MICROBIAL ACTIVITY WITH CINNAMON ESSENTIAL OIL IN ARTISTIC WORKS AND PACKAGING

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Background
The artwork and the packaging system used in all transfers and storage is a good substrate for the growth of microorganisms that can produce irreversible changes, as destruction or alteration of substrates on which microorganisms can multiply and that as a result of secondary metabolism can accumulate among them the indelible pigment.

Objectives
The aim of this study is to provide a preventive solution to the development of microorganisms without risk to the integrity of the artwork.

Methods
We proceeded through sterile swabs to collection of samples from artwork and packaging deposited in the MACBA. Samples were plated on TSA to determine the presence of bacteria and AS with antibiotic to determine fungi. All artwork and packaging were submit an aspiration synthetic pallet and vacuum equipped with a HEPA filter.
Then, we applied by spraying a mixture of alcohol: water: cinnamon and elsewhere, to compare the results for the traditionally used mixture formed alcohol: water. After a month of the application of the products we collected samples from all the treated areas in order to evaluate the efficacy of treatments. In order to assess the effect of treatment in cases of contamination, proceeded experimentally inoculated packaging materials and pieces of artwork who underwent both treatments indicated and to expose the action of an environment contaminated items artistic and packaging that did not have initially any pollution.

Conclusions
The results indicates that the mixture comprising cinnamon oil has a strong antimicrobial effect and avoid contamination of sensitive material in contact with elements altered by microorganisms.
THE EFFICACY OF PRTA TO PREVENT STREPTOCOCCUS PNEUMONIAE INFECTION

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Background

Streptococcus pneumoniae is a respiratory pathogen contributing to severe lung infection and/or bacteremic complications. Worldwide efforts were made to develop a conserved protein vaccine against S. pneumoniae and to provide a maximal and memory protection.

Objectives

The pneumococcal surface protein, PrtA, was identified by convalescent patient serum screening on a pneumococcal genomic expression library. The prtA gene is prevalent and conserved among S. pneumoniae strains; however, there was no report describing its protective efficacy.

Methods

Since IL-17A elicitation can improve the clearance of pneumococcal colonization, we combined a recombinant PrtA fragment (aa 144-1041) with curdlan, a strong inducer of IL-17A, to vaccinate three-wk old CD1 mice intranasally once a week for three weeks. It showed an elevated PrtA-specific Ab titer in mucosa and increased IL-17A production by splenocytes compared with sham group. Finally, the PrtA vaccination can successfully reduce S. pneumoniae D39 colonization in lung within 24 hr but failed to reduce the mortality after 12 days of infection. All of the experiments were approved by the Institutional Animal Care and Use Committee of the National Defense Medical Center Laboratory Animal Center. The animal experiments were performed in strict accordance with the Taiwan regulations of the Animal Protection Act and the course on Animal Care and Use in Research and Education from the American Association for Laboratory Animal Science.

Conclusions

Combined with PrtA fragment and curdlan adjuvant can enhance PrtA-specific humoral and cellular immunity; however, the vaccination was not sufficient to protect against S. pneumoniae infection.
ANTI-TOXOPLASMA ACTIVITIES OF METHANOLIC EXTRACT FROM ZEA MAYS AND ERYNGIUM CAUCASICUM

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Background

Toxoplasma gondii is an obligatory intracellular parasite that infects a wide range of warm-blooded animals and humans.

Objectives

Considering the severity of toxoplasmosis, side effects of current treatments, and the contribution of the ethnopharmacological knowledge for the treatment of parasitic infections, the aim of the present study was to investigate the efficacy of methanolic extracts of Zea mays and Eryngium caucasicum against tachyzoite of T. gondii.

Methods

Four concentrations (5, 10, 25 and 50 mg ml\(^{-1}\)) of extracts were incubated with infected macrophages for 30, 60, 120 and 180 min and then the viability of the tachyzoites were evaluated by trypan blue staining.

Conclusions

Concentrations of 10 and 25 mg ml\(^{-1}\) of Z. mays after 180 minutes, and also concentration 50 mg ml\(^{-1}\) after 120 minutes killed 100% of the tachyzoites. Also high anti-Toxoplasma activity was seen using E. caucasicum extract. The anti-Toxoplasma effects of the methanolic extracts from Z. mays and E. caucasicum did not show any significant difference in comparison with pyrimethamine (positive control). Findings of this research indicate that these extracts have acceptable anti-Toxoplasma efficacies \textit{in vitro} and it can be used as candidate extracts for further research on treatment of toxoplasmosis \textit{in vivo}.
Background

Toxoplasma gondii is a protozoan with worldwide distribution and in spite of increasing information about its biology, treatment of toxoplasmosis is restricted to a few drugs and unfortunately using of each of drugs is associated with significant side effects in patients.

Objectives

This study was designed to evaluate the efficacy of cromolyn sodium and ketotifen as alternative drugs for the treatment of toxoplasmosis.

Methods

In vitro; in case group, concentrations of 1, 5, 10 and 15 lg/ml of ketotifen and cromolyn sodium were added to RPMI medium containing peritoneal macrophages. After 1 h incubation and adding tachyzoites to medium, efficacy rate of these drugs in entrance inhibition of Toxoplasma tachyzoites into macrophages were evaluated after 30 and 60 min. In vivo; case groups received ketotifen and cromolyn sodium with different concentrations at various times. Control groups received none of drugs.

Conclusions

We found that in vitro; after 60 min the best efficacy of these drugs in inhibition of cell entrance of Toxoplasma was observed at 15 lg/ml (78.9 ± 1.70 and 91.97 ± 0.37 %, respectively) (P<0.05). In vivo; after 60 min ketotifen at 2 mg/kg in 3 h before tachyzoite injection (69.83 ± 2.25 %), and cromolyn sodium, at 10 mg/kg in 6 h before tachyzoite injection (80.47 ± 2/49 %) had the best effect on inhibition of Toxoplasma entry into the cells (P<0.05). Our findings show that ketotifen and cromolyn sodium are suitable drugs for entrance inhibition of tachyzoites into nucleated cells in vitro and in vivo.
THE NAGOYA PROTOCOL: REVOLUTION OR EVOLUTION? ACCESS AND STUDY OF MICROBIAL DIVERSITY IS SUBJECT TO NEW REGULATION

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Background

The Nagoya Protocol (NP) under the Convention on Biological Diversity (CBD) has entered into force on 12 October 2014. The NP covers the use of biological diversity in Research and Development. If not correctly translated by lawmakers or inefficiently apprehended by microbiologists it could heavily impact the activities of Culture Collections (CC) and their partners. The European lawmakers have already translated the NP into European Regulation.

[i] Nagoya Protocol on Access to Genetic Resources and the Fair and Equitable Sharing of Benefits Arising from their Utilization to the Convention on Biological Diversity. (http://www.cbd.int/abs/)


Objectives

The challenge is to organize efficiently the access and the conveyance of transfer and utilization of the microorganisms at affordable cost.

Methods

CCs have anticipated these legal developments and have presented solutions during the 1st NP’s Meeting of the Parties.
Conclusions

CCs propose TRUST\textsuperscript{[i]}, a system to help microbiologists dealing with this new set of rules on access and utilization of biological diversity, including micro-organisms.

\textsuperscript{[i]} Implementing the Nagoya Protocol in microbiology - Gaining trust, building TRUST. Note of the World Federation for Culture Collections (http://www.cbd.int/doc/meetings/abs/np-mop-01/information/np-mop-01-inf-08-en.pdf)
Background
Saccharomyces cerevisiae trehalose synthesis complex is composed by Tps1 and Tps2 and two regulatory subunits, Tsl1 and Tps3. To protect membranes trehalose must be present on both sides of the bilayer.

Objectives
Our goal was to elucidate the protective mechanism of trehalose.

Methods
The absence of Tsl1 abolished the increase in Tps1 activity and accumulation of trehalose in response to a 40°C treatment, whereas deficiency of Tps3 only reduced Tps1 activity and trehalose. In extracts of heat stressed cells, Tps1 was inhibited by trehalose-6-phosphate and by cAMP. In contrast, cAMP-dependent phosphorylation did not inhibit Tps1 in tps3 cells, which accumulated a higher proportion of trehalose-6-phosphate after heating. Agt1 deficiency led to a reduced tolerance to heat shock, without interfering in trehalose accumulation, probably because the outside of the lipid bilayer was unprotected. ath1 cells still showed a high trehalose levels when they were shifted back from 40 to 28°C, as well as, increased tolerance to a subsequent heat stress.

Conclusions
We conclude that Agt1 plays a crucial role in transporting endogenous trehalose to the outer side of the plasma membrane when yeasts face adverse conditions. With the end of the stress, the intracellular trehalose pool would be hydrolyzed by Nth1 whereas the external molecules would be hydrolysed by Ath1. To stop readily trehalose synthesis during stress recovery, Tps3 would be phosphorylated by cAMP-dependent protein kinase, decreasing Tps2 activity and, consequently, increasing trehalose-6-phosphate levels which, in turn, would inhibit Tps1. We believe that these concepts are of great importance for medical and biotechnological applications.
INVESTIGATION OF ROsmarinus OFFICINALIS L EXTRACT ON THE BACTERIA CAUSES GENITAL INFECTION

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Background
Some microbial causes in creating vaginitis are streptococcus group B and Gardnerella vaginalis which result in irrecoverable harms and increasing fatality, and providing infection in pregnancy period and transfer to the baby.

Objectives
the antibacterial effects some plants like Rosemary (Rosmarinus officinalis) on vaginitis we can identify these kinds of plants and then give them to the people in the form of plant drugs. This research is an investigation of Rosemary and its antibacterial effect on the bacteria and also comparing this effect with common antibacterial.

Methods
This study of 96 women who suffered from this disease, all samples confirms with phenotyping method. The Disk diffusion method was used for comparing the effect of antibacterial total extract with common antibiotics.

Conclusions
The MIC result for each strain showed respectively; streptococcus 1/64, Listeria 1/128, Candida Albicans 1/2, staph aureus 1/128, Gardenerella vaginalis 1/64. In the disk diffusion method, the results were in this form which streptococcus group B, staph aureus and Gradenerella vaginalis were more sensitive than antibiotic disk in comparing with extract, While Listeria monocytogenes was more sensitive than antibiotic disk to the extract.
Background

Lignin is one of the major components of plant cell wall and has significant impact on animal digestion of plants. It is known that Phanerochaete secretes a series of extracellular oxidases, which are encoded in multiple genes, to degrade lignin. However, Phanerochaete needs more stringent conditions to ferment. For animal husbandry application, it is challenge in genetic engineering to transfer all required oxidase genes from Phanerochaete into other strains that can ferment easily. Therefore, we chemically fuse protoplasts between Pleurotus ostreatus, a Phanerochaete strain, and Saccharomyces cerevisiae. To gain strains with the excellent characteristics of both Pleurotus ostreatus and Saccharomyces cerevisiae.

Objectives

We chemically fuse protoplasts between Pleurotus ostreatus, a Phanerochaete strain, and Saccharomyces cerevisiae to gain strains with the excellent characteristics of both Pleurotus ostreatus and Saccharomyces cerevisiae.

Methods

At first, the enzymatic composition, duration and temperature of emzymolysis, the types of osmotic stabilizer and the age of strains were screened to gain optimal protoplasts preparative condition through $L_{16} (4^5)$ orthogonal test. Next, Pleurotus ostreatus and Saccharomyces cerevisiae protoplasts were inactivated by heat and UV, respectively. Duration and temperature of inactivation were screened for optimal conditions. Then, the optimal chemical fusion conditions, including fusion temperature and duration, pH value, Ca$^{2+}$ concentration and polyethylene glycol-4000 concentration, were screened through $L_{16} (4^5)$ orthogonal test. At last, antagonism, morphological comparison of hypha and Random Amplified Polymorphic DNA were used to characterize the potential fusants, which was analyzed using NTSYS-PC software.
Conclusions

183 regenerative strains were obtained and, among them, 14 strains were proved to be acquired from fusing *Pleurotus ostreatus* and *Saccharomyces cerevisiae* protoplast.
ARE DOG ENZYMES INVOLVED IN THE FORMATION OF THE YEAST TPS2 DELETION MUTANT'S RESIDUAL TREHALOSE?

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Background
Trehalose is a disaccharide of which the concentration correlates very well with cellular stress resistance in yeast. It is synthesized in two enzymatic steps, were the intermediate trehalose-6-phosphate (Tre6P) is converted to trehalose by the enzyme Tps2. However, deletion of this gene does not completely abolish trehalose production. A possible explanation for this observation is that unspecific sugar phosphatases take over the function of the Tre6P phosphatase in a tps2Δ mutant. Screening the Saccharomyces cerevisiae proteome for enzymes containing the phosphohydrolase motifs present in Tps2 identified Dog1 and Dog2 as possible candidates. These enzymes are known to dephosphorylate 2-deoxyglucose-6-phosphate, but the natural substrate remains unknown.

Objectives
Are Dog1 and Dog2 Tre6P-phosphatase that are active in the absence of Tps2?

Methods
We generated S. cerevisiae DOG1 and DOG2 deletion and over-expression mutants in the wild type and a tps2Δ background and determined heat stress tolerance and trehalose levels in these strains.

Conclusions
The results obtained with the deletion strains seem to support our hypothesis. The dog1Δ dog2Δ tps2Δ mutant is more sensitive to heat stress than the tps2Δ mutant, which would indicate lower trehalose levels in the absence of the DOG genes. However, the results obtained with the DOG1 and DOG2 over-expression strains do not support the hypothesis, as trehalose levels and Tre6P-phosphatase activities are not increased in these strains. These results seem to indicate that the Dog1 and Dog2 may be involved in the expression of trehalose metabolism genes are may affect the enzymatic activity of these genes.
Background

One of the challenges of this century is to transform our economy into an eco-friendly and self-sustaining system. An innovative approach is the use of mycelium based materials.

Objectives

The mycelium of the mushroom forming fungus *Schizophyllum commune* has great potential for developing such materials.

Methods

Mycelium of *S. commune* was grown as a floating layer on liquid medium in the light or in the dark under ambient or a high concentration of CO$_2$. Mycelia were dried and subjected to increasing tensile force. Their tensile modulus (E), elongation (ε) and stress of the sample at breaking point (σ) were measured. These values were rather low for the wild-type strain 4-39 that had been grown in the light at low CO$_2$ (E=0.03±0.02 GPa, ε=0.75±0.28 %, σ: 0.44±0.31 MPa), resembling a crumble material that breaks easily. The properties of the wild-type strain 4.8b that had been grown under the same conditions resembled a flexible material (E: 0.22±0.08 GPa, ε=1.8±0.56 %, σ=3.5±0.9 MPa). Strain 4-39 and 4.8b grown in the light at high CO$_2$ had plastic-like and even stiffer properties, respectively (4-39: E=0.26±0.12 GPa, ε=1.15±0.49 %, σ=point 2.59±1.30 MPa; 4.8b: E=0.50±0.10 GPa, ε=1.13±0.21 %, σ=5.58±1.65 MPa). Similar results were obtained with 4.8b and 4-39 when grown in the dark, regardless of CO$_2$ concentration.

Conclusions

Taken together, physical properties of the mycelium is strain dependent and are affected by environmental growth conditions.
CHARACTERIZATION OF AIF5A (AEF5): AN EVOLUTIONARY CONSERVED TRANSLATION FACTOR

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Background

Protein synthesis is a complex process of fundamental importance in all cells during which a nucleotide sequence is translated into an amino acid sequence. The translation apparatus includes mRNA, ribosomes, aminoacyl-tRNAs and a number of translation factors, which assist all stages of the process. Some of the translation factors are conserved among Bacteria, Archaea and Eukarya. One of the universally conserved factors is eIF5A, highly conserved between Eukarya and Archaea, with a homologue (EF-P) in bacteria. It was initially identified as a translation initiation factor, but recent studies have established a role for both eIF5A and EF-P in translation elongation specifically in promoting the translation of polyproline containing proteins. The two proteins are both characterized by particular post-translational modifications, which are essential for their function: hypusination for eIF5A and lysinylation for EF-P.

Objectives

Eukaryal and bacterial proteins have been extensively studied, while information on the archaeal homologues are extremely limited so in order to fill this gap we have started the characterization of the archaeal protein in Sulfolobaceae.

Methods

We have produced the recombinant protein from S. solfataricus using two different expression systems: E. coli and S. acidocaldarius and the purified protein was then used in several in vitro functional tests.

Conclusions

Results clarifying its function in the translation process will be presented and a molecular modelling will shed light on its interaction with the ribosome. In addition, we failed to obtain a knock-out strain in S. acidocaldarius proving that, as for the eukaryal and bacterial proteins, the gene product aIF5A is essential.
AN INVESTIGATION OF BURKHOLDERIA CENOCEPACIA LIPOPOLYSACCHARIDE (LPS) MODIFICATION DURING CYSTIC FIBROSIS (CF) CHRONIC INFECTION AND ITS IMPACT ON HOST RESPONSE MODULATION

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Background

Bacterial species within the CF microbiome demonstrate adaptive strategies including altered expression of LPS. The structure of LPS governs its immuno-stimulatory potential. It is thus important to elucidate how LPS is modified during chronic infection and to determine its effects on host response modulation.

Objectives

We are investigating LPS extracted from four \textit{B. cenocepacia} sequential isolates cultured from a CF patient during a 3.5 year infection to determine 1) if \textit{B. cenocepacia} LPS is structurally modified during chronic infection and 2) the effect of any modification on the host response to LPS.

Methods

The LPS was extracted using hot phenol extraction and analysed via SDS-PAGE. The LPS was incubated with CFBE41o-\textsuperscript{-}, 16HBE14o-\textsuperscript{-} and HL60 cells. The pro-inflammatory cytokines were subsequently analysed by ELISA. The LPS mutant was generated by deletion of BCAL2405, an O-antigen biosynthesis gene using a tri-parental mating method.

Conclusions

The O-antigen domain was present in the LPS from the first isolate but absent from the three later isolates. The LPS from the later isolates induced a significantly greater pro-inflammatory response from CFBE41o-\textsuperscript{-} cells (\(P < 0.001\)), 16HBE14o-\textsuperscript{-} cells (\(P < 0.05\)) and HL60 cells (\(P < 0.001\)) when compared to LPS from the initial isolate. These data suggest that the loss of O-antigen in the later isolates may be responsible for the increased pro-inflammatory response. To verify this, we have successfully deleted the BCAL2405 gene from \textit{B. cenocepacia} K56-2 which yielded LPS without the O-antigen and are currently investigating the pro-inflammatory potential of the mutant strain.
Background

Millions of species of microorganisms on the earth, a large amount of them still undiscovered, are the source of biological diversity. Microorganisms are not only indispensable for life; they are basic for many human activities. Microorganisms are an essential part of the day-to-day work not only for scientists, but also for people working in the area of e.g. biotechnology, agriculture, industry, or medicine. Therefore, their economic value is very high. A large percentage of this microbial material is preserved in culture collections (CCs), in which their stability and reproducibility can be protected and through which they are (in most cases) available for other researchers.

Objectives

The implementation of the strategy on improved communication between user and provider has to be organised in a coordinated way. By arranging a major part of the CCs pro-active communication through the MIRRI portal the users will get a facilitated access to all requested information.

Methods

The main objectives of CCs and microbial resource centres (MRCs) are, besides conserving biodiversity, sharing microbial resources and associated data.

Conclusions

MIRRI has prepared strategy for communication between providers, which are Culture Collections with status of Microbiological Resource Centre (MRC) and users of microbiological resources. MRC are public resource centers specialized on microbial
raw material. They are capable of providing for users with culturable organisms such as micro-organisms, plant, animal and human cells, replicable parts of these (genomes, plasmids, viruses, cDNAs), as well as databases containing molecular, physiological and structural information relevant to these collections and related bioinformatics.
ELECTROTAXIS AND DUROTAXIS INFLUENCE THE BEHAVIOUR OF SINGLE CELL PSEUDOMONAS AERUGINOSA

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Background

P. aeruginosa motility plays a pivotal role in the colonization of surfaces, especially during the formation of antibiotic resistant biofilm structures. Endogenous electrical signals represent an important signaling mechanism that guides cell migration during wound healing, ultimately regulating the direction of cell movement. We report an approach to study the motility of single P. aeruginosa cells in microfluidic channels possessing different structural geometries, all with the flexibility of being able to manipulate chemical concentration gradients and electric fields to investigate changes in motility in response to specific stimuli.

Objectives

This study is designed to examine the interplay between surface rigidity, mechanical, and electrical cues to pave the way for improvements in the design of anti-fouling surfaces for biomedical applications and to identify new ways to inhibit bacterial biofilm growth through motility restriction.

Methods

Three different microfluidic devices with varying geometries (circular, semi-circular shell, and square cut out respectively) were developed using soft-lithography and nanofabrication tools for the analysis of durotaxis and galvanotaxis of single cell P. aeruginosa.
Conclusions

With the use of fabricated microfluidic platforms, we studied the impact of spatial restrictions, rigidity of the substrate medium, and the impact of external electrical fields on Pseudomonas aeruginosa motility. Spatial restrictions resulted in a significant reduction in single-cell velocity and the distance travelled, but did not affect the trajectory directness. Pseudomonas aeruginosa single cells showed a preference for an environment with increased stiffness, which was independent of the time of exposure to the electric field.
MARINE-DERIVED PENICILLIUM IN KOREA: DIVERSITY, ENZYME ACTIVITY, AND ANTIFUNGAL PROPERTIES
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Background

In Korea, approximately 100 Penicillium species have been recorded. Many of these species were isolated from soil, and some were found to be associated with post-harvest diseases of plant products. However, the diversity of marine-derived Penicillium in Korea is poorly understood relative to terrestrial species.

Objectives

We explored the diversity of marine-derived Penicillium in Korea by isolating Penicillium species from various marine substrates and identifying them using a multigene phylogenetic approach. We also evaluated the biological activity (extracellular enzyme activity and antifungal activity against the plant pathogens) of the strains.

Methods

The diversity of marine-derived Penicillium from Korea was investigated using morphological and multigene phylogenetic approaches (ITS, BenA, RPB2). We tested for the extracellular enzyme activity of alginase, endoglucanase, and beta-glucosidase, and antifungal activity against two plant pathogens (Colletotrichum acutatum and Fusarium oxysporum).

Conclusions

A total of 184 strains of 36 Penicillium species were isolated, with 27 species being identified. The most common species were P. polonicum (19.6 %), P. rubens (11.4 %), and P. chrysogenum (11.4 %). The diversity of Penicillium strains isolated from soil and marine macroorganisms was higher than the diversity of strains isolated from seawater. While many of the isolated strains showed alginase and beta-glucosidase activity, no endoglucanase activity was found. More than half the strains (50.5 %) showed antifungal activity against at least one of the plant pathogens tested. The
relatively high proportion of strains that showed antifungal and enzyme activity demonstrates that marine-derived *Penicillium* have great potential to be used in the production of natural bioactive products for pharmaceutical and/or industrial use.
Background
In fish farming the benefits of probiotics have been usually inferred appraising the effects observed on the host, namely growth, survival and immune response, and not through the direct assessment of probiotics dynamics and role in the host gut microbiota. Currently, there is the urgency to develop accurate tools to assess and weigh up the dynamics of autochthonous bacteria and probiotics in the gut microbiota, in order to better understand the effects of probiotics, prebiotics and synbiotics in aquaculture production.

Objectives
This work aimed to identify novel taxa-specific DNA markers for \textit{B. licheniformis}, one of the most used probiotics in aquaculture, and develop culture-independent methods for detection and quantification of these bacteria and understand their modulation role of fish-gut microbiota.

Methods
Several putative \textit{B. licheniformis}-specific DNA markers were selected using the
CUPID and Insignia databases. These utilities, combined with comprehensive genomic studies and dedicated experimental validation, allowed the selection of four DNA markers used for detection and quantification of *B. licheniformis* by multiplex PCR and qPCR, respectively, in digesta samples of fish gut samples.

**Conclusions**

The selected DNA markers were successfully applied for detection and quantification of *B. licheniformis* in fish gut. These results are a contribution for a better understanding of *B. licheniformis* dynamics in gut microbiota of farmed fish in response to different diets, but also during disease outbreaks. This knowledge is an added-value to optimize and implement best practices for aquaculture.
ICLIKVAL: COMMUNITY RESOURCE FOR CURATING THE VAST WEALTH OF MICROBIOLOGY-RELATED LITERATURE THROUGH THE POWER OF CROWDSOURCING

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Background

There are currently over 24 million citations in PubMed, including many related to microbiology research. Searching this vast resource does not always give desirable or complete results due to several factors. Ideally, every citation should include a complete set of keywords that describe the original article in detail; however, this would require countless hours of manual curation.

Objectives

Our goal is to make manual curation 'fun', social, and self-correcting, thus enriching resources like PubMed so that users are able to extract more valuable and relevant results. We developed a web-based open-access tool for manual curation of PubMed articles, and other media types, using a crowdsourcing approach. We encourage the use of ontology terms and support them as auto-suggest keyword terms, but we do not restrict users to these so as not to impose any limitations on the annotation types.
Non-English annotation is also supported.

**Methods**

We constructed a cross-browser and platform-independent application using a NoSQL database. Users perform searches, mark articles for review, load PDFs into the viewer, select annotations (values) within the text, and add appropriate keywords (keys). Article-specific comments can be made, key-value pairs can be edited and rated, live chats between users can be conducted, annotations can be added via Twitter, etc.

**Conclusions**

The more annotations that accumulate in the database the more reliable the results. We implemented a REST API to make the annotations easily accessible to the research community. We hope this will become the default resource for community-based curation of all online microbiology-related and scientific literature.
THE RISK OF SERIOUS INFECTIONS WITH THE USE OF RITUXIMAB IN RHEUMATOID ARTHRITIS. RESULTS FROM A META-ANALYSIS OF RANDOMISED CLINICAL TRIALS.

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Background

The association between serious infections (SINFs) and the use of rituximab in patients with non-Hodgkin’s lymphoma has been clearly confirmed; but, uncertainty remains about the association of SINFs with rituximab in rheumatoid arthritis (RA).

Objectives

Our study, a systematic review and meta-analysis (SR&MA), aimed to provide reliable assessment of the risk of SINFs and rituximab in RA (RTX-RA).

Methods

This SR&MA was registered with the PROSPERO database (CRD42014015655) as protocol for a complete evaluation of the safety profile of RTX-RA. The search strategy involved randomised clinical trials using rituximab in RA, published from January 1990 to December 2014 in Medline, EMBASE and Cochrane Library databases. The outcomes evaluated were: the number of adverse events and SINFs reported as outcomes of interest. Odds ratio (OR) analysis, 95% confidence intervals (CI95%) and \( p \) values (chi-squared) were calculated; heterogeneity was assessed using the \( I^2 \) test. Sub-analysis by doses (500 vs 1000 mg) and type of infection were performed.

Results: Eight publications (3,272 subjects) were selected for review and 5 publications (2,249 subjects) were included in the meta-analysis; the number of events of SINFs in rituximab+ methotrexate (RTX-MTX) group vs control (placebo+ methotrexate) were 35/1,465 vs 27/784. We found no evidence of association between RTX-MTX and SINFs (OR 0.64; CI95% 0.39-1.04; \( p \)=0.74; \( I^2=0.000 \)). Similarly, our sub-analysis by RTX doses (500 vs 1000 mg) and type of infection, showed no evidence of association.
Conclusions

Our results suggest no evidence of association between RTX-MTX in RA and SINFs. Further studies are needed to confirm our findings.
AUDIT OF NOTIFIABLE DISEASE REPORTING IN A GENERAL MEDICINE DEPARTMENT OF A DISTRICT GENERAL HOSPITAL IN THE UK

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Background

"The purpose of notification [of infectious diseases] is to enable the prompt investigation, risk assessment and response to cases...that present a significant risk to human health. Notification has the secondary benefit of providing data for use in…epidemiological surveillance..."¹ Doctors in England have a duty to notify diseases as a legislative requirement.

Objectives
To establish whether general medical doctors in a hospital in London are appropriately reporting notifiable diseases according to 2010 health protection guidance.

Methods

Notifiable diseases managed in adult secondary care were selected: acute meningitis, invasive group A streptococcus, enteric fever, meningococcal septicaemia, malaria.

Adult patients diagnosed with these diseases between May and October 2013 were identified via the microbiology database and clinical coding. The notes were reviewed for documentation of discussion with public health or the notification form, and the time this occurred relative to diagnosis/clinical suspicion.

Conclusions

24 cases of notifiable diseases were identified and of these 17 notes were located (7 malaria, 9 viral meningitis, 1 Salmonella typhi). There was 1 documented notification (6% of 17 notes), with no identifiable date.

Possible causes of the low notification rate include: poor awareness among doctors that it is their responsibility, lack of knowledge which diseases are notifiable, uncertainty which clinician/clinical team has responsibility for notification, and difficulty in locating forms on the hospital intranet.
Measures to improve notification include implementation of electronic notification reminders with results, staff education and more easily accessible information/notification form on intranet.

CHARACTERIZATION OF THE LIPOPOLYSACCHARIDE TRANSPORT MACHINE IN ESCHERICHIA COLI: IN SEARCH OF A FUNCTION FOR THE ELUSIVE COMPONENT LPTC

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Background
The lipopolysaccharide (LPS) transport (Lpt) system is responsible for transferring LPS from the periplasmic surface of the inner membrane (IM) to the outer leaflet of the outer membrane (OM), where it plays a crucial role in OM selective permeability. In E. coli seven essential proteins form the Lpt trans-envelope complex: LptBFG form the IM ABC transporter; LptDE form the OM translocon for final LPS delivery; LptC, an IM-anchored protein with a periplasmic domain, interacts with the IM ABC transporter, the periplasmic protein LptA, and LPS(1). Although essential, LptC can tolerate several mutations and its role in LPS transport is unclear.

Objectives
To clarify the functional role of LptC in the Lpt machine.

Methods
By plasmid shuffling we obtained viable mutants missing lptC; genome sequencing of ΔlptC mutants revealed single amino acid substitutions at a unique position (*) in the IM component lptF; in complementation tests, lptF* mutants suppress lethality of both ΔlptC and lptC conditional expression mutants.

Conclusions
Our data show that a specific mutation in LptF can compensate the lack of LptC and suggest that LptC may serve as a chaperon of the Lpt machine assembly and/or activity rather than an essential structural component. Studies are in progress to understand the structure and function of the six-protein Lpt* system.

A NOVEL SCENARIO OF OPRF REGULATION IN PSEUDOMONAS AERUGINOSA

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Background

OprF, the major porin of *Pseudomonas aeruginosa*, has an immunogenic potential and a large set of information regarding its functions, including its role in the outer membrane structure, interaction with lipopolysaccharide and association with virulence and quorum sensing. In the prototype strain PAO1, transcription of *oprF* was reported to be dependent on sigma 70 and the alternative ECF sigma factor SigX.

Objectives

As we have shown that SigX overexpression in PA14 leads to a growth defect, we assessed *oprF* transcription regulation in order to verify if this effect is due to OprF overexpression and to characterize *oprF* regulation in PA14.

Methods

*oprF* promoter activity was determined via -*lacZ* reporter fusions integrated in the chromosome or by relative *oprF* mRNA levels (qRT-PCR) in wild-type PA14, *sigX* null mutant or in PA14 overexpressing *sigX* from an inducible promoter in a multicopy plasmid (ALB04 strain). *oprF* and *sigX* promoters activity was also assessed in the PA14*rpoS* mutant by β-galactosidase assays.

Conclusions

Surprisingly, we found that all *oprF-lacZ* promoters activity decreased in ALB04 and the amount of OprF did not increase in a proteomic analysis. The *sigX* null mutant has wild-type levels of *oprF* mRNA, contrasting to other SigX-induced genes. We have also found that transcription of *oprF* occurs mainly via the region containing the putative sigma 70-dependent promoter. Although *oprF* transcription regulation is independent of SigX in PA14, both *oprF* and *sigX* transcription is dependent on the general stress sigma factor RpoS. The data presented here discloses a novel
scenario of *oprF* regulation in *P. aeruginosa*. 
TRIGGER ENZYME PEPA FROM E. COLI, A TRANSCRIPTIONAL REPRESSOR THAT GENERATES POSITIVE SUPERCOILING

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Background
Hexameric \textit{E. coli} Leucine-aminopeptidase A (PepA) is a trigger enzyme endowed with catalytic activity and DNA-binding properties prominent in the resolution of ColE1 and pSC101 plasmid multimers and transcriptional regulation of the \textit{carAB} operon, encoding the unique carbamoylphosphate synthase of \textit{E. coli}. Previous studies by DNase I footprinting and atomic force microscopy (AFM) had both indicated a pronounced deformation of the \textit{carAB} control region upon PepA binding, suggestive of DNA wrapping. On the basis of this observation and previous work, PepA was believed to play a major, although merely architectural role in the formation of the synaptic complexes involved in site-specific DNA recombination and in the elaboration of a higher order regulatory protein-\textit{carP1} DNA complex.

Objectives
Here we further investigate the molecular mechanism of PepA-mediated transcriptional regulation of the \textit{carP1} promoter.

Methods
To this aim we use single-round \textit{in vitro} transcription and DNA topology assays.

Conclusions
The \textit{in vitro} transcription assays performed with supercoiled template and purified components demonstrate that PepA is a repressor in its own right, which specifically inhibits transcription initiation at \textit{carP1}. Furthermore, DNA topology studies performed on artificial DNA mini-circles by means of various topoisomerases with different substrate specificities demonstrate that PepA binding induces positive supercoiling. Such topological changes may serve as a regulatory mechanism that allows swifter response and multi-layered control of promoter activity in concert with other regulatory components known to be involved in the control of \textit{carP1}.
FEMS-1148
Bacterial cell biology

OFF THE WALL: FROM FILAMENTOUS GROWTH TO PRIMORDIAL CELLS AND BACK AGAIN
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Background
- Objectives
- Methods
- Conclusions

Streptomyces are filamentous bacteria that grow by apical tip extension. This process is orchestrated by the tropomyosin-like protein DivIVA, which is present at hyphal tips. DivIVA interacts with various proteins, among which the cellulose synthase-like protein CslA. This protein synthesizes a β-(1,4)-glycan, which is thought to protect growing apices that are continuously being remodeled. To obtain further insight in the role of DivIVA and CslA in polar growth and morphogenesis, we have recently generated so-called Streptomyces L-forms that can grow without peptidoglycan. As a consequence, such cells are round and lack any obvious form of polarity. L-form cells have recently been suggested to resemble primordial cell, based on the observation that their growth and proliferation do not require the canonical cytoskeletal or cell division proteins. Instead, their proliferation can merely be explained by physical processes. However, our work on Streptomyces L-forms suggests that these cells require glycans, such as those formed by CslA, for their growth. Such glycans might have served for protection of early life forms, before the modern cell wall was invented. We have recently isolated an L-form mutant strain, which readily switches back and forth between mycelial and L-form growth. This mutant with the capability to re-synthesize peptidoglycan is crucial to understand which genes play an essential role in proliferation of L-forms, but also to unravel the mechanism underlying filamentous growth.
THE ESCHERICHIA COLI MEMBRANE PROTEIN INSERTASE YIDC ASSISTS IN THE BIOGENESIS OF Penicillin Binding Proteins

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Background
Membrane proteins need to be properly inserted and folded in the membrane in order to perform a range of activities that are essential for the survival of bacteria. The Sec translocon and the YidC insertase are responsible for the insertion of the majority of proteins into the cytoplasmic membrane. YidC can act in combination with the Sec translocon in the insertion and folding of membrane proteins. However, YidC also functions as an insertase independently of the Sec translocon for so-called YidC-only substrates. In addition, YidC can act as a foldase and promote the proper assembly of membrane protein complexes.

Objectives
Here, we investigate the effect of Escherichia coli YidC depletion on the assembly of Penicillium Binding Proteins (PBPs), that are involved in cell wall synthesis.

Methods
Active PBPs were assessed by substrate binding and total protein amount was determined by immunoblotting.

Conclusions
YidC depletion hardly affects the total amount of the specific cell division PBP3 (FtsI) in the membrane, but the amount of active PBP3 is strongly reduced. Similar reductions in the amounts of active PBP2 and PBP4 were observed, while the levels of active PBP1A/1B and PBP5 were essentially similar. PBP1B and PBP3 disappeared from higher Mw bands upon YidC depletion, indicating that YidC might play a role in PBP complex formation. Taken together, our results suggest that the foldase activity of YidC can extend to the periplasmic domains of membrane proteins.
Background

Neisseria gonorrhoeae is an obligate human pathogen causing gonorrhea. It mainly employs homologous recombination to repair double stranded breaks in DNA. The involvement of mismatch repair helicase, UvrD in homologous recombination is debatable till date. Evidences have implied that it could be an anti-recombinase or promote RecFOR pathway. In this study, a detailed biochemical characterization of UvrD from N. gonorrhoeae and its interaction with RecJ exonuclease (NgoRecJ) which is involved in resection of DNA ends during RecFOR pathway of recombination has been carried out.

Objectives

1. To biochemically characterize UvrD from Neisseria gonorrhoeae (NgoUvrD).
2. To study the physical interaction between NgoUvrD and NgoRecJ exonuclease.
3. To elucidate the functional role of this interaction, if any.

Methods

Both the proteins were purified by Ni$^{2+}$-NTA followed by Heparin-Sepharose chromatography. The unwinding and exonuclease activities were monitored using gel based assays. ATPase assays were carried out using TLC. Protein-protein interaction studies were done using Far-western analysis.

Conclusions

Biochemical analysis suggested that NgoUvrD behaves as a dimer in solution, exhibits 3’-5’ polarity on ssDNA and can unwind blunt end duplex DNA as well as different recombination intermediates such as overhang containing DNA and Holliday junction in ATP dependent manner. It was observed that NgoRecJ and NgoUvrD
physically interact with each other. Interestingly, the ATPase activity of NgoUvrD shows 5-fold stimulation in presence of NgoRecJ. This observation for the first time shows that UvrD in complex with RecJ exonuclease might be involved in the initial step of RecFOR pathway, thereby promoting recombination.
Background
Programmed cell death (PCD) in bacteria has long been controversial due to the belief that only multicellular organisms would benefit from this kind of altruistic behavior. However, over the past decade, compelling experimental evidence has established a consensus that such pathways exist in bacteria.

Objectives
Recently, we discovered that expression of a mutant isoform of an essential GTPase, ObgE, causes rapid cell death in *Escherichia coli*. The physiological changes that occur upon expression of this mutant protein (ObgE*) and the genetic basis of this cell death pathway were investigated.

Methods
Besides a clear loss of viability, fluorescence microscopy and FACS analysis revealed that a large array of apoptotic markers – including chromosome condensation, DNA fragmentation, loss of membrane potential and exposure of phosphatidylserine on the cell surface – can be found upon expression of ObgE*, indicating that ObgE* triggers a PCD pathway. Previous reports of bacterial PCD attribute essential roles to either RecA and its function in the SOS response or to the toxic effect of the mazEF TA module. However, by analysing viability and physiological changes upon ObgE* expression in recA and mazEF mutant strains, all currently known bacterial PCD mechanisms could be excluded.

Conclusions
ObgE* triggers a PCD pathway in *E. coli* that differs fundamentally from all other previously described bacterial PCD pathways. Further research into the genetic and molecular basis of this new PCD mechanism may prove useful in unraveling the function and mechanisms of genetically encoded cell death pathways in bacteria.
MUTATIONAL ANALYSIS OF LPTA, AN ESSENTIAL LPS-TRANSPORT PROTEIN IN ESCHERICHIA COLI

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Background

Lipopolysaccharide (LPS) is the major outer leaflet constituent of the Gram-negative bacteria outer membrane (OM). In E. coli LptA protein is a periplasmic component of the LPS transporter (Lpt) made of seven components (LptABCDEF), which ferries LPS from the inner membrane (IM) to OM. LptA interacts with LptC and chaperons LPS through the periplasm. The crystal structure of LptA has been solved and some residues involved in binding LPS and other Lpt proteins have been identified.

Objectives

1. To characterize LptA structure-function.
2. To identify interactions between genes/proteins implicated in OM functionality.

Methods

1. We generated by site-directed mutagenesis lptA41, a quadruple mutant in four conserved amino acids potentially involved in LPS or LptC binding; the mutant exhibited increased sensitivity to hydrophobic compounds (Shc phenotype). This suggests that lptA41 impairs LPS transport thus leading to OM permeability defects.

2. We selected suppressors of lptA41 Shc phenotype and sequenced the genomes of two mutants (Sup102 and Sup103) to identify the suppressing mutations.

Conclusions

Sup102 harbors a small in-frame deletion in vacJ, which codes for a putative OM lipoprotein involved in maintaining OM asymmetry. Sup103 harbors an additional amino-acid substitution (M112I) in LptA41 (lptA42 allele) and a nonsense mutation in mdoH, a gene encoding a glycosyltransferase involved in periplasmic membrane-derived oligosaccharides synthesis.
These results reveal two different strategies adopted by the cell to overcome OM defects caused by defective LPS transport.
Background

Accumulating evidence has shown that Hfq is a RNA chaperone involved in post-transcriptional regulation, mediating the interaction of small RNAs with their target mRNAs. However, besides its regulatory role in the translation of bacterial mRNA, in \textit{Escherichia coli}, Hfq is also a DNA-binding protein and is associated with nucleoid DNA. Bcc bacteria are among the few prokaryotes that encode in their genomes two distinct and functional Hfq-like proteins, the 79 aa Hfq and the 188 aa Hfq2. Both proteins bind RNA, however bioinformatics predictions suggested that Hfq2 is able to bind DNA by means of its C-terminal domain.

Objectives

The present study aims to unveil the biological significance of Hfq2 C-terminus predicted ability to bind to DNA.

Methods

A DNA binding domain was bioinformatically predicted for Hfq2 and DNA binding assays were performed using purified His-tagged Hfq or Hfq2 and the \textit{araC} promoter of pMLBAD plasmid. Electrophoretic Mobility Shift Assays indicates that Hfq2 protein, but not Hfq, is able to bind this DNA fragment. Hfq2 derivatives with complete, partial and absent C-terminus extension were produced to confirm if this region confers the ability to bind DNA. To identify the DNA sequences to which Hfq2 specifically binds, a ChIP-seq strategy is currently being performed, using \textit{B. cenocepacia} J2315 fragmented total DNA.

Conclusions

Contrasting with Hfq, a putative DNA binding domain was identified in Hfq2 and our results showed that this protein is able to bind DNA. Ongoing work is being performed to identify the DNA sequences and the specificity of DNA-Hfq2 interactions.
KNOCKOUT OF MINC LEADS TO BRANCHING IN XANTHOMONAS CITRI

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Background

Xanthomonas citri (Xac) is the causal agent of citrus-canker, a disease that affects citrus plants worldwide, leading to low productivity, culminating in considerable economic losses to orange growers. There is no treatment for citrus-canker, where the eradication of plants is the only reliable method to prevent the spread of Xac to areas considered free of the pathogen. Recently, we reported on the use of new cell-division inhibitors as an alternative to prevent Xac growth, and plant colonization (Silva et al., 2013 JBacteriol 195:85). To further extend the characterization of cell division in Xac, we started the investigation of Xac minC.

Objectives

To study minC encoded by Xac and its function in site division selection at cell division

Methods

The minC gene was deleted using allele exchange. The knocked-out strain was complemented with a replicative vector carrying minC under the control of the arabinose promoter. Pathogenicity tests were carried out by infiltration of sweet orange leaves (Citrus sinensis (L.) Osbeck). Cell morphology was evaluated using DIC microscopy.

Conclusions

Xac deleted for minC exhibited the classic delta-minC phenotypes: production of minicells, cell filamentation, and asymmetric/misplaced division constrictions along the rods. Surprisingly, Xac delta-minC produced branches that resembled those observed for Escherichia coli deleted for PBP5 and other low-molecular-weight penicillin-binding proteins. Complementation of Xac delta-minC with minC supplied in trans restituted the wild-type division phenotype. Irrespective of the presence or absence of minC, Xac was competent to colonize the host citrus and produce disease symptoms. Altogether, data shows that minC is involved in cell division in Xac.
Background

The CsrA protein (Carbon Storage Regulator A) is a global post-transcriptional regulator controlling carbon fluxes and group behaviors in bacteria. Our preliminary SEM analyses revealed that an E. coli mutant deleted for the csrA gene displays a strongly altered morphology. Cells are of small size as compared to the wild-type cells and present an irregular surface and blebs.

Objectives

Cell envelope is the first line target for physicochemical injuries arising from the environment. This complex structure also supports fundamental processes such as energy production and nutrient acquisition. To preserve envelope homeostasis bacteria have evolved intricate stress response pathways. The objective is to evaluate whether envelope stress responses are induced in ΔcsrA mutant and if it is the case, what are the underlying molecular mechanisms.

Methods

Expression of effectors belonging to the 5 well-described envelope stress response pathways (bae, psp, cpx, sigmaE, rcs) were analyzed by RT-qPCR. Envelope integrity was evaluated by measuring sensitivity to specific compounds.

Conclusions

We have shown that in the ΔcsrA mutant, the Psp and Rcs pathways are constitutively induced. Rcs pathway monitors outer membrane integrity while Psp that of the inner membrane. This mutant also shows a high sensitivity to deoxycholate and vancomycin, indicating an increased permeability of the outer membrane. In addition, the ΔcsrA mutant accumulates high ROS concentration, which might participate to psp activation. ROS accumulation might thus represent the link between metabolism and envelope defects. This hypothesis is currently under investigation.
IDENTIFICATION OF THE MULTI-COMPONENTS DNA UPTAKE SYSTEM OF NATURALLY TRANSFORMABLE LEGIONELLA PNEUMOPHILA USING TRANSCRIPTOMIC ANALYSIS.

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Background

Natural transformation is a major mechanism of horizontal gene transfer that is already well-studied in historical Gram-positive models whereas knowledge on Gram-negative bacteria remains sparse. Legionella pneumophila is a Gram-negative bacterium that develops, under specific growth conditions or stress, competence for natural transformation, a genetically programmed state allowing bacteria to take up free exogenous DNA and integrate it in their genome. The L. pneumophila genome shows a large repertoire of virulence-associated eukaryotic-like genes. Natural transformation offers a plausible route for acquisition of foreign genes contributing to the emergence of this pulmonary pathogen.

Objectives

The DNA uptake system allowing L. pneumophila to take up DNA and recombine it into its chromosome remains unknown. The aim of our study was to identify this system and determine its contribution to the infectious process.

Methods

We used a transcriptomic analysis to identify genes up-regulated in an hypercompetent mutant of L. pneumophila. We performed a systematic targeted mutagenesis of these up-regulated genes to test their involvement in transformability and DNA uptake. We used immunofluorescence microscopy to visualize components of the DNA uptake system.

Conclusions
Our work allowed us to identify the main components of the *Legionella* DNA uptake system which involves a type IV transformation pilus dedicated to natural transformation. We propose a model of the DNA uptake system that unexpectedly relies on the actin-like protein MreB. These progresses pave the way for a more detailed analysis of the DNA uptake system dynamics and more broadly for a better understanding of Gram-negative bacteria transformation systems.
Background

TnrA is a master transcriptional regulator of nitrogen metabolism in *Bacillus subtilis*. Its activity was suggested to be regulated by binding to the feedback-inhibited glutamine synthetase (GS). Recently we showed that TnrA also binds to the PII-like GlnK protein.

Objectives

Understanding of TnrA activity regulation via interaction with GlnK or GS

Methods

*In vivo* cross-linking, Surface Plasmon Resonance, ITC

Conclusions

We found that *in vivo* TnrA binds both proteins, but in contrast to the suggested mechanism, TnrA binds GS also under nitrogen-poor conditions. SPR analysis demonstrated that GS in the absence of feed-back inhibitors efficiently binds TnrA, but 1 mM glutamine strengthens the binding 2-fold. By contrast, ATP strongly repressed TnrA-GS interaction, whereas 1 mM glutamine completely neutralized this negative effect. In presence of L-methionine sulfoximine (MSX), a transition-state analogue of GS, GS–TnrA interaction was also stimulated. However, in presence of MSX and ATP, where MSX is phosphorylated and irreversibly fixes GS in transition state, GS binding to TnrA was completely abolished. Isothermal titration calorimetry revealed competitive binding of glutamine and ATP to GS. Therefore, we suggest two conformations of GS for TnrA interaction: (1) a TnrA-binding "N-state", stabilized by glutamine or its analogue MSX and (2), a non-binding "A-state", which reflects the conformation of the enzyme in the catalytic transition state or bound with ATP in
absence of glutamine. We propose that GS changes between the “N-state” and “A-state”. In vivo, this transition depends on the intracellular glutamine:ATP ratio and thus regulates the amount of active TnrA.
FEMS-3056
Bacterial cell biology

A VIBRIO VUNIFICUS ORTHOLOG OF ESCHERICHIA COLI RRAA CONTRIBUTES TO DIFFERENTIAL RNASE E-MEDIATED MRNA DECAY
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Background
Endoribonuclease E (RNase E) plays an important role in the degradation and processing of RNAs in Escherichia coli. Expression levels and enzymatic activity of RNase E are tightly controlled by self-cleavage of rne mRNA and regulators of RNase activity, RraA and RraB. The marine pathogenic bacterium Vibrio vulnificus also contains homologs of RNase E and RraA, designated as RNase EV, RraAV1, and RraAV2.

Objectives
In this study, we show that RraAV1 actively inhibits the enzymatic activity of RNase EV on a subgroup of substrate RNAs.

Methods
Notably, RNase EV cleavage on rne mRNA encoding RNase EV was greatly affected by RraAV1 in vitro and in vivo, contributing to autoregulation of both expression and activity of RNase EV.

Conclusions
Our findings suggest that RraAV1 plays an active role in differential RNase E-mediated mRNA decay.
Background

Gram-negative bacteria produce outer membrane vesicles (OMVs) containing biologically active proteins. Despite of extensive information about OMVs biogenesis available, this process is still not enough clear. Gram-negative bacterium Lysobacter sp. XL1 forms OMVs containing bacteriolytic endopeptidase L5, – one of five bacteriolytic enzymes secreted by this microorganism. Knowledge about OMVs biogenesis by bacteria of Lysobacter genus is rather restricted.

Objectives

The objective of our work was the study of factors causing biogenesis of OMVs produced by Lysobacter sp. XL1.

Methods

Methods were used: fractionation of OMVs using sucrose density gradient centrifugation, electronic microscopy including immunocytochemistry with the protein A-gold, SDS-PAG electrophoresis, Western blotting assay, protein and 2-keto-3-deoxyoctonate assays, thin-layer chromatography, determination of lytic OMVs action.

Conclusions

Enzyme L5 was found to be localized inside of the lightest OMVs fraction which differed in protein composition from other fractions. The L5 protein was revealed at certain loci of bacterial outer membrane (OM). OMVs were produced from these loci. Thereby, secreting enzyme L5 took part in biogenesis of OMVs. OM was shown to contain cardiolipin, phosphatidylethanolamine, phosphatidylglycerol. OMVs contained the only one major phospholipid, – cardiolipin. Thus, OMVs of Lysobacter sp. XL1
were formed from loci enriched with cardiolipin. It has been offered the model of vesicles biogenesis produced by *Lysobacter* sp. XL1. Besides, OMVs containing protein L5 were shown to lyze broad range of opportunistic and pathogenic bacteria including multidrug-resistant strains. The data obtained are the basis for development of artificial vesicular structures – liposomes – containing bacteriolytic enzymes of *Lysobacter* sp. XL1 as a new effective antimicrobial preparation.
Background

Despite its essential role as a trace element, copper becomes toxic at higher concentrations, prompting bacteria to develop a tight regulation of copper homeostasis in order to survive.

Objectives

The alphaproteobacterium Caulobacter crescentus lives in poor and contaminated environments and gives rise to two distinct cell types upon cell division: a motile swarmer cell and a sessile stalked cell. The swarmer cell is thought to seek optimal environments in order to differentiate into a stalked cell, which will in turn replicate its DNA and divide. This dimorphic cell cycle likely provides a better adaptation to various stresses such as an increase of copper concentration.

Methods

We found that the swarmer and the stalked cells respond differently to a toxic copper concentration. Copper slows down swarmer to stalked cell transition and the further initiation of DNA replication, whereas copper-treated stalked cells undergoes a normal DNA replication. Accordingly, copper concentration in swarmer cells is 3 fold higher than in the stalked cells. We showed that copper homeostasis in the stalked cells is ensured by the conserved PcoAB\textsubscript{Cc} system, which is also able to sustain copper detoxification in the swarmer cells when constitutively expressed in trans, thereby restoring a proper swarmer to stalked cell transition.

Conclusions

Both C. crescentus cell types regulate copper homeostasis differently in order to progress throughout the cell cycle.
Background
Endoribonuclease G (RNase G) is involved in rRNA processing and degradation of a subgroup of mRNAs in *Escherichia coli*. However, little is known about physiological role of this enzyme.

Objectives
We observed RNase G expression-dependent alterations in the survival rate of *Salmonella* Typhimurium cells in both macrophages and mice. These alterations were strongly associated with expression levels of *rbs* mRNA encoding ribose transport system, which was identified as a substrate of RNase G.

Methods
Analysis of polycistronic *rbs* mRNA identified two RNase G cleavage sites in the 5'-untranslated region of *rbsA*. The induced *rbs* expression during macrophage infection coincided with decreased rates of bacterial infection and survival.

Conclusions
Our study shows that RNase G mediates a previously uncharacterized pathway that involves ribose transport system as a key factor for the survival and virulence of *S. Typhimurium* in host cells.
IDENTIFICATION AND CHARACTERIZATION OF A Corynebacterium glutamicum Mutant Impaired in the Biosynthesis of Peptidoglycan

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Background

Bacteria belonging to Corynebacteriales, an order of actinomycetes group that includes corynebacteria, mycobacteria, nocardia and rhodococci possess an atypic, complex multilayered envelope. This envelop is composed of a heteropolymer of peptidoglycan (PG) and arabinogalactan (AG) covalently associated to an outer membrane mostly composed of mycolic acids. We explored the biogenesis of Corynebacteriales cell envelope by using a genome-wide transposon mutagenesis approach targeting Corynebacterium glutamicum. For this purpose, we developed an effective immunological screen that allows us to rapidly identify bacteria exhibiting an altered cell envelope.

Objectives

We wanted to identify missing actors of the Corynebacteriales envelope biosynthesis pathways. One interesting mutant of our library was chosen and characterized in this study.

Methods

We identified the gene inactivated by the transposon and constructed a strain in which this gene was deleted. We performed analyses of the main cell wall compounds of this mutant strain, i.e. PG, AG and mycolic acids. A physiological study was also conducted and in particular a screen for the sensitivity of this mutant to various antibiotics.

Conclusions

Our analyses revealed that this mutant was impaired in the PG biosynthesis pathway and more specifically in a modification of this polymer that also occurs in mycobacteria species and in other Corynebacterium pathogens species such as Corynebacterium jeikeium. Lack of this modification led to morphologically-altered
cells with high susceptibility to lysozyme and to antibiotics of the b-lactam family.
TOWARDS COMPREHENSION OF THE ROLE OF KEY GENES DURING LACTOBACILLUS CASEI ESTABLISHMENT IN THE GUT.

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Background

Beneficial interactions between gut microbiota and the host have been revealed but little is known at the molecular level. To identify bacterial factors that account for these symbiotic interactions, we recently carried out reverse genetics of Lactobacillus casei which allowed identification of 47 key genes for L. casei establishment in the gut (Licandro-Seraut et al. 2014. PNAS).

Objectives

Now, we aim to decipher the functions encoded by these key genes in the gut context.

Methods

Some of the L. casei mutants have been investigated individually with regard to their morphology, metabolism, gene regulation...

Conclusions

A mutant for the predicted cysteine synthase was investigated to demonstrate the metabolic role of the cysteine synthase. Also, it was less resistant than the wild type to several stresses. Three mutants for three genes in an operon encoding a two-component system and a penicillin-binding protein were characterized. Analysis of primary data suggests that these genes are involved in the cell surface modulation and particularly in the regulation of peptidoglycan hydrolases which are required for the bacteria protection in the intestinal environment.
Role of PprM in catalase gene expression of the radiation-resistant bacterium, Deinococcus radiodurans

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Background: Deinococcus radiodurans (D. radiodurans) is a poly-extremophilic organism, capable of tolerating a wide variety of different stresses such as gamma/UV radiation, desiccation, and oxidative stress. It is known that Pprl (DR0167) is a global regulator, which is essential for its extreme resistance, and cold shock protein homologue PprM (DR0907) is presumably controlled by Pprl. Pprl mutation decreases the catalase activity of D. radiodurans, but, the effect of PprM on catalase has not been studied.

Objectives: We investigated the role of PprM, which is involved Pprl-mediated signal transduction, in catalase gene expression.

Methods: First, we performed an in-gel catalase activity assay to observe activity of three catalases (DR1998, DRA0146, and DRA0259) independently in wild type, pprM, pprl, and pprl/pprM double mutant strains. We also constructed catalase mutants of D. radiodurans to identify the catalase that is affected by PprM and purified catalases to check the activity in vitro. The mRNA and protein levels of catalase genes were monitored by using real-time PCR and western blot assays in various mutant strains.

Conclusions: Among three catalases, we confirmed that DRA0146 does not have catalase activity. Deletion of pprM decreased catalase activity and protein level of DR1998, but not its mRNA level. We could not observe an additional reduction of the DR1998 protein in the pprl/pprM double mutants compared to pprl or pprM single mutants. These suggest that Pprl can affect DR1998 via PprM, and PprM may exert its effect on DR1998 at the post-transcriptional level.
Background

Acinetobacter baumannii is an emerging pathogen involved in severe infections and outbreaks mainly occurring in clinical settings. Multi-drug resistance and ability to form biofilm could play a role in its persistence in hospital settings.

The present study analyzes an A. baumannii clinical strain, belonging to the SMAL clone, isolated in an Italian hospital.

Objectives

The aim of this work is to study the effect of glucose, at a concentration similar to human blood, on production of biofilm determinants in A. baumannii.

Methods

Genome sequence and transcriptome analysis were performed by using deep sequencing and RNA-seq techniques (Illumina).

Transcription of selected genes were measured by using transcriptional fusions with the reporter gene lacZ.

Exopolysaccharide (EPS) production was analysed by EDTA extraction and phenol-sulfuric acid quantification.

Lipopolysaccharide (LPS) production was measured using tricine SDS-PAGE.

Cytokine production was tested by ELISA analysis.

Conclusions
Genome sequence of the analysed strain showed identity with the sequence type 78, epidemic in several Mediterranean countries.

Growth in the presence of glucose showed higher EPS production; consistently, transcriptome analysis suggested a deep modification of gene expression in presence of glucose. In particular, several genes involved in EPS and LPS biosynthesis and modification were up-regulated in presence of glucose. Remarkably, *A. baumannii* grown in presence of glucose appears to release LPS, which in turn stimulates cytokine production in macrophages.

Our results suggest that glucose at physiological concentrations might act as a trigger for LPS production and virulence in *A. baumannii*. 
FEMS-1759
Bacterial cell biology

ACTION OF MOLECULAR CHAPERONES DEDICATED TO THE MATURATION OF REDUCTIVE DEHALOGENASES
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Background
Reductive dehalogenases (RDases) are key enzymes in organohalide respiration (OHR) as they catalyze the reduction of halogenated compounds such as tetrachloroethene (PCE), a major environmental pollutant. RDases contain a corrinoid and two iron-sulfur cofactors and require cofactor assembly and maturation prior to transport across the cytoplasmic membrane via the Twin-arginine translocation (Tat) pathway. Recently, a new dedicated Tat molecular chaperone, PceT, has been identified that seems to act in the maturation of the PCE RDase (PceA) of Dehalobacter restrictus and members of Desulfitobacterium hafniense.

Objectives
This work intends to study the molecular action and specificity of PceT and other members of the RdhT chaperone family on the maturation of RDases.

Methods
Heterologous co-expression of the pceA and pceT genes will be investigated in E. coli in various growth and induction conditions in order to evaluate the effect of PceT on the production, stability and cofactor load of PceA. Additional general molecular chaperones will be also considered here. A detail characterization of the protein-protein interaction between PceT and PceA will be obtained by a combination of in vitro biochemical approaches. Site-directed mutagenesis of conserved residues of RdhT proteins should allow identifying key amino acids in the recognition of, i.e. action on, PceA.

Conclusions
Preliminary data already suggested that PceT mainly recognizes the Tat signal peptide of PceA, therefore resembling the paradigmatic Tat specific chaperones (TorD, NapD,...) as they play a major role in quality control of their cognate redox enzyme allowing sufficient time for cofactor assembly and folding.
IDENTIFICATION OF A PUTATIVE CHROMOSOMAL REPLICATION ORIGIN
FROM BDELOVIBRIO BACTERIOVORUS AND ITS INTERACTION WITH THE
INITIATOR PROTEIN DNAA
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Background
Bdellovibrio bacteriovorus is a small Gram-negative, obligate predatory bacterium
that attacks and invades other Gram-negative bacteria, including pathogens such as
Helicobacter pylori or Pseudomonas aeruginosa. Its life cycle consists of two stages –
non-replicative attack phase, wherein predator searches for prey and replicative
growth phase, wherein it actively divides in host periplasm. Initiation is the first and
strictly regulated step of bacterial chromosome replication, which leads to duplication
of the genetic material in bacterial cells prior to their division.

Objectives
Our aim was to identify and characterize the key elements of initiation of chromosome
replication in B. bacteriovorus: origin of chromosomal replication (oriC) and initiator
protein.

Methods
Using in silico analysis, we identified the oriC region, which is located downstream of
the dnaA gene; the B. bacteriovorus oriC (BdoriC) contains eight putative DnaA
boxes. Comprehensive in vitro studies using EMSA, DMS footprinting and SPR
revealed that the DnaA protein specifically binds all eight DnaA boxes. By P1
nuclease assay we localized the DNA unwinding elements (DUE) where DNA
replication starts. In addition, we compared the architecture of the DnaA–oriC
complexes (orisomes) in homologous (oriC and DnaA from B. bacteriovorus) and
heterologous (BdoriC and DnaA from prey, E. coli or P. aeruginosa) systems.
Interestingly, we demonstrated that DnaA proteins from preys (relatively distantly
related from B. bacteriovorus) not only specifically bind BdoriC, but also unwind DNA
at the DUE.

Conclusions
To conclude, we identified the oriC of B. bacteriovorus and characterize in details its
interaction with the replication initiator protein.
Background

*Brucella*, an ALPHA2-proteobacteria, is the etiological agent of brucellosis that generates a worldwide zoonosis. It can infect human and will lead, if not treated, to a chronic infection with severe complications.

To proliferate, bacteria need to get carbon sources from their environment and the phosphotransferase system (PTS) contributed to this goal. It displays regulating functions in carbon metabolism and forms a phospho-relay, ending with the entry and concomitant phosphorylation of a sugar.

*Brucella abortus* possesses a paralogous system called Nitrogen PTS (PTS\textsuperscript{Ntr}), acting the same way except that no sugar entry occurs. Starting from the phosphoenolpyruvate, the phosphoryl group is successively transferred on histidine residues of Enzyme I (E\textsubscript{I}Ntr) (ptsP gene), then to the NPR (ptsO) protein and finally to EII\textsubscript{A}Ntr (ptsN) or EII\textsubscript{A}MAN (ptsM).

Objectives

We were interested in the identification of a link between central metabolism and the *B. abortus* PTS\textsuperscript{Ntr}.

Methods

We have generated the knockout mutant of those 4 protagonists and studied their bacterial growth behavior on media containing different single carbon source. We found that *B. abortus* was mainly able to grow on sugar entering the pentose phosphate pathway. Interestingly, compared with the wild type strain, the ptsP (E\textsubscript{I}Ntr), ptsO (NPR) and ptsN (EII\textsubscript{A}MAN) mutants shows a growth defect on those sugars while the ptsM (EII\textsubscript{A}Ntr) mutant grows better. Moreover, point mutation study on NPR protein shows that phosphorylation state of the histidine residue seems to be the main actor controlling growth behavior.

Conclusions

It suggests a role of the *Brucella* PTS\textsuperscript{Ntr} in the regulation of central metabolism.
ENRICHMENT FOR G1-ARRESTED DAUGHTER CELLS OF BRUCELLA ABORTUS

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Background: In HeLa cells and RAW264.7 macrophages, the facultative intracellular pathogen B. abortus blocks its growth and its cell cycle at the G1 stage during the first hours of infection (M. Deghelt et al.). G1 bacteria are proposed to have an increased ability to invade these host cells. Therefore, it would be interesting to prepare a bacterial population enriched with G1-arrested daughter cells to investigate B. abortus cell cycle in culture and in interaction with the host cells.

Objectives: Searching the best method for synchronizing the cell cycle of B. abortus at the G1 stage. Then, testing G1 bacteria for their potential invasion of host cells.

Methods: Nutrient downshift method was used to produce G1-arrested cells of Sinorhizobium meliloti (De Nisco et al.). Our results indicate that this is not an effective way to synchronize G1-arrested daughter cells of B. abortus. However, culture of B. abortus in rich medium until the early stationary phase resulted in a reproducible enrichment in G1 bacteria (about 70%), as assessed by flow cytometry. Another method called “baby machine”, that immobilize a mixed population of bacteria labelled with NHS-biotin (N-hydroxysuccinimido-biotin) by streptavidin-coated magnetic beads, was set up to recover only G1 bacteria. We expected that the unipolar growth will be resumed on the beads in rich medium, allowing the enrichment of daughter cells at the G1 stage of the cell cycle.

Conclusions: Culture of B. abortus in rich medium yielded a partial and transient enrichment of G1 bacteria. It seems that the unipolar nature of growth is no more detectable when bacteria are labelled with NHS-biotin. Other labelled compounds for “baby machine” method are currently tested and analysed to achieve G1 bacteria of B. abortus.
CHARACTERIZATION OF LMO1521 AND LMO2591 MUREIN HYDROLASES FROM LISTERIA MONOCYTGENES

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Background

*L. monocytogenes* is a gram-positive, foodborne pathogen causing listeriosis with a high mortality rate (1). This species is widespread in nature which is connected with a high number of surface proteins. Very important class of these proteins are murein hydrolases (autolysins), involved in several crucial processes, including: cell growth, turnover of cell wall components, cell separation and division, biofilm formation, protein secretion, autolytic activity of some antibiotics and pathogenicity (2). This work was supported by a grant from the National Center of Science 2013/09/B/NZ6/00710

Objectives

Determination of the activity and substrate specificity of two autolysins from *L. monocytogenes*: Lmo1521 and Lmo2591.

Methods

Lmo1521 and Lmo2591 proteins with C-terminal hexa-His-Tag were expressed in *E. coli* BL21 using expression vector pET-28a (Novagen). These proteins were purified on Ni-NTA Agarose column (Qiagen), and the isolation of the correct proteins was verified by immunoblotting. The effect of overexpression of the proteins on host cells morphology (*E. coli* BL21) was tested using scanning microscopy. To investigate activity we have performed spectrophotometric and zymographic assay (using different purified bacterial cell walls).

Conclusions

Bioinformatic analysis showed the presence of a N-acetylmuramoyl-L-alanine domain in Lmo1521, N-acetylmuramidase domain in Lmo2591 and GW-motif in both. Our results indicate muralytic activity of the studied proteins against different bacterial cell walls. We have also determined the substrate specificity of these two autolysins.
THE EFFECT OF RSD, THE ANTI-SIGMA FACTOR OF SIGMA 70, ON BIOFILM FORMATION AND MOTILITY IN ESCHERICHIA COLI

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Background

In bacteria, σ subunit of RNA polymerase (RNAP) directs transcription initiation. The regulation of σ activity is important for fine tuning of gene expression. σ activity is determined by their cellular level, affinity for core RNAP, and interactions with regulatory proteins. In Escherichia coli, housekeeping σ factor, σ⁺⁷₀, has the highest affinity for core RNAP and is the most abundant s factor. Rsd, regulator of sigma D, binds specifically to σ⁺⁷₀ and it has been known as an anti-σ factor of σ⁺⁷₀.

Objectives

Even though Rsd is known as an anti-σ factor, no specific phenotype has been associated with deficiency or overexpression of Rsd to date. In this study, we found new phenotypes of the rsd mutant.

Methods

Cell-to-cell autoaggregation assay
Biofilm formation assay
Semi-solid agar assay for assessment of motility

Conclusions

An rsd-deficient mutant cell sank much faster than wild type. In spite of its increased cell aggregation, biofilm formation decreased in the rsd mutant. We found a protein of ~43 kDa whose expression was significantly higher in the rsd mutant compared to wild-type. Peptide mass fingerprinting revealed that this enriched protein is antigen 43 (Ag43) encoded by agn43, whose transcription is σ⁺⁷₀-dependent. Also, one protein band increased in wild-type and Rsd-overexpressing strains compared to the rsd mutant. Mass spectrometry identified this protein band as the flagellin protein FliC. Based on these results, we propose that Rsd decreases the transcriptional level of agn43 through the regulation of σ⁺⁷₀ activity and consequently it influences the biofilm formation and motility.
Background

Horizontal Gene Transfer (HGT) is an evolutive mechanism by which entire genes are transferred among bacterial cells, thus enabling an immediate adaptation to new environmental conditions. HGT has obvious implications for human health as it is used by pathogenic microorganisms – even among different species– during the spread of virulence factor and antibiotic resistance. Three main routes can be distinguished: conjugation (plasmid transfer through the direct interaction of two bacterial cells), transduction (bacteriophage-mediated DNA transfer) and transformation (uptake of naked DNA from the environment). The term DNA Uptake Pump (DUP) refers to the translocating machinery used by bacteria to incorporate naked DNA from the environment to the cytoplasm during bacterial transformation.

Objectives

The specific aim of this project is to determine the structure of the DNA uptake pump responsible for bacterial transformation in gram-positive bacteria using X-ray crystallography. This machinery is mainly formed by a DNA receptor, a transmembrane channel that mediates DNA translocation across the cytoplasmic membrane and an ATPase which pulls DNA into the cytoplasm at the expense of ATP.

Methods

We adopted a high-throughput approach to achieve our goals. We designed 96 constructs from five different species with several solubility-enhancing fusion partners and affinity tags. The constructs can be divided into three groups: putative soluble constructs, fragments containing trans-membrane helices and full-length integral membrane proteins.

Conclusions

We already have cloned the target genes and have tested them for soluble expression in E. coli. Large scale purification and preliminary crystallization trials are in progress for the soluble constructs.
Background

Why bacteria have evolved and maintained their specific shapes is one central question in bacterial cell biology. Rod-shaped bacteria cells are remarkable in keeping their geometry and shape. How this shape is regulated is still mostly uncharted territory.

Objectives

We are setting out to understand how and if rod-shaped bacteria are maintaining their rod shape when subjected to mechanical deformation. We explore how mechanical force changes the bacteria morphology and the consequences for bacteria after the mechanical force is released. The behaviour of proteins, that are involved in maintaining rod-shape, e.g. MreB, Mbl, is analyzed microscopically to understand if mechanical deformation can influence protein localization.

Methods

With the help of soft lithography we are applying microfabrication tools to create microenvironments to manipulate the shape of bacteria. To analyse the shape morphology and protein localization we are using different types of microscopy and automated image analysis tools.

Conclusions

Rod-shaped bacteria are maintaining their rod shape in response to mechanical deformations. Cells return to their original shape with a fast rate in cell wall recovery. It appears that the rod shape is a strong evolutionary feature and tightly controlled.
Background

Capnocytophaga canimorsus (Cc) are Gram-negative bacteria that are part of the normal flora of dogs' mouth and can cause rare but severe infections in humans which have been in contact with a dog.

Cc grow in co-culture with eukariotic cells by harvesting surface glycans via the PUL5 encoded Gpd complex. PUL5 mutant bacteria show a reduced growth and display coccoid forms, which strongly resemble E. coli mutant bacteria that are unable to synthesize aminosugars, and are deficient in peptidoglycan synthesis.

We thus speculated that Cc’s arrest of growth and cell rounding would be due to starvation of aminosugars.

Objectives

Here we address the question why Cc rely on host aminosugars and how they retrieve them from their niche, the dog’s mouth.

Methods

Cc strain 5 (Cc5) genome analysis revealed that Cc5 could not synthesize N-acetylg glucosamine (GlcNAc) because of the lack of two enzymes which convert glucosamine into GlcNAc. Heterologous expression of these enzymes as well as supplementation with GlcNAc completely restored the growth and abolished the formation of coccoid forms.

Conclusions

We show that C. canimorsus cannot synthesize GlcNAc and thus rely on exogenous GlcNAc which they retrieve by foraging glycans from mucin and N-linked
glycoproteins through two different apparatuses, Muc and Gpd, both related to the \textit{Bacteroides} starch utilization system (Sus).

We propose that the GlcNAc auxotrophy reflects the adaptation of \textit{C. canimorsus} to its host and ecological niche, the dog's mouth.
STREPTOMYCES PEUCETIUS VAR. CAESIUS AND ITS GLUCOKINASES: WHAT THEIR PHYSIOLOGICAL RELEVANCE IS?

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Background

*Streptomyces peucetius* var. *caesius*, a relevant industrial strain overproduces the antitumor, doxorubicin. As reported for other secondary metabolites, doxorubicin production is also repressed by glucose. In streptomycetes, the glucokinase (ATPGlk), participates in the process of carbon catabolite repression (CCR). However, the mechanism by which this enzyme exerts its regulatory role, has not been elucidated. In this bacterium, we have reported the presence of an additional glucokinase (PPGlk), which uses polyphosphate as its phosphate donor group. Contrary to what is observed for other *Streptomyces* such as *S. peucetius*, *S. coelicolor*, *S. lividans* and *S. thermocarboxydus*, in *S. peucetius* var. *caesius*, PPGlk activity is higher compared to that of ATPGlk. A similar situation has been reported for a chlortetracycline overproducer strain of *Streptomyces aureofaciens*. So far, the implications of the presence of two glucokinases in development and physiology of the genus *Streptomyces* are unknown.

Objectives

To evaluate the role of both glucokinases in the physiology of *S. peucetius* var. *caesius*, their participation in the CCR, morphological development and synthesis of anthracyclines.

Methods

ATPGlk::aac(3)IV and PPGlk::aadA mutants were obtained by replacing the genes for an apramycin and a streptomycin resistance cassette (*aadA*), respectively. Strains were grown in different carbon sources, and their ATPGlk, PPGlk, b-galactosidase and glycerol kinase activities were determined. Total anthracyclines and morphological differentiation (microscopy) were also evaluated.

Conclusions

The ATPgk::aac(3)IV mutant showed inability to grow in high glucose concentrations and exhibited deregulation in CCR. In regard to the parent strain, the mutant growth and production of anthracyclines, was delayed on solid medium.
Background

The genus *Streptomyces* comprises soil-dwelling, filamentous bacteria having great biotechnological significance since these bacteria produce 75% of the known antibiotics. Autoregulatory molecules play a key role in controlling antibiotic production and morphological differentiation. In *Streptomyces griseus* A-factor (2-isocapryloyl-3R-hydroxymethyl-y-butyrolactone) proved to be significant as the A-factor negative (AFN) mutant had a non-differentiating phenotype.

Objectives

Previous studies showed that the A-factor biosynthetic *afsA* gene was functional and transcribed in the AFN strain. Our aim is to understand the background of the AFN phenotype.

Methods

In this study the production of AfsA protein, aerial mycelium, spores, extracellular protease and antibiotics was followed on rich and minimal medium. To detect the presence of AfsA by Western blotting we produced the AfsA as a GST-fusion protein in *Escherichia coli*. The purified AfsA was used for immunization of rabbits.

Conclusions

The AFN strain did not form aerial mycelium on rich medium due to the lack of A-factor but spores were formed from the substrate mycelium. Aerial mycelium emergence and sporulation were observed on minimal medium in AFN although the production of spores, extracellular protease and antibiotics were decreased compared to the control strain. The AfsA protein was detected in the AFN strain. These data confirm the hypothesis that the presence of AfsA is not enough for normal A-factor production and A-factor is essential for the aerial mycelium but not for spore formation on rich medium. Moreover our data support the existence of a crosstalk between the A-factor mediated and the starvation induced pathway.
Background

The biogenesis of bacterial cell-wall polymers (e.g., peptidoglycan, lipopolysaccharides, teichoic acids) requires undecaprenyl-phosphate (C55-P). The glycan units are linked to this lipid at the inner face of the membrane. Thereafter, the membrane intermediate is translocated in order to transfer the glycan unit to the periplasmic side, where the glycan moiety is transferred to the growing polymer. The process releases undecaprenyl-pyrophosphate (C55-PP). C55-P originates from the dephosphorylation of C55-PP, itself generated by de novo synthesis and recycling.

Objectives

Identification and biochemical characterisation of bacterial C55-P phosphatase.

Methods


Conclusions

Two families of membrane proteins exhibit C55-PP phosphatase activity: BacA and members of the PAP2 super-family. *Escherichia coli* possesses one BacA and three PAP2 proteins (PgpB, YbjG and LpxT), raising the question of the significance of such a multiplicity. LpxT catalyses the transfer of the C55-PP phosphate group onto the lipid A moiety of the lipopolysaccharides (LPS), yielding C55-P and lipid A-1diP. The role of this LPS modification is under investigation (e.g., antibiotic resistance, ion homeostasis, innate immunity). This raises a tantalising hypothesis that the other C55-PP phosphatases could also exhibit such a phospho-transferase activity with different acceptor molecules. Our research project is aimed at the complete functional and structural characterisation of these multiple membrane phosphatases.
Background
Bacterial cell division is carried out at mid-cell position by a group of cell division proteins referred to as the divisome. For the Gram-negative model bacterium *Escherichia coli*, ten of these proteins (FtsZ, -A, -K, -B, -L, -N, -Q, -I, -W and ZipA) are considered as the core players of division. Thus far, interaction networks between individual divisome proteins have been investigated extensively with use of various biochemical and genetic techniques and many interactions between cell division proteins have been found. Still, no oligomeric interactions have been demonstrated that show that the divisome is present as a large multi-protein machinery.

Objectives
To detect whether or not a complete ‘divisome’ assembly is present in the cell.

Methods
Clear Native gel Electrophoresis on mildly solubilized *E. coli* cells, followed by 2D Electrophoresis and Western Blotting.

Conclusions
We describe the finding of a large 1 MDa cell division protein complex in *E. coli*, that includes at least 8 essential division proteins; FtsZ, ZipA, FtsK, FtsQ, FtsB, FtsL, ans FtsN. The complex is present in cultures of rapidly dividing cells, but not when cells are not dividing. Also, we found that a slight overexpression of *ftsQ D237N* (encoding interaction-impaired FtsQ), prevents formation of this 1 MDa complex. Combined, our findings indicate that a large protein complex containing cell division proteins indeed exists.
GANGLUTAMYLTRANSPEPTIDASE EXPRESSION IN CAMPYLOBACTER JEJUNI IS NOT ONLY REGULATED BY THE RACRS TWO-COMPONENT SYSTEM

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Background

Campylobacter jejuni is an asaccharolytic human pathogenic Gram-negative bacterium that utilizes amino acids as carbon and energy source and colonizes the gastrointestinal tracts of warm-blooded animals. Some strains of C. jejuni possesses the enzyme gamma-glutamyltranspeptidase (GGT), which enables growth on glutathione and glutamine by generation of glutamate in the periplasm and contributes to persistent colonization of the avian gut.

Objectives

We have shown that the RacRS two-component system is involved in cytoplasmic glutamate anabolism by regulating the gltBD genes, here we investigated whether the RacRS system also regulates the periplasmic glutamate anabolism by regulating the ggt gene.

Methods

Our results show that RacRS regulates the expression of ggt under low oxygen conditions. Under these conditions, higher transcription levels of the ggt gene and enhanced GGT activity are observed in the wt compared to the racR mutant strain. By using different ggt truncated promoter elements in EMSA and luciferase reporter assays we show that RacR binds directly on a specific location of the ggt promoter. Furthermore we show that under high oxygen conditions ggt expression peaks around end-log phase and GGT activity is highest in stationary phase and is not dependent on RacR. GGT expression and activity is repressed by addition of glutamine to the medium or glutamine catabolic products.

Conclusions

In conclusion we show that the C. jejuni GGT activity is dependent on multiple factors and one of them is the RacRS two-component system.
EFFECT OF HUMAN ANTIMICROBIAL PEPTIDES AGAINST LEGIONELLA PNEUMOPHILA

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Background
Antimicrobial peptides (AMPs) are natural antibiotics widespread throughout the animal kingdom, from bacteria to mammals. They are important components of both innate and adaptative immunity, providing protection against a broad spectrum of pathogens, such as viruses, bacteria, fungi and parasites.

Objectives
Investigate the effect of two human AMPs (HBD-3 and LL-37) on Legionella pneumophila.

Methods
In this study, we investigated the action of synthetic LL-37 and HBD-3 on both extracellular and intracellular lifestyle of L. pneumophila.

Conclusions
We showed that both peptides exhibit a phase dependent bactericidal effect on extracellular L. pneumophila. We also observed that LL-37 and HBD3 inhibit intracellular replication of L. pneumophila in macrophages and pneumocytes. We showed by colony counting assays that the adherence and internalization of L. pneumophila was not affected in presence of LL-37, but was stimulated with HBD-3, suggesting that this two peptides restrain the bacterial replication by different mechanism of action.

Human AMPs LL-37 and HBD-3 seems to be involved in innate immunity against L. pneumophila by acting on the extracellular bacteria and on its intracellular replication.
SINGLE-MOLECULE IMAGING REVELS HETEROGENEOUS MOBILITY OF MEMBRANE PROTEINS IN LIVE ESCHERICHIA COLI
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Background
Membrane proteins perform vital cellular functions like respiration, signaling and nutrient uptake. For proper functioning, conformational dynamics, complex formation and ability to diffuse in the membrane are vital parameters. Despite a lot of work on model membranes, little is known about lateral mobility of proteins in bacteria.

Objectives
To investigate how molecular crowding, protein size and membrane fluidity affect the mobility of membrane proteins in living E. coli bacteria
Methods

Here we use single-molecule wide-field epi-fluorescence microscopy to track the
lateral mobility of seven integral membrane proteins of different size fused to green fluorescent protein in living E. coli. We apply a novel method, IPODD to extract accurate diffusion coefficients from the 2-D projected diffusion trajectories along the 3-D curved bacterial membrane.

**Conclusions**
The diffusion coefficients we find are significantly lower than those reported in *in vitro* studies of isolated membrane proteins in GUVs. Our results indicate that molecular crowding in the plasma membrane of E. coli substantially slows down transmembrane protein diffusion. Surprisingly, all seven proteins studied diffuse heterogeneously: they all show a faster and a slower moving component. Our interpretation of the heterogeneity is that it reflects heterogeneity in membrane composition. To further support this interpretation, we stained E. coli inner membrane with a lipid-mimicking dye Dil-C12 and tracked its lateral mobility. Strikingly, Dil-C12 diffuses heterogeneously yet again reflecting the dynamic heterogeneous nature of the plasma membrane. Therefore, from these observations, we propose that the plasma membrane of E. coli might contain micro-domains with different fluidity that could play key roles in specific localization of proteins.
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Bacterial spores

WHEN MICROBIAL SURVIVAL STRATEGIES NEVER STOP TO WONDER: SPORULATION OUTSIDE FIRMICUTES

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Background

Spore or spore-like structures are only found in four bacterial phyla: Actinobacteria, Cyanobacteria, Proteobacteria and Firmicutes. These structures provide resistance to adverse conditions. The ability to form spores is not, however, a widely spread characteristic and it is restricted to only some orders within those phyla. For example amongst Proteobacteria, solely δ-proteobacteria can produce spore-like fruiting bodies, or so we knew. A novel γ-proteobacterium, Serratia ureilytica str. Lr5/4, was found to produce spores that not only resemble structurally to those produced by endospore-forming Firmicutes, but also provide heat-resistance.

Objectives

The aim of this study is to describe this novel strain and its Firmicute-like spores and to reveal the molecular pathway of this sporulation procedure in comparison to those that are already known.

Methods

Physiological, biochemical, carbon source assimilation and antibiotic resistance tests were performed. Morphology of vegetative cells and spores was described by phase contrast microscopy, SEM, and TEM. Moreover, spores of Lr5/4 were revived after heat-shock tests and shown to contain dipicolinic acid (DPA). These two characteristics were so far unique to the heat-resistant endospores found in Firmicutes. Sequencing and annotation of its full genome has been performed in order to reveal the relationship of spore formation in Lr5/4 to other known sporulation pathways.

Conclusions

It has been previously proposed that the properties of spore formation in non-sporulating species were due to molecular gene transfer. However, the present study rejects this scenario and demonstrates a novel mechanism for the formation of the described spores of S. ureilytica.
Background
Biofilm proficiency has a great impact in bacteria-host interactions. Caenorhabditis elegans and the spore-forming bacteria Bacillus subtilis are both common soil inhabitants which are used as models of lifespan studies. However, the role of in vivo biofilm development during gut colonization and its effects on animal lifespan are poorly documented.

Objectives
Here, we investigated the ability of B. subtilis to form a biofilm in the intestine of C. elegans and its effects on gut colonization and lifespan of the nematode.

Methods
The results showed that wild-type NCIB3610 strain was able to colonize and persist in the gut of C. elegans more efficiently than the laboratory strain JH642. The ability to make a biofilm was essential for the observed behavior because FICT-labelled NCIB3610-derived biofilm-mutant cells, but not FICT-wild-type cells, lost the capacity to persist and colonize the nematode gut. In situ fluorescence microscopy and beta-galactosidase expression, driven from the promoter of surfactin harbored by B. subtilis engineered cells and used to feed the worm confirm the higher ability of NCIB3610, instead of JH642 cells, to colonize the gut of C. elegans. The ability to make a persistant biofilm correlated well with a positive lifespan effect of the bacterium on the nematode.

Conclusions
B. subtilis spores are able to germinate, growth, make biofilm and persist in the intestine of C. elegans. In addition, we provide evidences showing that the intestinal life cycle of B. subtilis making a biofilm is beneficial and required for the increase for the lifespan of C. elegans.
CHARACTERIZATION OF SPORE-FORMING BACILLUS, ISOLATED FROM THE ANDES MOUNTAINS, EXHIBITING HIGH LEVELS OF RESISTANCE TO ARSENIC AND UV RADIATION

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Background

The North-Western part of Argentina is particularly rich in wetlands located in the Puna at an altitude between 3,600 and 4,600 m above sea level. Here, incidence of high levels of UV radiation and contamination with high contents of toxic metals, particularly arsenic (As) is common. We hypothesize that one strategy to contribute to the remediation of these area could be the use of Bacillus spp.

Objectives

We collected soil and water samples from different parts of the Puna and isolated native spore-forming Bacilli. The novel isolates were examined at different growth temperatures and osmolarities on their ability to grow and make biofilms in the presence of toxic concentrations of As and high UV radiation.

Methods

The Bacilli communities were characterized by pyrosequencing 16S rRNA gene amplicons derived from specific PCR oligonucleotides. The Andean Bacillus isolates displayed a greater capacity to make persistent biofilms, colonize surfaces (swarming motility) and tolerance to high temperatures, salinity and UV-B / UV-C radiations at levels that were lethal to common laboratory strains.

Conclusions

The present results allow the potential biotechnological application of the isolated Bacillus strains with bioremediation potential of pristine environments contaminated with metals.
NOVEL GENES INFLUENCING THE GERMINATION RATE OF SPORES OF BACILLUS SUBTILIS FOOD ISOLATES

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Background

Bacillus subtilis responds to starvation with formation of endospores. Spores can easily contaminate food materials and their resistance impedes their removal from food products. In response to various triggers, spores reinitiate vegetative growth via germination. Vegetative cells can subsequently cause food spoilage. Accurate prediction of germination behavior is difficult as differences in germination responses are observed even among various strains of one species. Thus, data obtained for the model strain, B. subtilis 168, cannot be directly applied to food-spoiling strains.

Objectives

The purpose of this study was to elucidate germination behavior of spores of B. subtilis strains that cause food spoilage and to couple the observed germination phenotypes with the strains’ genomic contents.

Methods

Genomes of thirteen B. subtilis industrial isolates were sequenced. Germination of their spores was induced under various conditions and analyzed via absorbance measurements and phase-contrast microscopy. The spore germination responses of different strains were linked with their genomic content using gene-trait matching software.

Conclusions

Strains were divided into two phenotypic groups based on the ability of their spores to germinate, in particular their germination rates. Slower germination correlated with the presence of specific genes in industrial isolates, which are absent in the laboratory strain B. subtilis 168. Insertion of these genes into B. subtilis 168 decreased the rate of germination of its spores. Thus, new genes influencing B. subtilis germination responses were identified. The function of these genes is currently being elucidated. These genes can be used in the future as biomarkers indicating slow germination.
UNDERSTANDING BACTERIAL SPORE RESISTANCE TOWARDS PLASMA STERILIZATION

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Background

Background: Spores of Bacillus subtilis have been used extensively as biological indicators for industrial purposes such as sterilization or decontamination. For several years a number of activities in the field of sterilization of heat sensitive materials by means of non-thermal plasmas have been known. Plasma sterilization methods are characterized by the use of gas or gas mixtures that are partially excited by an applied electric field. Plasma discharges contain a high degree of UV/VUV-radiation, as well as charged particles and free radicals, which exert detrimental effects on microorganisms by damaging genetic material, outer cell layers and proteins.

Objectives

Objectives: We are interested in understanding the protective attributes and molecular mechanisms involved in the B. subtilis spore resistance to plasma sterilization.

Methods

Methods: In a systematic manner different B. subtilis spores varying in their protection (e.g., coat, crust, DPA, SASP formation) and DNA repair capabilities were studied on their impact towards plasma sterilization.

Conclusions

Conclusions: We will present our recent findings on the protective attributes and molecular mechanisms involved in the spore resistance to plasma sterilization. The multi-layered proteinaceous spore coat, being the first barrier to environmental influences, was shown to be a major factor contributing to spore resistance towards plasma treatment. During spore revitalization, DNA damage accumulated during the dormant spore stage is the subject of a variety of different repair systems. Deficiencies in repair genes of non-homologous end-joining, spore photoproduct
lyase, or nucleotide excision repair led to a significant decrease in spore resistance to low pressure plasma sterilization.
Background

*Bacillus weihenstephanensis* is a spore forming food pathogen known for its ability to sporulate, germinate, grow and produce toxins at lower temperatures than its close relatives. As such, it presents an increased risk to food safety where standard procedures rely on cold-inactivation of microbes. Properties of the infective form Bacilli, the spore, are known to vary with the conditions under which the spores were produced. For instance, lower sporulation temperature has been indicated to result in a lower wet heat resistance. This resistance is in part attributed to the spore coat layers, proteinaceous layers surrounding the core which contains the bacterial DNA.

Objectives

The aim of this study was to determine the proteomic basis for the heat resistant phenotype. As no specific singular protein can be indicated to confer heat resistance, this supposedly acquired trait can only be appreciated by analysis of the spore proteome as a whole.

Methods

Analysis of the proteome of spores of *B. weihenstephanensis* strain WSBC10204 produced at either 12°C or 30°C by mass spectrometry provided insight into variations in the protein content of the spore coat at different sporulation temperatures.

Conclusions

Interestingly, only minor differences were observed, suggesting wet heat resistance does not depend on the identity of the proteins making up the coat layers. As wet heat resistance was indeed lower for the spores produced at lower temperatures, low temperature might instead affect the rate of spore maturation instead, a process where spore coat proteins are cross-linked after they have been deposited onto the spore.
FEMS-1185  
Bacterial spores  

CARBON FIXATION AND ALLOCATION IN A FILAMENTOUS CYANOBACTERIUM DURING AKINETE DIFFERENTIATION USING NANOSIMS AND MICROSCOPY  
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Background  
Akinetes are resting cells of species members of the Nostocales and Stigonematales orders of cyanobacteria. These are spore-like, non-motile cells that differentiate from vegetative cells and serve a perennating role. During their differentiation, akinetes vary their metabolic activities and cellular pools.  

Objectives  
Here we follow carbon assimilation in the cyanobacterium Aphanizomenon ovalisporum and accumulation of cyanophycin during akinetes differentiation and maturation.  

Methods  
Photosynthetic assimilation of H₁³CO₃⁻ was studied by NanoSIMS technology and variations in abundance and distribution of cyanophicin globules were followed by transmission electron microscopy and histochemistry techniques.  

Conclusions  
NanoSIMS results clearly indicate that during their differentiation, akinetes maintain metabolic activity and assimilate inorganic carbon via photosynthesis. Towards advanced stages of differentiation and maturation, the metabolic activity of akinetes reduced and many of the free akinetes lost their photosynthetic capacity as indicated by changes in ¹³C/¹²C ratio. Nevertheless, a small group of free (mature akinetes) still maintains their photosynthetic capacity. Structural changes during akinetes differentiation observed by transmission electron microscopy and histochemistry techniques, included the accumulation of cyanophycin bodies in akinetes. The rate of cyanophycin accumulation and carbon allocation into this protein storage is currently being investigated in a complementary NanoSIMS analysis.
Background

In response to nutrient limitation Gram positive organisms like *Bacillus subtilis* form dormant spores. These cellular entities are survival capsules, resistant to chemical and environmental assaults. They pose challenges to the food and medical sectors. Upon contact with germinants spores return to vegetative life through a process called 'germination and outgrowth'. The vegetative cells may cause food spoilage and food borne diseases.

Objectives

To perform a detailed quantitative time-resolved proteomics study of (dormant)spores and their secretome during germination and outgrowth

Methods

Using a quantitative proteomics approach a study of the time resolved break down of spore proteins during early stages of germination was initiated. Spores were germinated in minimal medium supplemented with a germinant mixture (L-Asparagine, D-glucose D-fructose and Potassium Chloride). The spore layers and the secretome were analyzed separately by Ion Trap LC-MS/MS analysis.

Conclusions

During germination the action of proteases, lytic enzymes and peptidases in the spore results in proteolysis, cortex hydrolysis and break down of spore dormancy. Partially digested proteins as well as intact proteins are released from the spore into the medium. Study of these proteins by our method allowed us to understand molecular details of the germination process. Current analyses are aimed at correlating the germinating spore proteome to its transcriptome under control as well as thermal and organic acid stress conditions (Keijser et al. 2007 and Ter Beek et al., our unpublished observations). We aim at deconvoluting heterogeneous germination and outgrowth profiles using our live imaging technology (Pandey et al., 2013).
Background

Undomesticated spore formers isolated from the environment or from processing facilities commonly show higher diversity, resistance and more heterogeneous behaviour compared to widely used reference strains domesticated in laboratories and that may have lost relevant traits. Analysis of such undomesticated isolates therefore provides valuable insights in sporulation capacity, spore resistance, germination and outgrowth efficiency that determine survival and fitness in different environmental niches including food and food processing environments.

Objectives

In our project we aim to obtain insight in the diversity among pathogenic Bacillus cereus strains with focus on spore properties. To this end, B. cereus reference strains ATCC14579 and ATCC10987 were analysed and compared to behaviour of strains isolated from food and food production facilities.

Methods

Growth, sporulation and germination capacity of 20 B. cereus strains isolated from different food products and two reference strains was characterized under different conditions. Spore stress resistance and germination characteristics were assessed upon exposure to disinfectants, heat and mild processing technologies. Behaviour of untreated and treated spores was analysed both at population level as well as at the level of individual spores by using Flow cytometry (FCM) in combination with sorting (FACS).

Conclusions

The B. cereus strains displayed a highly diverse response with respect to growth, sporulation, spore resistance and germination capacity. Additionally, spore population behaviour showed substantial differences with some strains displaying a
homogenous and other a heterogeneous performance. We have demonstrated the relevance of including environmental isolates in the studying behaviour of pathogenic spore former *B. cereus* along the different steps of life cycle.
Background

*Bacillus licheniformis, B. amyloliquifaciens* and *B. thermoamylovorans* can cause significant non-sterility issues in high heat treated foods due to survival of heat resistant spores that can subsequently germinate followed by outgrowth during the shelf life.

Objectives

To gain improved control of spores of these species in food manufacturing, a genetic basis for spore heat resistance and germination efficiency was investigated by combining spore phenotypes and genome information of individual strains.

Methods

The heat inactivation kinetics of spores of different isolates were determined. In addition, the genomes of over 20 strains within these species were sequenced to investigate a genetic basis for different spore heat resistances.

For *B. licheniformis* (10 strains) and *B. amyloliquifaciens* (9 strains) significant differences in heat resistance of spores were found for different isolates. Strains producing heat resistant spores contained a subset of genes belonging to a cluster that confers high heat resistance in *B. subtilis*.

For spores of *B. thermoamylovorans* (4 strains) spore heat resistance, nutrient germination, non-nutrient germination, and detailed spore counts were determined. Non-nutrient germination using calcium-dipicolinic acid induced good germination of spores of all strains. Poor germination on regular rich cultivation media resulted in gross underestimation of the viable spores truly present.
Conclusions

The above findings shed light on different phenomena that can contribute to spore survival. Such information is important to devise appropriate strategies to enumerate spores in food ingredients and to inactivate spores, leading to improved control of heat resistant spores in foods.
SSGB COORDINATES SEPTUM SITE LOCALIZATION WITH CHROMOSOME ORGANIZATION DURING STREPTOMYCES SPORULATION

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Background
Streptomyces are multicellular bacteria with a complex life cycle. During the reproductive phase aerial hyphae transition into chains of hundred or more spores via a highly orchestrated cell division process. Coordination of sporulation is regulated by the SLP proteins (SsgA-like proteins). SsgA forms an array of foci along the hypha and provides an anchor for SsgB to multimerize on the cell membrane, after which SsgB recruits FtsZ and stimulates FtsZ polymer formation.

Objectives
SsgB is a highly conserved protein that forms trimers. The C-terminal α3 helix is important for trimerization and interacts with α1 and α2 of the neighboring SsgB monomers. To elucidate how SsgB interacts with SsgA and FtsZ, we have created an SsgB mutant library.

Methods

These ssgB variants were introduced in the ssgB mutant strain, after which strains were automatically scored for restored sporulation via a scanner based imaging approach. Over 500 variant SsgB a single amino acid change and these were further analysed.

Conclusions

All sporulating samples were examined with transmission electron microscopy (TEM), which revealed effects varying from changes in spore wall thickness, size distribution, DNA segregation to affecting spore shape. In contrast wild type spores are similar sized, have condensed DNA in the center of the spores and have a thick spore wall.

We have shown that single amino acid changes in SsgB can facilitate many division related phenotypes. and ultimately dictate septum orientation resulting in longitudinal division. The latest results in elucidating the function of individual amino acids will be presented.
EVALUATION OF TWO SCREENING METHODS FOR DETECTION OF BIOFILM FORMATION AND ITS EFFECT ON ANTIBIOTIC SUSCEPTIBILITY AMONG STAPHYLOCOCCI SPECIES

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Background

Biofilm formation is a hallmark characteristic of staphylococcal infections as it allows attachment to solid surfaces, exchange of genes that can result in more virulent strains, production of high concentration of toxins, evasion of immune defenses and antimicrobial resistance.

Objectives

This study was conducted to evaluate two different screening methods for detection of Biofilm formation and its effect on antibiotic susceptibility among the clinical isolates of different Staphylococci species.

Methods

Seventy five isolates of staphylococci were tested by modified tissue culture plate method (MTCP), congo red agar (CRA) method and tube method (TM) for their ability to form biofilm. Antibiotic susceptibility testing was done for thirty isolates, 15 biofilm-producers and 15 biofilm nonproducers, using broth microdilution assay.

Conclusions

The rate of detection of biofilm formation among the staphylococcal isolates by the MTCP method, the CRA method and the TM were 48%, 38.7% and 25.3% respectively. Comparing to the results of MTCP method as a gold standard, the sensitivities of the CRA method and tube method were 80.6% and 52.8% respectively. Both showed 100% specificity. Regarding the antibiotic susceptibility of the biofilm producers and non-producers, statistically significant differences were detected for oxacillin, ceftriaxone and augmentin susceptibility (P<0.02). All the
isolates were sensitive to vancomycin. The MTCP takes 3 days and many steps. The TM is observer dependent. Congo red agar method can be used as a screening method for detection of biofilm formation among staphylococcal isolates as it is cheap, rapid with no subjective errors and requires less expertise.
INFRARED SPECTROSCOPIC ANALYSIS OF XYLELLA FASTIDIOSA BIOFILMS

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Background
Xylella fastidiosa (Xf) affects plant xylem vessels, causing diseases such as Pierce’s disease. As abundant plants of economic relevance may be affected by Xf, it is essential understanding the biofilm formation behavior of Xf, whose genome has been sequenced partly by our collaborators for developing appropriate countermeasures.[1]

Objectives

The appearance of Xf infected olive trees in Italy is of substantial concern due to the first appearance of this species within Europe. Considering the role of biofilms within many human diseases, the investigation of the adsorption behavior of bacteria along with the involved biomolecules is evident.[2][3] Particularly interesting is the formation
of disulfide bonds appearing essential during biofilm formation.[4]

Methods
This study focuses on the analysis of biomolecules involved in Xf biofilm formation via vibrational spectroscopy. In a unique instrumental combination, infrared attenuated total reflection (IR-ATR) spectroscopy and atomic force microscopy (AFM) were applied to characterize the growth media and biofilms associated with Xf.

Conclusions
Experimental and analytical procedures will be discussed providing first insight on the involved molecular processes during biofilm formation and maturation.

SUPPLEMENTARY MICROBIOLOGICAL ANALYSIS OF FILLING MATERIALS REVEALS NEW INSIGHT INTO THE ECOLOGY OF PERSISTENT ENDODONTIC INFECTIONS

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Background

Persistent root canal infections correlate with endodontic treatment failures. Microorganisms have been found in 35%-100% of previously treated root canals. The analysis of the microbiota of endodontic infections is prerequisite for improvement of treatment methods of infected root canals.

Objectives

1. The microbiological analysis of infected root-filled teeth with periradicular lesions using culture-dependent and -independent approaches

2. The analysis of the microorganisms adhered to the retrieved endodontic filling materials (EFM) and the comparison with the results gained from the root canal samples alone.

Methods

Twenty patients were enrolled in this study and samples from the corresponding twenty root-filled teeth were taken according to standard protocols. In addition to these samples, the EFM were also analyzed microbiologically using culture and culture-independent methods and transmission electron microscopy. The culture-independent technique was conducted by cloning and sequencing of 16S rRNA amplicons.

Conclusions

Results: Bacteria were revealed in 15 root canal samples and in 18 EFM samples at a concentration ranging from $10^3$ to $10^7$ cfu/ml. 56 different species were found in association with the root canal filling materials. In contrary, only 47 species could be identified in the root canal samples. 20 species were detected solely in the EFM samples including Aggregatibacter actinomycetemcomitans and Eubacterium nodatum. The most abundant species belonged to the phyla Firmicutes, Actinobacteria and Proteobacteria.
**Conclusions:** The analysis of microorganisms associated with root-canal filling materials in addition to the root canal samples taken after revision reveals a better insight into endodontic infections and should not be neglected.
ANALYSIS OF INTERACTIONS BETWEEN NEISSERIA STRAINS IN BIOFILMS
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Background
The human nasopharynx is colonized by different species of the genus Neisseria, including commensals, such as N. lactamica, but also an important pathogen, i.e. N. meningitidis. In situ, these bacteria are organized in micro-colonies, structures that resemble biofilms and that may represent a defense mechanism for the bacteria to survive in a hostile environment. In vivo, these species must interact and compete for the same niche.

Objectives
Here, we designed novel constructs to generate green and red fluorescent Neisseria strains to facilitate the discrimination between different microorganisms by microscopy in several assays.

Methods
These constructs were stably integrated in a highly conserved region on the chromosome of both species, as evidenced PCR assays, and allowed for constitutive expression of the fluorescent proteins. We next optimized biofilm formation assays and studied the organization of biofilms of independent and mixed Neisseria strains.

Conclusions
Our results showed that indeed biofilms of monocultures of N. meningitidis and N. lactamica are organized as micro-colonies with intervenient spaces. However, the organization of these spaces and the distribution of the biomass varied between strains of the same species. Mixed biofilm assays revealed that both species can differently interact, altering the organization of the biofilms. This work is the first report of the interaction between both species and allows for speculation about the process of human colonization.
Changes in Sewer Biofilm Microbial Communities Related to Downstream Nitrate Dosage

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Background

Nitrate (NO\textsubscript{3}\textsuperscript{-}) is an effective and widely used chemical in sewer systems to reduce biogenic sulfide (H\textsubscript{2}S) and methane (CH\textsubscript{4}) produced in anaerobic rising main pipes. However, the main limitation of this approach is that anoxic conditions must be continuously kept through the whole pipe, otherwise H\textsubscript{2}S-producing biofilms prevail.

Objectives

The aim of this study is to determine the effects of the Downstream Nitrate Dosage strategy (DND) on anaerobic sewer biofilms with regards to sulfide oxidation and, for the first time, on methane production.

Methods

Effectiveness of the strategy was assessed on H\textsubscript{2}S and CH\textsubscript{4} abatement on the effluent of specially designed laboratory-scale systems that mimics a real sewer biofilms. A combination of process (normal functioning monitoring and batch tests) and molecular (by 454-pyrosequencing) methods were used to investigate the impacts and microbial activities related to the nitrate addition.

Conclusions

Results showed a complete abatement of H\textsubscript{2}S generated, due to the growth of a Sulfide-Oxidizing Nitrate-Reducing population. Methane was reduced to 50% while nitrate was added, due to the CH\textsubscript{4} oxidation in the anoxic conditions established at the end of the pipe. An increase of microorganisms of the genera \textit{Simplicispira}, \textit{Comamonas}, \textit{Azonexus} and \textit{Thauera} was detected during nitrate addition. Regarding anoxic methane oxidation, only one Operational Taxonomic Unit (OTU) was identified, which is likely related with this metabolism. Both sulfidogenic and methanogenic activities resumed upon cessation of NO\textsubscript{3}\textsuperscript{-} dosage. Obtained results are relevant for the optimal management of nitrate dosage strategies in sewer systems.
Background

Transport of wastewater along sewer pipes facilitates the formation of microbial biofilms that grow attached to their inner surfaces. Anoxic conditions in sewers favour the production of both sulfide (H₂S) and methane (CH₄) as end products of anaerobic microbial metabolisms, i.e. sulfate-reduction and methanogenesis, respectively. The build-up of H₂S and CH₄ in sewerage causes different detrimental effects such as odour, corrosion and toxicity.

Objectives

The aim of this study was to investigate the initial stages of development of microbial biofilms in sewer systems with a special focus on the interactions between sulfate-reducing bacteria (SRB) and methanogenic archaea (MA).

Methods

The work was carried out using a laboratory sewer pilot plant fed with wastewater that mimic the functioning of a real anaerobic pressured sewer. Biological activities and phylogenetic community composition were investigated during sewer colonisation using a combination of molecular techniques (DGGE, qPCR and 16S rRNA tag sequencing) and process data (H₂S and CH₄ production).

Conclusions

Results showed that the SRB community was established and active after two weeks of biofilm development, causing substantial sulfide accumulation in the pilot plant. In turn, growth of MA and methane production were low during initial stages of biofilm colonisation but readily increased over time after the replacement of initial MA species derived from human faeces (Methanobrevibacter spp.) by other MA representatives more adapted to sewer conditions (Methanoseta spp.). Altogether, our results pointed to different dynamics and activity of key community members from sewer biofilms, greatly affecting the production kinetics of H₂S and CH₄ along time.
Background
To better understand the molecular events that occur during *Vibrio cholerae* biofilm formation in vivo, we did a previous proteomic analysis of planktonic and biofilm cells, after 14, 24 and 48h of growth under conditions that mimic the intraintestinal environment, namely, pH 7.4 at 37°C, Pi limitation and the presence of the bile salt, sodium deoxycholate (DOC). The groups of differential proteins identified indicated that free cells were at stationary phase, whereas, those in biofilms were at exponential culture phase, after 48h growth. Moreover, in planktonic cells at all times, alkaline phosphatase was produced, suggesting Pho regulon activity, which was not observed in the biofilms cells.

Objectives
To perform a lipidomic analysis to explore differences in lipid profiles between planktonic and biofilm cells of *V. cholerae*.

Methods
Cells were grown overnight in MGLP (MOPS, salts, glucose, pH 7.4 with KH$_2$PO$_4$ at 0.65μM-low phosphate) without (free cells) or with 0.2% DOC (biofilm) for 14-48hs, at 37°C, without agitation. The lipids were extracted according to Bligh and Dyer 1959 and analyzed by Q-TOF LC/MS.

Conclusions
In the 14h planktonic cells, the great majority of phospholipids was replaced by ornithine containing lipids (OL). Between 24 and 48h, OLs were found in these cells, but in smaller amounts. In biofilm cells, phosphatidylethanolamine (PE) was the major lipid at all times, whereas OLs were detected in small amounts. These data suggest, for the first time, the involvement of PhoB regulated genes in the expression regulation of OLs and PE in *V. cholerae*. 
Background
Bacteria have the ability to attach to surfaces commonly found in the food processing environment, such as polystyrene and stainless steel and to become more resistant to environmental stresses and sanitizers. So the search for new fight strategies is needed.

Objectives
In this study, a strain of Enterococcus (En.) faecium, originally isolated from a milking machine inner-surface, was studied for its biofilm formation potential on plastic and stainless steel supports and its antibacterial and anti-adhesive potential against four pathogens (Escherichia coli ATCC 25922, Staphylococcus aureus ATCC 25923, Pseudomonas aeruginosa ATCC 27853 and Listeria innocua CLIP 74915).

Methods
The adhesion and the anti-adhesive potential of this strain were studied (at 25 and 37°C) using polystyrene microtiter plates and stainless steel 304 L coupons after their surface characterization by contact angle method.

Conclusions
The results revealed that the strain was a strong biofilm producer on polystyrene and stainless steel either at 25 and 37°C with the best adherence level observed at 25°C. En. faecium cells and culture supernatant showed a significant (p<0.05) inhibition potential of the pathogens most probably due to bacteriocins synthesis.
Background

Archaeal biofilms are interesting for biotechnological applications like whole cell biocatalysis, because they promise a higher tolerance towards toxic compounds, which is a characteristic property known for biofilms. In this study the effect of the organic solvent 1-butanol on biofilms of the thermoacidophilic, aerobic crenarchaeon *Sulfolobus acidocaldarius* with growth at 78 °C and pH 3 was investigated. *S. acidocaldarius* is adapted to extreme environments, genetically tractable and able to form biofilms.

Objectives

The aim of this study was to characterize the tolerance of *S. acidocaldarius* biofilms to 1-butanol exposure. The effect to 1-butanol on biofilm formation as well as on established biofilms was investigated.

Methods

Submerged biofilms were grown (4 d, 78 °C) in 96-well microtiter plates, Petri dishes or µ-dishes on either polystyrene or glass surfaces. 1-butanol was present during biofilm formation or added to already established biofilms. Analysis of biofilms was performed by microscopic methods (confocal laser scanning microscopy, epifluorescence microscopy, atomic force microscopy, environmental scanning electron microscopy), and microbiological and biochemical methods (determination of total cell counts, viable counts and aerobic respiration activity). Extracellular polymeric substances were isolated and quantified by colorimetric assays.

Conclusions

*S. acidocaldarius* tolerates 1-butanol concentrations up to 1.5 % [v/v] as validated by viability assays. 1-butanol exposure led to stress responses like aggregation of cells, a change of carbohydrate distribution and composition as well as carbohydrate and
protein amounts in the biofilm matrix. Butanol-exposed biofilm cells revealed alterations of shape and loss of cell appendages.
EFFECT OF THAI-PLANT EXTRACTS AND COMBINATION WITH ANTIMICROBIALS ON PSEUDOMONAS AERUGINOSA BIOFILM FORMATION

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Background
Pseudomonas aeruginosa is the most common Gram-negative bacteria causing fatal nosocomial infections in hospitalized patients especially with the immunocompromised subjects. Bacteria can resist a wide variety of currently available antibiotics. One of resistant mechanisms is by forming the biofilm in infected host preventing drugs to access to the respective targets of the bacterial cell. Novel strategy to overcome this antibiotic resistant mechanism is to use the combination of antibiotics with inhibitors of biofilm formation.

Objectives
The objective in this study is to investigate the effect of mulberry-leaf, fruit and goji berry on biofilm formation from P. aeruginosa.

Methods
The result was shown by specific biofilm formation (SBF) index and the ethanolic extract (256 mg/ml) and the ether extract (32 mg/ml) of mulberry-leaf presented strong activity (SBF index > 1.1) to reduce biofilm in 44.0 % and 44.8 %, respectively. The plant extract with anti-biofilm activity was tested further in combination with gentamicin (0.125 to 1024 µg/dl) by Checkerboard method indicated that there was no synergistic efficacy. The results of MBEC and CLSM revealed that the ethanolic extract (32 mg/ml) was effective on P. aeruginosa biofilm.

Conclusions
The extract of mulberry-leaf could be an alternative thai herbs for biofilm formation inhibiting the most common drug resistant bacteria.
BACKGROUND

In natural and man-made ecosystems, anammox bacteria demonstrate a strong tendency for attached growth and biofilms formation.

OBJECTIVES

We studied biofilms structure and formation by microbial community with dominant group of anammox bacteria, enriched in anaerobic vertical upflow bioreactor with nitrogen load 5 g N/l per day and upflow vertical concentration gradient of nitrogen substrates and pH.

METHODS

Biofilms structure and microbial community composition were studied via microscopy (light, electron, confocal and atomic force) and molecular genetics methods (sequencing and FISH). Biofilm formation de novo was studied on microscopic slides submerged in the upper part of the bioreactor.

CONCLUSIONS

Several groups of microorganisms are involved in biofilm formation. Anammox bacteria belong to three different species. Their satellites are: trichal forms (presumably Chloroflexi), coccoid and rod-shaped cells (likely different species of nitrifier and denitrifier communities involved in nitrogen removal and trace amounts of oxygen consumption). Chloroflexi community includes 3 phylotypes of Anaerolineae and Dehalococcoidia strains responsible for biofilm spatial structure formation and trace amounts of organics elimination.

Primary biofilms are discovered on day 10-12. Subsequent growth of biofilms appears due to increasing the number of cells involved in biofilms and extracellular polymeric matrix accumulation. Mature biofilms (80-90 days) include cells of the same morphotypes as mature biofilms inside the reactor: clustered cocci with typical
anammox ultrastructure, interstitial rods and remarkably long trichal cells (more than 100 µm long).
Surface microrelief images of such biofilms were obtained via atomic force microscopy.
ASSSESSMENT OF BIOFILM DESTABILIZATION AND REMOVAL BY LAUNDRY AGENTS

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Background

The formation of biofilms is a well known phenomenon in a variety of technical and natural environments such as rivers, water systems or washing machines. They can harbor a potential risk to human health serving as habitat for pathogens. Furthermore, biofilm growth within washing machines leads to biocorrosion and unpleasant odours. Therefore, the removal or prevention of biofilm growth is of great interest.

The relevance of biofilm EPS in terms of bacterial adhesion to surfaces has been indicated in several studies. Therefore, EPS can be one working point for laundry agents for the destabilization and detachment of biofilms.

Objectives

The objective of this investigation was to test the efficacy of different laundry agents on biofilms.

Methods

Biofilms grown in rinse water were treated with cleaning bases in a flushing chamber. Several concentration and fluxes of cleaning bases were tested. The attack of the cleaning agents on EPS components was analyzed by confocal laser microscopy after simultaneous staining of bacteria, proteins and polysaccharides.

Conclusions

Our findings showed that cleaning agents did not contribute an attack on specific polymer-groups of the EPS. The selection of the detergent as well as the applied shear forces were of rather great impact. Laundry agents containing oxidative bleach were much more effective than agents containing low or non bleach. CLSM images revealed the compact biofilm structure changed into a small-sized structured biofilm. Worst results were obtained by using liquid detergents. Furthermore, shear forces were of greater impact than the flushing duration and supported the effects of cleaning agents.
RESPONSE OF SESSILE CELLS OF CANDIDA SPECIES TO OXIDATIVE STRESS

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Background

The biofilms of Candida species are associated with high indices of hospital morbidity and mortality [1,2]. Sessile cells are highly resistant to antifungals and to the host mechanisms of defense [1,3-5]. Though the ability of some Candida species to form biofilms on various medical implants has been evaluated as well as their response to oxidative stress [4,6], it is however important to investigate how the reactive oxygen species impacts on sessile cells of different Candida species.

Objectives

To evaluate the response of oxidative stress on biofilm formation by C. albicans, C. glabrata, C. krusei and C. parapsilosis.

Methods

Sessile and control cells of Candida species were exposed to increasing concentrations of H2O2. Biofilms were observed by scanning electron microscopy and quantified by the XTT assay.

Conclusions

To our knowledge, this is the first report describing the effect of oxidative stress on biofilm formation by four Candida species in vitro. Our results suggest that candidiasis in vivo is a multifactorial and complex process where the human immune system and the adaptability of the pathogen should be considered altogether to provide an effective treatment of the patient.
Background

The *fap* operon is present in the majority of *Pseudomonas* species and encodes for the Fap functional amyloid system, which is associated with cell aggregation and enhanced biofilm formation. Six genes make up the operon.

Objectives

However, it is so far unknown how the *fap* operon is regulated.

Methods

In this study, a biotinylated dsDNA probe was constructed in order to isolate potential DNA binding molecules, which specifically associate with the promoter region of the *fap* operon of the amyloid expressing *Pseudomonas* sp. UK4 and potential transcription factors were identified using LC-MS/MS taking advantage of a closed genome sequence. Two global regulators (MvaU and MvaT) were found to interact specifically, with the *fap* promoter. Specific binding of the alternative sigma factor (RpoN) was also observed. The latter observation is interesting as a gene encoding an RpoN activator protein is located next to the *fap* operon. Therefore, we created overexpression and deletion mutants of *Pseudomonas* sp. UK4 with the RpoN activator protein and the two general transcription regulators MvaU and MvaT.

Conclusions

A hydrolysis probe based qPCR assay was developed in order to determine the effect of all these gene regulatory players on the *fap* operon. Consequently, the effect of elevated and decreased levels of c-di-GMP was also studied and experiments will show how these regulators affect transcription of the *fap* operon. This study contributes with an increased understanding of the transcriptional machinery that
control Fap-mediated biofilm formation. In the future, this regulatory system could be an important target in antibiofilm strategies.
EFFECT OF DENTAL BIOMATERIALS ON STREPTOCOCCUS MITIS/HUMAN GINGIVAL FIBROBLASTS CO-CULTURE

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Background

The biocompatibility of oral biomaterials depends not only on the biomaterial properties, but also on the host and oral indigenous microbiota response.

Objectives

A co-culture model of Streptococcus mitis/Human Gingival Fibroblast (HGF) was set up to evaluate the biocompatibility of: 2-hydroxyethylmethacrylate (HEMA), triethyleneglycoldimethacrylate (TEGDMA) and new synthesized bioactive lactose-substituted chitosan-silver-nanoparticles (Chitlac-nAg) in colloidal solution and as thermosets coating.

Methods

S. mitis was co-cultured with HGFs with the above mentioned materials in presence of saliva in terms of bacterial adhesion and aggregation, HGF cell toxicity, expression of apoptosis-associated proteins and intracellular signaling.

Conclusions

HEMA treatment showed an increased aggregation and adhesion of S. mitis on HGFs, mediated by the PKCa/integrin b1 signalling system. HEMA treatment decreased viable eukaryotic cell that was balanced by the presence of bacteria and saliva.

Concerning TEGDMA, S. mitis growth was not decreased and large bacterial aggregates were present on HGFs. Moreover, S. mitis and saliva reduced the HGFs oxidative stress and apoptosis induced by TEGDMA treatment.

Chitlac-nAg colloidal solution, in the HGF/S. mitis co-culture, induced less numerous viable bacteria adhering to the HGFs. Moreover, S. mitis and saliva reduced the HGFs oxidative stress and apoptosis induced by TEGDMA treatment.

Taken together, our results shed new light on the interaction of dental biomaterials/Streptococci/HGFs occurring in oral environment, underling the
importance of the co-culture model, resulting closer to the *in vivo* situation, to obtain a more realistic outcome.
Background

Biocides are commonly used to control unwanted biofilm development in cooling tower. The more environmentally friendly biodispersants can also be used, but little information is available on the efficiency of their application, likely affected by the biofilm structure, composition and growth phase.

Objectives

The objective of this work was to study the structure and diversity of biofilms developed from makeup water used in a full scale industrial cooling tower. The efficacy of the biodispersant was further evaluated on biofilms at different growth phase.

Methods

Biofilms were grown on glass slides, in a batch system, using as source community the cooling tower makeup water under four different nutrient conditions. The taxonomic bacteria affiliation was investigated by Fluorescence In Situ Hybridization (FISH) and diversity index was calculated. EPS were detected in situ by epifluorescence microscopy after staining with fluorochromes. A biodispersant commonly utilized in full scale cooling towers was then used and removal efficiencies at different stage of development were evaluated.

Conclusions

FISH allowed to identify about 90% of total bacteria, mainly belonging to Proteobacteria. Starting from the source community, biofilms showed an initial increase in biodiversity likely due to the functional niche diversification. Then, diversity decreased, mainly due to the adhesion of eukaryotes and to the concurrent reduction in bacteria taxon richness. The removal efficiencies of biodispersant decreased over time likely due to the higher biofilm stability at the late growth stage, where an increase of α and β glucans, mainly found surrounding eucaryotic and
alphaproteobacterial cells, was observed.
ΔppiB ENHANCED MOTILITY AND BIOFILM FORMATION ABILITY IS POSSIBLY MEDIATED THROUGH CHANGES IN THE PPIB INTERACTIONS WITH SOME OF ITS PREY PROTEINS

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Background

E. coli ppiB is a peptidyl-prolyl cis/trans isomerase with chaperone activity. ΔppiB strain shows enhanced swimming and swarming motility as well as biofilm formation ability.

Objectives

We hypothesize that the effects of ppiB on such cellular behaviors are likely accomplished through its protein interactions. Using a candidate approach we aim to identify multicopy suppressors of the ΔppiB phenotypes and to characterize protein interactions with putative ppiB prey proteins.

Methods

Based on data provided by IntAct database we considered available ppiB prey proteins as putative interacting partners. To address whether these proteins could be direct effectors inhibiting motility or biofilm formation we over-expressed each of them in the ΔppiB cells and checked for restoration of the corresponding phenotypes. Further, we examined possible protein interactions with ppiB under native or denaturing prey conditions. To explore the potential involvement of the PPIase active site on the binding of ppiB to these proteins, we determined the inhibitory effect of each of them on the PPIase activity of ppiB.

Conclusions

Both the hyper-motility and enhanced biofilm formation ability of ΔppiB strain was reversed by over-expression of certain prey proteins albeit different in each condition. With very few exceptions, we were not able to detect stable protein complexes, probably an indication for weaker transient interactions. However, under denaturing prey conditions, more interactions were revealed suggesting that ppiB is able to recognize denatured preys. A decrease in the ppiB PPIase activity, in the presence of many of them, further supports many of the putative associations.
EFFECT OF SHEAR STRESS ON PSEUDOMONAS AERUGINOSA ISOLATED FROM THE CYSTIC FIBROSIS LUNG

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Background

Chronic \textit{Pseudomonas aeruginosa} lung infections are the main cause of morbidity in cystic fibrosis (CF) patients. Previously, we have sequenced the genome of the transmissible Belgian epidemic strain CF_PA39. This CF-adapted strain has been present in the UZ Brussel hospital environment for at least ten years. Furthermore, earlier experiments showed that culturing \textit{P. aeruginosa} PAO1 in a low fluid shear environment, obtained by means of the Rotating Wall Vessel (RWV) technology, leads to the formation of a biofilm phenotype comparable to that observed in the CF lung.

Objectives

To study the effect of fluid shear on biofilm formation by an adapted \textit{P. aeruginosa} CF strain in artificial sputum medium (ASM) at the genotypic, transcriptomic, and phenotypic level.

Methods

In this study, an RWV experiment was performed that closely resembled the \textit{in vivo} situation by inoculating this adapted CF isolate in ASM in the RWV either with or without two glass beads, simulating high and low fluid shear conditions, respectively. Scanning electron microscopy (SEM) was utilized to determine biofilm formation in both conditions, while RNAseq and qPCR was used to study gene expression in both conditions.

Conclusions

Increasing fluid shear in the RWV model disrupted biofilm formation of an adapted \textit{P. aeruginosa} CF isolate.

In accordance with the biofilm disruption, several genes involved in denitrification, tryptophane synthesis, choline metabolism, and alginate biosynthesis were down-regulated in the high fluid shear condition, resembling the planktonic stage of growth. Furthermore, we identified small RNAs that are differentially expressed.
Background
It has been shown that zinc oxide (ZnO) nanostructures can photocatalytically inhibit growth of fresh water bacterial and fungal strains under solar irradiation.

Objectives
The objective of this study was to investigate the prevention of formation of marine biofilms by ZnO nanorod coatings in the laboratory and out-door experiments.

Methods
ZnO nanorod coatings were fabricated on microscope glass substrates by a simple hydrothermal technique using equimolar solutions of 10 mM zinc nitrate and hexamethylenetetramine. In laboratory conditions, upon 5 h of white light irradiation (light intensity= 100 Klx from a tungsten-halogen lamp), nanorod coatings significantly reduced the growth of marine bacterium Acinetobacter sp. and inhibited biofilm formation as compared to coatings kept in the dark (not exposed to light). In the continuous 7 days out-door experiment conducted in an aquaria (volume = 70 L) with sea water collected from Oman sea (Muscat, 23°34′55″ N, 58°36′27″ E), the nanorod coatings significantly reduced density of bacteria in comparison to the control (no coatings). In the absence of sunlight, test and control slides were equally colonized by bacteria. De novo sequencing of bacterial biofilms on MiSeq system® demonstrated that different communities were formed in the presence and absence of light on nanorod coatings.

Conclusions
Our study suggests that ZnO nanorod coatings effectively prevent biofilm formation and can be used as a novel green antifouling technology.
Background

*Acinetobacter baumannii* is a pathogen with a high quality to form biofilm which can be isolated from different environmental sources. In addition, pathogenicity and biofilm formation associated with infectious agent.

Objectives

In this study, we investigated that degradation of biofilm formation by *Acinetobacter baumannii* is determined with amylase enzyme from *Bacillus amyloliquefaciens*.

Methods

*Bacillus amyloliquefaciens* is amylase enzyme producer which was obtained from Hacettepe University Department of Biology/Biotechnology. The biofilm forming ability in clinical isolates of *Acinetobacter baumannii* was determined by both microscopic and spectrophotometric analyses with using 24-well polystyrene plates.

Conclusions

According to results, amylase which obtained from *Bacillus amyloliquefaciens* was observed to inhibit biofilm formation in *Acinetobacter baumannii*. In continuous, amylase enzyme prevent to increase *Acinetobacter baumannii* cell population. As well as, in most of pathogen bacteria like *Acinetobacter baumannii*, amylase enzyme is considered to degradation of biofilm formation. In a result, amylase enzyme can be used as a drug supplement.
EFFECT OF SUBINHIBITORY CONCENTRATION OF SILVER NANOPARTICLES ON ADHESION OF GRAM-POSITIVE AND GRAM-NEGATIVE BACTERIA ONTO POLYSTYRENE SURFACES

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Background
In biofilms, bacterial cells adhere irreversibly to a solid surface by extracellular polymeric substances. This way of life provides several advantages over the planktonic mode, as responses to environmental stress.

Objectives
The objective was determined the effect of subinhibitory concentration of silver nanoparticles (Ag-NPs) and silver nitrate (AgNO₃) on adhesion of gram-positive and gram-negative bacteria.

Methods
Ag-NPs were synthesized by reduction of AgNO₃ with sodium citrate. Bacterial strains used were gram-positive Staphylococcus aureus ATCC 6538, Bacillus cereus, isolated from milk cooling tank and Enterococcus faecalis ATCC 51299 and gram-negative Escherichia coli ATCC 11229, Salmonella enterica serovar Typhimurium ATCC 13076 and Pseudomonas aeruginosa ATCC 15442. The minimum inhibitory concentration was determined by macrodilution technique according CLSI, 2003. Müller-Hinton broth supplemented with subinhibitory concentration of Ag-NPs or AgNO₃ were transferred to 96-wells polystyrene microtiter plates, and inoculated with 5.0 x 10⁵ cfu/ml of cell suspension. After 48 h at 37°C, the optical density was determined at 600 nm and supernatant was discarded. The surface-attached cells were stained with crystal violet for 30 min and washed three times with water. The absorbance at 590 nm was determined by addition of ethanol.

Conclusions
Adhesion was estimated by calculating the ratio between the absorbance of violet crystal and the optical density of cells. Subinhibitory concentrations of Ag-NPs and AgNO₃ increased the adhesion of the gram-positive bacteria, probably, as response to stress. The adhesion of E. coli and S. Typhimurium was not affected by presence of Ag-NPs or AgNO₃. However, P. aeruginosa adhesion decreased in the presence of Ag-NPs.
CANDIDA GLABRATA BIOFILMS RESPONSE TO AMPHOTERICIN B

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Background

Candida species are responsible for recurrent human infections, mostly in immunocompromised patients, due to their high vulnerability. Candida glabrata has been showing to have a major role in these infections being the second most prevalent species involved in human fungemia. Amphotericin B (AmB), a common antifungal drug, is a hospital-environment exclusive polyene, normally being efficient when used to fight candidiasis.

Objectives

The main goal of this work was to infer about the influence of AmB in Candida glabrata biofilms formation and its effect on matrix composition and ERG genes expression.

Methods

Candida glabrata biofilms were formed in the presence of AmB and analyzed by dry weight. Moreover, ERG genes expression was evaluated by qRT-PCR and matrix was analyzed in terms of composition in carbohydrates, proteins, beta-glucans and a new finding: ergosterol.

Conclusions

In addition to an inefficient reduction of the C. glabrata biofilms, this work showed that ERG genes seem to be less involved than the matrix composition in C. glabrata biofilms response to AmB. Specifically, C. glabrata biofilms matrices respond with an increase of carbohydrates, particularly beta-1,3 glucans, and with a decrease of total proteins. The ergosterol values did not expressively changed in the presence of AmB.

The present work support the theory of multifaceted mechanisms developed by C. glabrata biofilms as response to the presence of AmB.
FLOW CELL ANALYSIS OF THE PROCESS OF BACTERIAL CELL ADHESION MEDIATED BY THE ADHESIVE NANOFIBER PROTEIN ATAA UNDER SHEAR STRESS

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Background

Elucidation of bacterial adhesion is important for preventing detrimental biofilm formation and for applying it to microbial immobilization. An environmental strain, Acinetobacter sp. Tol 5 shows nonspecific, remarkably high adhesiveness to solid surfaces through AtaA (Acinetobacter trimeric autotransporter adhesin) nanofibers.

Objectives

We aim to examine the effects of flow rate on the adhesion and autoagglutination of resting Tol 5 cells to reveal adhesion property mediated by AtaA.

Methods

Bacterial adhesion was analyzed through using a flow cell system and the shear stress of the flow was controlled by flow rates and the viscosity of cell suspension supplied with PEG-400. The biomass of attached cells was quantitatively measured by staining with crystal violet. The thickness of the cell clump adhered on the surface was measured by confocal laser scanning microscopy.

Conclusions

Higher shear stress in the flow cell decreased bacterial adhesion and the thickness of cell clump formed on the flow cell surface. By contrast, the bacteria adhesion to well-plates was increased by mild agitation compared with that in a static condition due to enhanced autoagglutination of the cells. These results suggest that there is the optimum flow rate for cell adhesion in the balance between enhancement of autoagglutination due to the increase in the collision probability of bacterial cells and detachment of cells and/or inhibition of cell-cell or cell-surface interaction by shear stress.
HRPB DEAD-BOX RNA HELICASE IS IMPORTANT FOR MOTILITY, BIOFILM FORMATION AND CANKER DEVELOPMENT IN XANTHOMONAS CITRI SUBSP. CITRI

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Background

DEAD-box RNA helicases are proteins highly conserved, present in all eukaryotic cells and in many bacteria. DEAD-box RNA helicases are involved in various processes of RNA metabolism and play important roles, e.g., translation initiation, ribosome biogenesis and RNA decay. Several bacterial DEAD-box proteins have been studied extensively due to their importance in *E. coli*, *Bacillus subtilis* and *Staphylococcus*. Interestingly, in *Staphylococcus aureus*, the DEAD-box protein CshA was involved in biofilm formation via modulation of *agr* mRNA stability. However, the function of DEAD-box RNA helicases remains unexplored in relation to adhesion, biofilm formation or pathogenicity in *Xanthomonas citri*.

Objectives

To characterize the function of DEAD-box RNA helicase (XAC0293/HrpB) in relation to motility, biofilm formation and citrus canker development in *Xanthomonas citri* subsp. *citri*.

Methods

We conducted assays for analysis of sliding motility, biofilm formation on biotic surface and pathogenicity of the wild type and the *hrpB* mutant of *X. citri* subsp. *citri*. Expression of different genes between the wild type and mutant strains was determined by quantitative RT-PCR.

Conclusions

Deletion of the *hrpB* gene reduced the expression of type IV pili genes, which are required for motility, biofilm development and adherence. Our data show that HrpB regulates the mobility and adherence of the *X. citri* in host. HrpB hereby is the first DEAD-box RNA helicase in *Xanthomonas citri* that is implicated in the regulation of
pili genes and that it is involved in motility, biofilm formation and citrus canker development.
COMPARATIVE GENOMICS OF IRON TRANSPORTING SYSTEMS AND FUNCTIONAL ANALYSIS IN BACILLUS CEREUS STRAINS FOR GROWTH AND BIOFILM FORMATION

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Background

Human pathogens such as \textit{Bacillus cereus} are in constant competition for iron with other microbes and their host. In most environments, including food and mammal hosts, iron is bound to complex compounds and is not readily available. However, pathogenic bacteria developed mechanisms to scavenge iron from a range of sources.

Objectives

The objective of this study was to investigate the use of different complex and non-complex iron sources by \textit{Bacillus cereus} reference strains ATCC14579 and ATCC 10987, and 20 undomesticated food isolates, for which growth phenotypes were linked to genotypes.

Methods

Genome sequences were analysed for the presence of putative iron transport systems. Growth and biofilm formation were monitored in LB with and without iron scavenger Bibyridine and different iron sources. Transcriptome studies in \textit{B. cereus} ATCC10987 revealed upregulation of many putative iron transport systems upon depletion of iron.

Conclusions

All 22 \textit{B. cereus} strains could effectively use Fe citrate and FeCl\textsubscript{3} for growth, and formation of air-liquid interface biofilms was promoted. Hemoglobin and Hemin were used by all, except one strain lacking functional petrobaictin and IIsA systems. Interestingly, Hb and Hemin triggered a submerged type biofilm formation by several strains. Ferritin and transferrin could be used for growth only by three and six strains respectively, but biofilm formation was inhibited. Lactoferrin did not restore the growth, however it supported submerged biofilm formation by several strains. Knowledge of iron transport systems and their functioning can serve as an indicator for bacterial fitness in different environments including usage of host-derived iron sources.
MIcroscopic analysis of the process of bacterial cell adhesion and autoagglutination mediated by the adhesive nanofiber protein AtaA

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Background

Elucidation of bacterial adhesion is important for preventing detrimental biofilm formation and for applying it to microbial immobilization. An environmental strain, Acinetobacter sp. Tol 5 shows nonspecific, remarkably high adhesiveness to solid surfaces through AtaA (Acinetobacter trimeric autotransporter adhesin) nanofibers.

Objectives

We aim to reveal the mechanism of cell adhesion mediated by AtaA and describe the cell adhesion process in a microscopic level.

Methods

The time profile of cell adhesion was analyzed through confocal laser scanning microscopy (CLSM) and flow cell analysis. The mechanism of autoagglutination was described by clustering process of colloidal particles by measuring the surface potential of Tol 5 cells with and without AtaA nanofibers.

Conclusions

By the CLSM, Tol 5 wild-type (WT) but not ΔataA mutant was revealed to form a thick cell layer (> 30 μm) with void structure on a glass surface. The process of the cell clumping was described by DLVO theory and CCA model, which were developed as colloidal aggregation models. The autoagglutination of Tol 5 cells was realized by Brownian motion and the collision of AtaA fibers, but not that of cell bodies. By flow cell analyses, it was demonstrated that autoagglutinated WT cells attached well to a glass surface, whereas its single cells hardly attached. These results suggest that autoagglutination greatly contributes to cell adhesion due to enlarged particle size.
Background

*Candida albicans* is a major fungal pathogen of humans. It is a commensal on the skin and mucosal surfaces, but can cause invasive and systemic infections particularly in the immunocompromised patients. *C. albicans* can easily form biofilm on supporting surfaces, such as mucosal surfaces and indwelling medical devices. Biofilms enhance *C. albicans* resistance to antifungals and may help cells to escape from the host immune systems.

Objectives

The identification and study of transcription factors related to biofilm formation can provide important insights into molecular mechanisms controlling biofilm formation.

Methods

The systems biology approach is a novel trend in biological research that focuses on the complex interactions among genes, proteins, and intracellular metabolites. Using systems biological approach, we identified several transcription factors that may involve in biofilm formation. *MSS11*-deleted mutant showed a defect in forming a mature biofilm and partially attenuates the virulence of *C. albicans* in a mouse model of infection. In contrast, deletion of the gene encoding another transcription factor enhanced cell adhesion and biofilm formation compared to wild type. Interestingly, this mutant also showed an increase in the expression of adhesin genes.

Conclusions

Taken together, our results support applying the systems biology approaches to study various aspects of fungal biofilm formation and pathogenesis.
INTERFERENCE OF BIOFILM FORMATION BY LIPOPOLYSACCHARIDE O-ANTIGEN AND THE CAPSULE-LIKE COMPLEX OF FRANCISELLA TULARENSIS

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Background
Francisella novicida forms a robust biofilm on most surfaces, whereas F. tularensis subsp. tularensis and subsp. holarctica form little or only a poor biofilm. F. novicida colonies are also dark red on Congo Red agar, whereas F. tularensis colonies are light salmon. However, colonies of F. tularensis lipopolysaccharide O-antigen mutants are also dark red in color. We hypothesized that the O-antigen, or other surface components, of F. tularensis interferes with biofilm formation.

Objectives
Our objectives were to determine if mutants of F. tularensis lacking O-antigen and/or a capsule-like complex (CLC) were more capable of forming a robust biofilm than parent strains.

Methods
F. novicida, F. tularensis (types A and B), and mutants were grown in polystyrene wells or on glass coverslips, and biofilms examined by crystal violet staining, confocal laser scanning microscopy, and scanning electron microscopy. Biofilm-associated exopolysaccharide (EPS) was isolated by enzyme digestion and phenol extraction, and composition determined by gas chromatography-mass spectrometry.

Conclusions
O-antigen mutants of F. tularensis made a more robust biofilm than parent strains within 10 days incubation. However, only a mutant lacking both O-antigen and CLC attached better and made significantly more biofilm than the parent after 15 days incubation. Some F. novicida O-antigen mutants also formed significantly more biofilm than the parent. Optimum biofilm formation was dependent upon the growth medium. A novel EPS was isolated and identified as glucan from the F. tularensis biofilm. We conclude that the O-antigen and CLC interfere with F. tularensis biofilm formation, likely through interfering with attachment.
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Biofilms

3D VISUALIZATION AND QUANTIFICATION OF BIOFILM IN POROUS MEDIA BY X-RAY TOMOGRAPHY
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Background

The development of reliable models of biofilm growth in biofilters at the industrial scale, suffers from a lack of 3D measurements at the scale of the biofilter’s constituents. Currently, only three experimental studies, using Synchrotron X-ray Microtomography, proposed protocols based on the use of different contrast agents to image such complex media at the pore scale.

Objectives

The aims of the present study, based on the adaptation of these protocols to laboratory X-ray microtomograph, are threefold:

(i) the 3D visualization and quantification of the interface between the biofilm and the bulk fluid. This will give us access to the shape, the spatial distribution and volume of the biofilm at the pore scale;

(ii) the numerical evaluation of physical properties such as permeability or diffusivity arising in macroscopic models using up-scaling methods;

(iii) the evolution of the aforementioned biofilm characteristics with time and according to the operating conditions (flow rate, nutrient conditions.)

Methods

Thus, the Pseudomonas putida biofilms were grown on glass beads in small biofiltration columns. A new protocol permitting to avoid the use of contrast agent was proposed and 3D X-ray images of the biofilms grown under various operating conditions were obtained (i,iii).

Conclusions
The 3D microstructural properties were quantified using appropriated image processing tools (i) and the physical properties were estimated using numerical tools (ii). The representativity of the imaged volume was checked for both structural and physical properties giving access to some of the missing elements to build the macroscopic model.
ANTI-BIOFILM FORMING EFFECT OF ESSENTIAL OILS ON LISTERIA MONOCYTOGENES IN MONO- AND MIXED CULTURES

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Background

Listeria monocytogenes can be found frequently in mono- or mixed culture biofilms. These make sanitization difficult and can lead to cross-contamination and food safety problems. Beside heat treatment, preservatives can be used to eliminate pathogens but sometimes they are not effective and can produce unpleasant by-products. Application of natural antimicrobial agents like essential oils (EOs) could be a potential solution.

Objectives

Our aim was to investigate the anti-biofilm forming effect of selected EOs and their major components on L. monocytogenes biofilms and on mixed-cultures of L. monocytogenes and E. coli.

Methods

MIC values were determined by micro-dilution method. Biofilm formation was conducted in microtiter plates; after 4 h adhesion time, fresh medium containing EOs or components was added in MIC/2 concentration. Biofilm formation was monitored by crystal violet staining. For mixed cultures L. monocytogenes and E. coli were mixed in 1:1 ratio and concentration intervals between the MIC values of the two bacteria were used.

Conclusions

In case of monocultures, cinnamon and cinnamaldehyde have the best anti-biofilm formation effect. For mixed cultures, the effect of the EOs was concentration dependent: α-pinene and linalool showed better anti-biofilm forming effect than the parent essential oil.
CHARACTERIZATION OF NaCl STRESS-INDUCIBLE PALMELLOIDS AND UNDERSTANDING THE ROLE OF A STRESS-RESPONSIVE GENE, BOLA FROM THE GREEN CHLOROPHYTE CHLAMYDOMONAS REINHARDTI.

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Background
Chlamydomonas reinhardtii, a unicellular green alga, is a preferred model organism for studying stress-induced responses. Depending on the stress dose and time of exposure, the flagella of this organism are paralyzed / lost and later cells manifest ‘palmelloids’ or apoptose. Palmelloids, akin to bacterial biofilms, are a stress-responsive temporary, dormant ‘colonial’ stage exhibiting conserved physiological changes.

Objectives
The current study deals with characterization of palmelloids formed by C. reinhardtii in response to NaCl stress and explores the role of a conserved stress-responsive gene family, C. reinhardtii bolA in biofilm / palmloid formation.

Methods
A detailed morphological and biochemical analysis of NaCl stress inducible palmelloids was performed. Drawing an analogy, it is known that the morphogene, bolA, significantly facilitates the formation of biofilms and induces morphological changes in response to stress in Escherichia coli. Chlamydomonas reinhardtii genome harbors five bolA-like genes. The ORF with highest homology to algal systems (CrbolA-4) was cloned and the protein over-expressed in Escherichia coli.

Conclusions
Palmelloids rapidly dissociated upon de-stress suggesting the reversibility of this phenomenon. The cells in palmelloids displayed conserved physiological features such as clustering of cells in a common envelope, decreased viability, accumulation of lipid and starch granules and presence of extracellular polysaccharides. An iTRAQ analysis of the spent medium from palmelloid-containing cells suggests a strong involvement of cell wall proteins. Further heterologous over-expression of CrbolA-4 in E. coli did not affect its growth; but, induced biofilm formation and changed its morphology, indicating functional conservancy. Its role in the NaCl-induced palmelloidy remains to be explored.
Background
Ornithine lipids (OLs) are interesting bacterial lipids that are widely found in outer membrane of many Gram-negative bacteria, but not detected in Eukarya and Archaea. Pseudomonas aeruginosa has olsBA operon encoding acyltransferases that functions the OL biosynthesis. OlsBA works in two steps for the OL biosynthesis, in which OlsB transfers an acyl group to ornithine to make lyso-ornithine lipid and OlsA converts the lyso-ornithine lipid into ornithine lipid by another acyl-group transfer.

Objectives
Acyltransferase genes of P. aeruginosa were originally screened for the effect on the virulence and biofilm formation of P. aeruginosa. The olsBA operon was found to have significant pleiotropic effects on the virulence related-phenotypes of P. aeruginosa. In this study, we addressed how OlsBA modulates the virulence and biofilm formation of P. aeruginosa.

Methods
1) The virulence was investigated by using two different host models, Tenebrio molitor, an insect and Caenorhabditis elegans, a nematode. 2) The effect of the olsBA overexpression on biofilm formation in P. aeruginosa was analyzed in flow cell system. 3) The host response to OLs was monitored through the expressions of COX-2 and iNOS using Western blot and real-time PCR analyses.

Conclusions
We found that the overexpression of olsBA operon modulated some virulence related-phenotypes of P. aeruginosa by reducing the quorum sensing response. In addition, the overproduced OLs directly alleviate the virulence of P. aeruginosa, enhances biofilm formation, and modulates the host physiology.
THE NITROGEN DIOXIDE INCREASES PSEUDOMONAS FLUORESCENS BIOFILM FORMATION: IDENTIFICATION OF THE BACTERIAL RESPONSE TO AIR POLLUTANT

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Background
Nitrogen dioxide is an air pollutant of increasing interest in biology. Exposure of animals to NO₂ results in several toxic effects, mostly lung injury.

Objectives
Aware of the effect of NO₂ on pseudomonal biofilm formation, we looked for the mechanisms of bacterial response that could explain their resistance to NO₂.

Methods
The confocal microscopy was used for biofilm studying, completed by motility assays. In a second time, in silico and in vitro genomics tools were used to narrow our research scope. Additionally, in order to characterize the mechanism of P. fluorescens resistance to NO₂ pollution these data were completed with proteomics and lipidomics studies. For this purpose, the MALDI-TOF MS Imaging was coupled to HPTLC and compiled with the traditional GC-MS.

Conclusions
When the NO₂ increases P. fluorescens biofilm formation, the bacterial motility decreases. In coherence, the level of cyclic di-GMP evolves in NO₂ exposed cells. The lipid study shows no drastic change in membrane charge and its composition in phospholipids and fatty acids. This suggests that NO₂ free radical could freely pass through membrane. In contrast, NO₂ promotes an extensive modification in protein production, notably the over-production of the proteins of stress response, involved in oxidative stress tolerance and iron transport/metabolism. Thus both iron up-regulation and C-di-GMP level could be signals for biofilm development. This knowledge should probably offer therapeutical solutions in antibacterial treatment.
Background

Bacteria adapt to unfavorable environmental conditions for survival. *Burkholderia glumae*, the causal agent of rice panicle blight, absolutely requires oxygen for respiration. For aerobes such as *B. glumae*, oxygen limitation is a serious problem for survival. Many bacteria including *B. glumae* possess genes encoding proteins with a PAS/PAC domain whose functions sense changing environments and manage gene regulation accordingly. Oxygen availability in bacteria is related to biofilm or pellicle formation. Cyclic-di-GMP synthesized by proteins carrying a GGDEF domain is a messenger for the pellicle formation.

Objectives

We aimed to investigate how *B. glumae* adapts to oxygen availability.

Methods

Overnight culture of *B. glumae* grown in LB broth was serially diluted with the same medium to give approximately $1 \times 10^7$ cells/ml. A portion of 200 µl was placed in 96 well cell culture plates and incubated at 37°C without shaking. Growth was monitored from day 0 to day 7 of the incubation period. Morphologically altered colonies were isolated, and a mutation responsible for colony variation was identified by whole genome re-sequencing.

Conclusions

Mutations in the *bpaA* (*B. glumae* pH associated protein A) gene encoding a protein possessing PAS/PAC and phosphatase domains conferred wrinkly colony types. A mutation in *bpaA* resulted in elevated expression of cellulose biosynthesis genes responsible for pellicle formation. Thus, mutations in *bpaA* and pellicle formation in oxygen limited conditions suggest how *B. glumae* has evolved to survive under unfavorable conditions.
ROLE OF EACH EPM GENE ON THE FORMATION OF A NOVEL BIOFILM-SCAFFOLD EXTRACELLULAR POLYMER FROM PSEUDOMONAS

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Background

Pseudomonas alkylphenolia KL28 forms unique three-dimensional biofilm bodies depending on the environmental stimuli. It forms highly branched mushroom-like aerial architecture during vapor p-cresol utilization and circular floating pellicles on standing LB medium. These indicate that P. alkylphenolia is a good bacterial sample for studying bacterial community formation and cell-cell interactions. An epm gene cluster consisting of eleven genes showing deduced amino acid sequence similarities to those encoded on alg genes has been identified to be responsible for formation of the special biofilm structures.

Objectives

In this study, the role of each epm gene was studied in relation to the biofilm formation and polymer processing.

Methods

For this purpose, mutants specifically deleted in epm genes from an epm-overproducing spontaneous mutant SG1-WC1-10⁷ were constructed. With those mutants, changes in surface-related phenotypes and polymer production were examined.

Conclusions

It was found that epm genes except epmL are required to form wrinkle colonies and mature biofilms including aerial architecture and circular floating bodies. Mutants in epmI, epmJ, epmX (acetylation, deduced from alg genes), epmG (epimerization), epmE, epmK (transporter) were still positive to polyuronic acid production. In addition, some of these epm genes cannot be substituted by alg genes. Thus, these results indicate that regardless of the similarities between Alg and Epm proteins the structure of Epm polymer derived from an epm gene cluster is different from alginate and the process of Epm polymer biosynthesis is different from that of alginate in Pseudomonas.
Background

Anthranilate and indole are alternative degradation products of tryptophan in nature. *Pseudomonas aeruginosa* produces anthranilate from tryptophan degradation. Previous studies showed that anthranilate and indole has an opposite effect on *P. aeruginosa* biofilm formation. While indole enhances the biofilm formation of *P. aeruginosa*, anthranilate was able to disintegrate the mushroom structure of biofilm.

Objectives

We addressed how indole could enhance the biofilm formation of *P. aeruginosa* in relation to anthranilate degradation.

Methods

*P. aeruginosa* biofilm formation was analyzed in flow-cell system. High Performance Liquid Chromatography was used to measure the level of anthranilate in *P. aeruginosa* culture supernatant. The effect of indole on AntR activity was analyzed by reporter fusion *in vivo* and gel shift assay *in vitro*.

Conclusions

In *P. aeruginosa*, the transcription of antABC operon encoding anthranilate dioxygenase complex that functions to degrade anthranilate, was activated by the indole treatment. Although indole alone failed to activate AntR, co-addition of indole with anthranilate boosted the activation of AntR by anthranilate. As consequence, the anthranilate level in *P. aeruginosa* culture supernatant decreased. Since anthranilate has a biofilm-disintegrating effect, we suggest that indole may enhance the biofilm formation of *P. aeruginosa* by reducing anthranilate level through the modulation of AntR activity.
TIMERIC AUTOTRANSPORTER ADHESINS (TAAS) - STRUCTURAL STUDIES OF THE TRANSLOCATION PROCESS, AND PERSPECTIVES FOR ITS INHIBITION

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Background
Trimeric autotransporter adhesins (TAAs) comprise a widespread family of adhesive molecules in Gram-negative bacteria, many of which are important virulence factors [1]. The prototypical trimeric autotransporter is the Yersinia adhesin YadA from Yersinia enterocolitica, which mediates attachment to collagen and other extracellular matrix (ECM) molecules, promotes serum resistance and mediates autoagglutination.

Objectives
The autotransport process depends on a conserved translocation domain and associated factors. Inhibiting this process is a possible antimicrobial strategy, as no homologous systems exist in Eukaryotes, and as many of the TAAs are important or even essential in host colonization by a broad range of Gram-negative pathogens.

Methods
We recently solved the structure of the Yersinia YadA translocation domain (using solid-state NMR [2]) and of a chaperone-like protein involved in the translocation of Salmonella SadA (using x-ray crystallography [3]). Conserved residues thought to be important for translocation are targeted by mutagenesis. Adhesion assays and antibody-based assays such as FACS are used to quantify translocation.

Conclusions
A better understanding of the translocation process on the molecular level is essential for efficient inhibition of the surface display of these important pathogenicity factors. The methods developed to quantify surface localization can be scaled up for systematic screens for antimicrobial substances.

[3] Leo et al., JBC 2014
DISSECTING METHANO- AND METHYLOTROPHIC BIOFILM FUNCTIONING IN A IODINE-RICH SUBSURFACE SPRING CAVERN

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Background

Massive microbial biofilms have recently been discovered in a subsurface spring cavern fed by iodine-rich deep formation water with high thermogenic methane loading. The biomass produced by the microbes and exopolysaccharide slime completely covers the walls and ceilings of the cave, the latter bearing snottites of up to 10 cm length.

Objectives

However, the nature of these unique biofilms as well as their ecophysiology are not yet understood.

Methods

Our primary taxonomic characterization of the biofilms revealed a surprisingly diverse polymicrobial community assembly, substantiating a complex network of bacteria and archaea interacting within the sticky matrix. Distinct methanotrophic and methylotrophic populations within the Alpha-, Beta- and Gammaproteobacteria dominated the biofilms. Methylophilaceae and Methylococcaceae were abundant on the walls and especially in the cavern water, but not so in ceiling biofilms. Here, ribosomal and functional gene analysis hinted at dominating methylotrophic and potentially also iodine-cycling populations, thus supporting a hypothesized cycling of iodomethane in this system. Geochemical gradients in the cavern were clearly reflected within community assembly, as well as elemental and stable isotope composition of the biofilms. These findings were further supported by distinct microaerophilic oxidation rate measurements for methano- and methylotroph substrates in spatially resolved biofilm samples. Ongoing work involves the elucidation of ecological niche partitioning between strict methanotrophs and potential methyl halide-oxidising populations in this unique habitat.

Conclusions
In conclusion, primary insights into a dominantly chemolithoautotrophic natural biofilm system are revealed, thriving on deep subsurface energy inputs but only a few meters under our feet.
EFFECT OF SUBINHIBITORY CONCENTRATIONS OF CEFOTAXIME ON BIOFILM FORMATION IN STREPTOCOCCUS PNEUMONIAE

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Background

Streptococcus pneumoniae forms biofilm to persist in the human nasopharynx. Biofilms are often associated with pneumococcal pneumonia and otitis media. β-lactam agents remain the first line antibiotic for the treatment of S. pneumoniae infections. Subinhibitory concentrations of antibiotics have been reported to enhance biofilm formation.

Objectives

This study aims to evaluate the effect of subinhibitory concentrations of a third generation cephalosporin, cefotaxime (CTX) on biofilm formation in S. pneumoniae.

Methods

The effect of CTX (at 1/2, 1/4 and 1/16 MIC) were tested against biofilm formation in R6 and four clinical isolates in 24-well microtiter plates. Crystal violet (CV) staining and colony forming unit (CFU) were used to quantify biofilm biomass. Similar experiments were carried out with erythromycin and ciprofloxacin.

CV staining results showed that 1/2 MIC CTX resulted in 52.3%, 64.9%, 68.2%, 69.5% and 84.5% (all \( p \) values <0.01) reduction in biomass of biofilm for R6 and the clinical strains tested. 1/4 MIC CTX led to 13.2% to 57.5% (\( p \) value <0.01) reduction of biofilm while the inhibitory effect of 1/16 MIC CTX on biofilm formation were insignificant (<20% inhibition). Living bacterial cells of R6 and two cefotaxime resistant strains recovered from biofilms unexposed to CTX were 1.5×10^8, 3.6×10^4 and 1.3×10^6 CFU/well, respectively and decreased to 4.7×10^7, 53 and 43 CFU/well, respectively when exposed to 1/2 MIC CTX. Subinhibitory concentrations of non-β-lactam antibiotics showed no significant influence on biofilm formation.

Conclusions

Subinhibitory concentrations of CTX significantly inhibited biofilm formation.
CHARACTERIZATION OF A NAPHTHALENE-DEGRADING MULTI-SPECIES BIOFILM ISOLATED FROM OIL-CONTAMINATED GROUND WATER

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Background
Environmental microorganisms exist predominantly as biofilms and gain high tolerance to environmental stresses. Biofilm systems are especially suitable for the treatment of recalcitrant compounds as polycyclic aromatic hydrocarbon (PAH) because of their high microbial biomass and ability to immobilize compounds. To our knowledge, reports on naphthalene degradation by biofilm-forming bacteria are limited. In this study we isolated multi-species biofilm degrading naphthalene from oil-contaminated ground water (oil refinery, Murcia, Spain).

Objectives
The aims of this work were to characterize the biofilm microbial community and the biofilm structure and to isolate the bacteria involved in the biofilm formation and degradation of naphthalene.

Methods
The microbial community composition was characterized by culture dependent and molecular methods. For a qualitative understanding of biofilm structure and composition we used Scanning Electron Microscope (SEM). We also designed degenerated probes to detect and amplify naphthalene dioxygenase nahA gene homologues from environmental DNA and bacterial isolates.

Conclusions
Molecular analyses showed that the initial community was dominated by Betaproteobacteria (79% *Rhodoferax*, 6% *Azovibrio*). After enrichment on naphthalene minimal medium we found Betaproteobacteria (54% *Variovorax*) and Alphaproteobacteria (43% of a *Xanthobacteraceae* strain) as dominant bacteria. We isolated members of *Microbacterium*, *Starkeya*, *Rhizobium*, *Brevundimonas*, *Variovorax*, *Pseudomonas*, *Pseudoxanthomonas*, *Epilithonimonas* and *Aquabacter*. Some of the isolates degraded naphthalene in pure culture and some other were able to produce biofilm. Homologues of *nahA* genes could be retrieved from the enrichment cultures and from some isolates.
NITRIC OXIDE (NO) DONORS TO DISPERSE BIOFILMS OF INDUSTRIAL SIGNIFICANCE FORMED BY SALMONELLA ENTERICA, PATHOGENIC ESCHERICHIA COLI AND LISTERIA SP.

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Background

Biofilms formed in post-harvest production facilities are recalcitrant reservoirs of pathogens, which are difficult to control. Pathogens in biofilms are resistant to common disinfectants and their mechanical removal is only partially effective. Therefore, novel approaches for controlling biofilms are needed. Recent discoveries of the function of nitric oxide in dispersing preformed biofilms offer an opportunity to test the feasibility of using this gas in industrial applications.

Objectives

Determine the effect of several nitric oxide donors on biofilm of Salmonella enterica, Escherichia coli and Listeria innocua preformed on surfaces of industrial interest.

Methods

Biofilms were pre-formed in appropriate media for 24 hours or 1 week in 96-well plastic plates (polypropylene, polystyrene). After incubation, medium was removed and nitric oxide donors were resuspended in Phosphate Saline Buffer (PBS) and added to the wells using different concentrations. Biofilms were exposed to Molsidomine and MAHMA-NONOate donors from 2 hours to 6 hours at 22°C and 4°C to mimic the post-harvest environments. Dispersal was measured by staining the remaining biofilms using crystal violet or using a GFP labeled Salmonella strain. The donors were also tested in association with a cellulose nanocrystals hydrogel (CNC).

Conclusions

Molsidomine and MAHMA-NONOate were able to disperse at least 50% of the biofilms preformed by human pathogens, Salmonella enterica 14028, Escherichia coli O157 (EHEC). The association of the two nitric oxide donors with cellulose nanocrystals was also effective in dispersing Salmonella preformed biofilms on polypropylene up to 15% of the total biomass. Our results show that nitric oxide donors expand the toolset of proactive solutions for removing industrial biofilms.
Effect of pomegranate peel extract on biofilm and planktonic cells of *Candida albicans*

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Background: Pomegranate peel is used by many people against diarrhea, ulcers, periodontal disease, stomatitis and pharyngitis. *Candida albicans* can infect any part of the body (gum, tongue, toes...) with tissue invasion thus inducing candidiasis. *Candida* infections are difficult to manage due to their persistent nature and the biofilm growth of this yeast serves as an increasing source of clinical infections.

Objectives: This study investigates the antioxidant activity of acetone peel extract of sour pomegranate peel named Quares (QE) and its effect against biofilm and planktonic cells of *Candida albicans*.

Methods: Acetone 50% (v/v) was used for phenolic compounds extraction. The antifungal activity is evaluated using three strains of: CA1, CA2 and CA3. Various concentrations have been used to determine the MIC. Preliminary screening of activity: The agar disc diffusion method was conducted. The concentrations tested are 10mg/disc to 0.5mg/disc. Two models were used for the biofilm formation inhibition of *Candida albicans* on micoplaque (Hamada et al., 1978) and on hydroxyapatite discs (Guggenheim, 2001). The samples were then viewed under a scanning electron microscope.

Results and discussion: The extract from the variety QE showed a high content of total polyphenols. The inhibition percent of H2O2 is 96.32 ± 0.56 %. The inhibition of *Candida albicans* biofilm formation on hydroxyapatite test showed strong inhibitory action against the strains in this order CA1 > CA3 > CA2.

Conclusion: This study reports the high content of phenolic compounds of pomegranate peel extract, antioxidiant activity and *Candida albicans* biofilm inhibition.
Background: Nowadays, mortality and morbidity due to infections caused by biofilm on medical devices such as catheter and implants are increasing and Candida has become one of the most common causes of nosocomial infections. One of the most important characteristics of biofilms is broad-spectrum resistance to antimicrobial drugs. However, too much using of this medicine can lead to the resistance of Candida species to fluconazole. Method: A total of 60 samples of medical devices attached to patients in ICU were investigated. Yeast colonies isolated from medical devices using routine laboratory procedures were identified. The MIC of fluconazole for all samples was carried out using Broth Microdilution according to CLSI. To assess biofilm formation by MTT, the experiment was carried out in a 96-cell microplate. Over half of the isolates of Candida had the ability to form biofilms, and the ability of biofilm formation was read by ELISA reader at a wavelength of 570 nm. Results: Out of 60 samples, 48 isolates of Candida with an abundance of Candida albicans (42%), C. glabrata (27%), C. kruise (17%) and C. tropicalis (14%) were identified. Among the isolates, 9 were susceptible to fluconazole (19%), 10 cases were dose dependent (21%) and 29 were resistant to fluconazole (60%). Candida isolates had the ability to form biofilms. C. albicans and C. tropicalis had the maximum and minimum power of biofilm formation respectively. Conclusions: The results showed that Candida was resistant to fluconazole. All resistant isolates had the ability to form biofilms.
MICROFLUIDIC MODEL FOR THE INVESTIGATION OF PSEUDOMONAS AERUGINOSA BIOFILMS

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Background

Pseudomonas aeruginosa is an opportunistic pathogen due to its capacity to form recalcitrant biofilm structures, while cohabiting with other harmful/pathogenic species and harboring the capability to release toxins that cause tissue necrosis. P. aeruginosa has been implicated in a number of microbially-induced or aggravated diseases, including cystic fibrosis and wound infections (e.g. burns, bites). Very little is known about the complex interactions that occur within polymicrobial communities and few tools exist for studying these interactions.

Objectives

In this study, we report on the development of a microfluidic model that mimics the relevant physiological properties of the wound microenvironment while incorporating materials present in the human extracellular matrix/wound environment.

Methods

We have validated the robustness of our model comparing traditional GFP-tagging to new fluorescent staining techniques to visualize/resolve individual species within a polymicrobial habitat. We have also demonstrated that chemotactic stimuli may be incorporated into our model through specialized ports in our chamber; we can readily monitor changes in motility in response to the introduction of amino acids, such as arginine and glutamine.
Conclusions

We show that our model can be used to investigate the spatio-temporal mechanobiological structures of the wound environment, which will significantly contribute to our understanding of the development and progression of polymicrobial biofilm infections and its interactions with the host environment. Through the implementation of microfluidic platforms, we pose complex and relevant questions that may enhance our understanding of microbial interactions, all with the goal of
identifying new and specific targets for more effective antimicrobial therapies.
IDENTIFICATION OF BIOFILM-RELATED GENES OF LISTERIA MONOCYTOGENES ISOLATED FROM SEAFOOD PROCESSING PREMISES

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Background

Listeria monocytogenes is a foodborne human pathogen which is commonly found in the environment and is capable of withstanding environmental threats by adhering to surfaces to form a biofilm. Biofilm formation is believed to be a major factor for cross-contamination in food processing premises and therefore, elucidating the molecular mechanism of biofilm formation in L. monocytogenes can contribute to secure food safety.

Objectives

To identify genes involved in biofilm formation of L. monocytogenes 15G01

Methods

A library of 6,500 mutants was constructed via transposon mutagenesis of L. monocytogenes 15G01, a persistent strain in New Zealand seafood processing premises, with the Himar1 mariner-based transposon. This library was screened for altered biofilm formation using a microtiter plate assay, using conditions defined as optimum through preliminary studies. Quantification of biofilm mass was undertaken by optical density measurement after staining with crystal violet. Mutants of interest were selected through statistical analysis (two-sample t-test, p≤0.05). The insertion site of the transposon was identified through semi-arbitrary PCR followed by DNA sequencing or by genome sequencing, and the inactivated loci analysed using BLAST.

Conclusions
Sequencing results revealed genes previously known to be involved in biofilm formation by *Listeria monocytogenes* and/or other bacterial species, which corroborates the method of transposon mutagenesis for functional analysis of biofilm formation. Furthermore, in this study, new genes influencing biofilm formation (either positively or negatively) have been identified. Further functional analysis of these genes is ongoing to elucidate their roles in biofilm formation.
Background

The opportunistic human pathogen *Pseudomonas aeruginosa* is able to adapt to a variety of often harmful environmental conditions due to different survival strategies including the formation of resistant biofilms.

Objectives

In this study, we investigated the adaptation of *P. aeruginosa* towards hypochlorite (HOCl), a strong oxidant used by human neutrophils to kill invading bacteria and which can be found in chronically inflamed host tissue, e.g. in the lungs of CF patients.

Methods

In static biofilm assays, we observed a significant increase of up to 3-fold in initial cell attachment in the presence of sub-lethal hypochlorite concentrations. Transcriptome analyses revealed a substantial upregulation of genes involved in oxidative stress response as well as biofilm formation in the presence of hypochlorite. Among others, we identified a 26-fold upregulation of ORF PA3177 coding for a putative diguanylate-cyclase (DGC), which catalyzes the synthesis of the second messenger c-di-GMP, and therefore influences motility, biofilm formation and persistence in *P. aeruginosa*. Subsequent LC-MS/MS analyses revealed a strong increase in c-di-GMP levels suggesting a key role of this second messenger in hypochlorite induced adhesion and biofilm development. The DGC PA3177 was further characterized in more detail demonstrating its involvement in motility, biofilm formation, antimicrobial resistance and persistence in *P. aeruginosa*. Using a subset of different mutant strains, we were able to show that both the pel and psl exopolysaccharides are effectors in the PA3177-dependent c-di-GMP network.
Conclusions
Our results demonstrate that host-derived antimicrobials are sensed by invading bacteria and exert a huge impact on bacterial pathogenesis including biofilm formation.
DISTINCT SAGA FROM HOSPITAL-ASSOCIATED CLADE A1 ENTEROCOCCUS FAECIUM STRAINS CONTRIBUTES TO BIOFILM FORMATION

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Background

Enterococcus faecium is an important nosocomial pathogen causing biofilm-mediated infections. Elucidating E. faecium biofilm pathogenesis is pivotal for development of new strategies to treat these infections. In several bacteria extracellular DNA (eDNA) and proteins act as matrix components contributing to biofilm development.

Objectives

In this study, we investigated biofilm formation capacity and the role of eDNA and secreted proteins in 83 E. faecium strains with different phylogenetic origin that clustered in clade A1 and clade B.

Methods

Although there was no significant difference in biofilm formation between E. faecium strains from both clades, addition of DNase I or proteinase K to biofilms demonstrated that eDNA is essential for biofilm formation in most E. faecium strains, while proteolysis primarily impacted on biofilm formation of hospital-associated E. faecium strains that cluster in clade A1. Secreted antigen A (SagA) was the most abundant protein in biofilms from clade A1 and B E. faecium strains, although localization differed between the two groups.

Conclusions

sagA is present in all sequenced E. faecium strains, with a consistent difference in the repeat region between the clades, which correlated with proteinase K susceptibility in biofilms. This indicates an association between the SagA repeat profile and the localization and contribution of SagA in E. faecium biofilms.
WCAJ DELETION ENHANCES BIOFILM FORMATION IN KLEBSIELLA PNEUMONIAE

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Background

The worldwide occurrence and recent rapid spread of multiresistant strains of Klebsiella pneumoniae is of great concern and has generated an urgent need to identify new drug targets. Bacterial adhesion and biofilm formation are important for the establishment of K. pneumoniae infections. Elucidation of mechanism of biofilm formation by pathogens is the first step towards the development of novel approaches for the treatment and prevention of biofilm related infections.

Objectives

The goal of this study was to gain additional insights into biofilm formation in K. pneumoniae MTCC 3384 by using mini-Tn5 transposon mutagenesis.
Disruption of Colanic acid synthesis in K. pneumoniae enhances biofilm formation

A

Specific Biofilm Formation

Wild Type
Kp3384 pDK6T
Kp3384 ΔwcaJ pDK6T
Kp3384 ΔwcaJ pDK6T-wcaJ

B

a) LB agar (without salt)

b) Colony morphologies of the strains

c) Calcofluor indicator plates

d) Congo Red indicator plates

Quantitative analysis of biofilm formation by Crystal violet (CV) staining
Methods

In this study, mini-Tn5 transposon insertion mutants were constructed and screened for their ability to form biofilms on abiotic surfaces by crystal violet staining. It was observed that disruption of wcaJ conferred high biofilm forming ability as compared to the parent strain. WcaJ is the initiating enzyme of colanic acid synthesis which loads the first sugar (glucose-1-P) on the lipid carrier undecaprenyl diphosphate. Its absence rendered the mutant defective in colanic acid, resulting in non-mucoid colonies. From the Calcofluor (binds cellulose) binding studies, it was evident that ΔwcaJ mutant does not produce much exopolysaccharide, thereby facilitating cell-cell interactions leading to the formation of closely packed structures. In addition, complementation of wcaJ in trans fully restored the wild type phenotype that confirms the observed effect of wcaJ on biofilm formation of K. pneumoniae.

Conclusions
Colanic acid expression in *K. pneumoniae* inhibits its cellular attachment to the substratum and biofilm formation, contradicting earlier studies that polysaccharides enhance biofilms in related species.
SALT STRESS INDUCES BIOFILM FORMATION IN CLOSTRIDIUM LJUNGDALII.

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Background

Clostridium ljungdahlii is a homoacetogen capable of producing organic commodities from carbon dioxide using electrical energy provided by an electrode, i.e. microbial electrosynthesis. The efficiency of this process likely depends on the biofilm formation on the electrode. So far, however, only limited attachment to electrodes has been obtained with C. ljungdahlii, while biofilm formation by C. ljungdahlii has not yet been reported.

Objectives

The presented work investigated whether adapted growth conditions can induce C. ljungdahlii biofilms and characterized the obtained attachment.

Methods

C. ljungdahlii was grown in 6-well plates using a tryptone containing medium. Different stress conditions (salt, antibiotic, temperature) were tested and biofilm formation was quantified using a crystal violet assay. The addition of NaCl (200 mM) most significantly induced biofilm formation, i.e. after 3 days the attachment was two order of magnitudes higher than in the control. Confocal laser scanning microscopy showed viable, 45 ± 5 µm thick biofilms, while without the addition of salt only a single cell layer (5 µm) was observed. Furthermore, a 15 ± 5 µm thick biofilm was obtained on a vertically placed piece of graphite, simulating an electrode, by adding NaCl to the medium. RNA sequence data of C. ljungdahlii grown with and without the NaCl addition are currently being analyzed to explore the genes involved in biofilm formation.

Conclusions

This work is the first to demonstrate biofilms in C. ljungdahlii and suggests that salt-induced biofilms of C. ljungdahlii should be tested for their microbial electrosynthesis potential.
CHARACTERIZATION OF XANA MUTANT OF THE XANTHOMONAS CITRI SUBSP. CITRI

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Background

*Xanthomonas citri* subsp. *citri* is the causal agent of citrus canker disease. This bacteria is able to form biofilm in vitro and on leave surface. The ability to form biofilm is important to citrus canker development, because it implicates to survival of bacteria on leaf surface of host plants, when it is sprayed on the plant. Cells in biofilm are embedded in a matrix composed for extracellular polysaccharide (EPS) and other proteins. In *Xanthomonas campestris*, the gene xanA is required for the synthesis of the exopolysaccharide xanthan, but participation of xanA in biofilm formation process was not evaluated in *Xanthomonas citri*.

Objectives

To characterize the function of xanA (XAC3579) in relation to adhesion, biofilm formation, motility, and epiphytic behavior of *X. citri* on leaves.

Methods

We realized assays for analysis of biofilm formation and adhesion in abiotic and biotic surface, EPS quantification, sliding and swimming motility to compare the mutant XAC3579 tn5 and *Xanthomonas citri* wild type.

Conclusions

The reduction of xanthan produced reduced the adherence and also the swimming and sliding motility. The presence of the xanthan was required for mature biofilm development on sweet orange leaf surfaces. This reduction can decrease the Xcc pathogenicity.
PROTEOMICS ANALYSIS IDENTIFIES PROTEINS ASSOCIATED WITH CURCUMIN-TREATED HELICOBACTER PYLORI BIOFILM

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Background
Helicobacter pylori is an important human pathogen, which is implicated as etiologic agent of peptic ulcer, chronic gastritis and gastric cancer. Recently, H. pylori is shown to have ability to form biofilm both in vitro and in vivo. Biofilm formation enables the bacterium to survive in adverse environments, to persist in patients causing chronic infection and confers enhanced resistance to antibiotics. Curcumin (diferuloylmethane), a natural compound found in turmeric, has been demonstrated for broad chemotherapeutic properties. We have previously reported that curcumin inhibits biofilm formation of H. pylori but its impact at a molecular level is unclear.

Objectives
This study was established to identify proteins involved in anti-biofilm activity of curcumin in H. pylori biofilm using proteomics analysis.

Methods
Biofilm of H. pylori ATCC43504 was established by pellicle assay. Two-dimentional gel electrophoresis (2-DIGE) and matrix-assisted laser desorption/ionization-time-of-flight (MALDI-TOF) were performed in order to investigate the difference in protein profile expression between curcumin-treated and -untreated H. pylori biofilms.

Conclusions
There were total 28 expressed proteins identified here. Eighteen proteins were regulated in both curcumin-treated and -untreated H. pylori biofilms. Proteins with decreased expression following exposure to 1/4 MIC of curcumin were associated with carbohydrate and nitrogen metabolisms, chemotaxis and motility, and electron transport. However, several chaperon proteins were up-regulated in response to a presence of curcumin. These data reveal a probable mechanism of curcumin that inhibit biofilm formation of H. pylori. Further studies need to be investigated in order to develop curcumin as a potential alternative medicine for treatment of H. pylori infections.
NOVEL METHODS FOR THE ASSESSMENT OF MARINE ANTI-BIOFILM TECHNOLOGIES.
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Background

Marine biofilms, mainly comprised of bacteria and diatoms, are now recognized to be a significant issue for a wide range of engineered structures such as ship hulls, energy systems, sensors, and oil/gas pipe lines. Biofilms constitute a major component of the overall marine growth on ship hulls and may lead to a 14 % increase in ship fuel costs and to an up to 18 % fuel consumption penalties. The scientific community is now placing significant effort towards the creation of environmentally benign surfaces with anti-biofilm capacity. However, there is significant gap in screening assays that test the bacterial/surface interface during initial attachment for anti-biofilm technologies while dynamic testing is often overlooked.

Objectives

In this work a range of bioassays have been developed to directly test bacterial attachment and biofilm growth on experimental anti-biofilm
surfaces, under both static and hydrodynamic conditions.

Methods

Following screening of several compounds for anti-biofilm efficacy, juglone (MIC at 5ppm) was chosen to act as model inhibitory agent, which was then successfully incorporated into a coating system and illustrated good anti-biofilm capacity. Novel microfluidic devices were developed for the assessment of bacterial attachments and biofilm formation in the presence and absence of juglone under hydrodynamic conditions. Juglone inhibited biofilm formation and following nucleic acid staining, toxicity against *Cobetia marina* was evident. This was not clear during the static assays.

Conclusions

The developed bioassays illustrated promising reproducibility and greater relevance for anti-biofilm technologies. New microfluidic approaches have been employed to reveal differences in results when comparing static vs. dynamic testing.
Background. Biofilm formation is the preferred growth mode and survival strategy of several bacteria for improving adaptation and viability under hostile conditions. For mucosal and health-promoting bacteria the biofilm formation is likely to play an important role in colonization and immunostimulation. We have previously shown that the probiotic paradigm Lactobacillus rhamnosus GG displays a high number of adhesive and immunostimulatory moonlighting proteins at the cell surface, and that surface-bound proteins are important mediators of biofilm formation of this strain.

Objectives. The aim of this study was to explore the effect of fourteen different fermentable carbohydrates on the biofilm formation of L. rhamnosus GG, and to uncover their impact on the surface protein/antigen composition and the adhesive properties of the strain during biofilm and planktonic growth.

Methods. Biofilm formation was assessed using the standardized polystyrene microtiter plate assay, and the viability and structure of selected biofilms were evaluated with LIVE/DEAD staining coupled with confocal microscopy analyses. Surface-associated proteins and antigens were identified and analyzed using in-house proteome and immunoproteome methods. Selected biofilm and planktonic cultures were further tested for their adhesion activities in vitro.

Conclusions. This study indicated that carbohydrate source plays a significant role in the protein-mediated biofilm formation and the antigenicity of the biofilm cells. We also show that moonlighting proteins are the major protein components of the biofilm matrix, and that certain carbohydrates affected the adhesion properties of biofilm cells.
Background

Biofilms are communities of microorganisms which live in a self-produced matrix of extracellular polymeric substances. Their infections are hard to treat due to their resistance to immune defense and antibiotics. Therefore it is important to develop tools to investigate biofilms for therapy improvement or to find new antimicrobial substances.

Objectives

The aim of this work is to establish a method for the fully automated large-scale screening of biofilms which could be used for examining the ability of biofilm formation of bacteria under different culture conditions and most importantly for determining the effects of different antibiotics and antimicrobial substances.

Methods

We enhanced our previous published VideoScan technology, which is based on fully automated fluorescence microscopy to perform high-throughput screening of biofilms. VideoScan enables the analysis of multiplex assays such as microbead or cell-based assays [Rödiger et al. 2013; Frömmel et al. 2013]. We use a 96 well plate format for the formation of biofilms, which are stained with Live/Dead staining followed by a VideoScan analysis. This analysis represents a two-step evaluation realized by our software and image processing. In the first step overview pictures of biofilms are taken and in a second step, a fine-grained analysis of the biofilm, bacteria are counted in different z-stacks of the biofilm.

Conclusions

With our VideoScan technology it is possible to study biofilms in a fully automated large-scale screening. In a next step a large panel of commensal and pathogenic E. coli strains will be screened for biofilm formation under different growth conditions and presence of biofilm-associated genes.
Background

The development of alternative strategy to conventional antibiotics in the struggle against pathogens and antibiotic resistance is a topical issue. The interference with the adhesion, the first stage of pathogenic process, can modulate virulence mechanisms like colonization, invasion of host tissues and biofilm formation. The curcumin can inhibit the adhesion of both Gram positive and Gram negative pathogenic strains and different mechanisms can be involved (inhibition of sortase A, Quorum-sensing interference, binding to curli, etc.).

Objectives

To obtain novel agents against pathogenic bacteria, targeting virulence mechanisms and biofilm formation (anti-virulence agents).

Methods

In this study, we synthesized a group of molecules derivatives of curcumin, and the inhibition of biofilm formation was evaluated at a screening concentration of 7.5 µg/ml against reference staphylococcal strains S.aureus ATCC 29213, 25923, 6238, S.epidermidis RP62A, and Gram negative reference strains E.coli ATCC 10536 and P.aeruginosa ATCC 15442. The activity as sortase A inhibitors was also screened in a high throughput assay by using the standard Dabcyl-QALPETGEE-Edans fluorescence resonance energy transfer (FRET)-peptide substrate for measurement of enzyme activity.

Conclusions

Some curcumin derived compounds resulted effective at lower concentration to prevent biofilm formation than the curcumin. The discovery of novel agents that target virulence mechanisms and biofilm formation offers new potential therapeutic strategies to treat chronic bacterial infections. They can act either alone or in combination with current antibiotics and could significantly impact on overcoming the problem of antibiotic resistance, which is recognized by WHO as one of the most important global challenges of our time.
UNRAVELING EROSION DYNAMICS: MICROBIAL STABILIZATION OF FINE SEDIMENTS

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Background
During the formation of aquatic biofilms microbes produce extracellular polymeric substances (EPSs) inter alia to attach to the sediment. Thus, sediment grains can be glued together and thereby are granted a significantly increased resistance versus erosive forces - a phenomenon called biostabilization.

Objectives
This process constitutes an essential ecosystem function (e.g. detainment of contaminated fine sediments in rivers) but is currently barely addresses in lotic waters. In our current DFG project engineers and biologists work together to elucidate the fundamental principles of biostabilization.

Methods
Straight flume mesocosm experiments were run at different seasons and under varied boundary conditions to investigate the effect of biotic and abiotic factors upon biofilm development and resulting biostabilization. Besides biofilms’ stabilization capacity (adhesiveness and erosion resistance) biochemical and biological features (e.g. microbial biomass, EPS production and microbial community composition) were investigated.

Conclusions
A general temporal increasing stabilizing effect could be detected - one of the first scientific proofs for this process in lotic waters. Furthermore, a strong effect of season and environment upon biofilm development and stabilizing capacity was detected, e.g. a significant maximum during spring but a minimum under minimal illumination. Besides the possible role of EPS protein content as proxy for biofilm stability the importance of successional processes were revealed: especially mutual adaption of algae and bacteria (e.g. expressed by the development of dynamics or functional organization within the bacterial community) is apparently a driving factor.
Background

Recent studies show that extracellular DNA is a major component of the EPS (extracellular polymeric substances) in biofilms and it was found to have widespread importance in biofilm formation. However, detailed knowledge about how it promotes biofilm formation and how it influences biofilm morphology is just starting to emerge (1).

The γ-Proteobacterium F8 was isolated from the South Saskatchewan River in Canada. It was found to produce huge amounts of extracellular DNA forming a filamentous network when grown on agar plates containing FBM medium (2, 3).

Objectives

The aim of this study was to examine this phenomenon for the first time under continuous flow conditions in a biofilm reactor.

Methods

F8 was grown under controlled conditions in a continuous flow biofilm reactor and eDNA-production as well as biofilm morphology was monitored over time. Cells and EPS components were visualized by differential staining with various fluorescent dyes and confocal laser scanning microscopy (CLSM).

Conclusions

F8 cells attached to glass slides and showed biofilm formation from day 1 on. Filamentous eDNA could be detected by staining with Syto 9 and Propidium Iodide, but not with DAPI. Results so far indicate that eDNA is emerging from living cells.
Background
In burn patients, *Pseudomonas aeruginosa* can lead to a great variety of systemic infections with higher mortality rate. In addition, treatment of *P. aeruginosa* infections is a serious medical challenge because these bacteria are associated to high resistance to antibiotics and ability to form biofilms, becoming a serious problem in burn wound center (BWC).

Objectives
The aim of this study was to investigate the antibiotic resistance and biofilm formation from 35 *P. aeruginosa* collected from wound burn patients and on the surface of the bath tank where balneotherapy was performed.

Methods
The antibiotic resistance was carried out by disk diffusion method, following CLSI guidelines and the evaluation of biofilm formation was achieved through microtiter plate assay incubated for 24h. Genotypic characteristics of these strains were determined by Pulsed-field gel electrophoresis (PFGE).

Conclusions
It was observed that the clone A, the most prevalent and presents in wound patients and on the surface of the bath tank, had a high capacity to form biofilm. These strains also showed resistance to drugs as ceftazidime, aztreonam, gentamicin, ciprofloxacin and even meropenem. We believed that this clone which is more able to accumulate drug resistance mechanisms and to form biofilm has greater conditions to adapt and to persist in the BWC.
MICRO-DIVERSITY OF ANAMMOX BACTERIA IN NITRITATION-ANAMMOX BIOFILMS

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Background
An emerging technology for nitrogen removal in wastewater is the Anammox process. However, questions about the stability of the process remain. It is generally believed that a higher diversity can increase ecosystem stability. Coexistence of two anammox species in a Moving Bed Biofilm Reactor (MBBR) was observed in a previous study (Persson et al. 2014).

Objectives
We studied if coexistence of different anammox populations is common in nitritation-anammox MBBRs.

Methods
Anammox micro-diversity was assessed with automated ribosomal intergenic spacer analysis (ARISA) using a nested PCR approach, and fluorescence in situ hybridization (FISH). Four pilot MBBRs at two different pilot wastewater treatment plants in Sweden with different running conditions and one full-scale integrated fixed-film activated sludge reactor (IFAS) were studied.
Conclusions
Preliminary ARISA results suggest that nitritation-anammox MBBRs usually harbor more than one anammox population and FISH from two MBBRs confirmed this (Figure 1). Possible explanations include presence of biofilm microenvironments and temporal variations in environmental conditions such as influent composition. However, one MBBR operated at the challenging conditions of low nitrogen concentrations and at low temperature (13°C) was dominated by a single species.

References
FUNCTIONALIZATION OF PDMS WITH CELLOBIOSE DEHYDROGENASE YIELDS ANTIBIOFILM SURFACE
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Background

Patients with medical indwelling devices (e.g. urinary catheters) are prone to acquiring infections caused by biofilm forming pathogens colonizing catheter surfaces. Catheter associated urinary tract infections (CAUTI) are among the most frequent ones. One of the polymers widely used for making urinary catheters is polydimethylsiloxane (PDMS) due to its favorable biocompatibility properties.

Objectives

In order to decrease the risk of infection, an antimicrobial enzyme, namely cellobiose dehydrogenase (CDH) was successfully grafted onto PDMS surface.

Methods

The system is based on the ability of CDH to use oxygen as electron acceptor and different oligosaccharides (e.g. cellobiose) as electron donors to produce H₂O₂. Several approaches of immobilizing CDH on PDMS surface were exploited including surface activation using oxygen plasma followed by covalent linkage of CDH. The success of the immobilization process was monitored by analyzing the change in the functional groups on the surfaces by FTIR measurements as well as measuring the ability of grafted CDH to produce H₂O₂. Antimicrobial activity of the immobilized enzyme was assessed against S. aureus by quantifying the amount of viable cells and total biomass formed on the enzyme treated sheets compared to the control.

Conclusions

CDH was successfully immobilized on the surface of PDMS as evidenced by H₂O₂ production in the presence of cellobiose. The modified surface showed up to 70% reduction of microbial growth as compared to the untreated control. The CDH modified PDMS catheters could help to prevent current problems of microbial colonization and multidrug resistant bacteria associated with catheters.
ENHANCED FLOC FORMING MUTANTS OF RHODOVULUM SULFIDOPHILUM OBTAINED BY UV MUTAGENESIS

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Background
A marine photosynthetic bacterium, Rhodovulum sulfidophilum, produces nucleic acids extracellularly¹. Interestingly, the nucleic acids thus produced outside cells are utilized as a sticky glue to make bacterial self-aggregates called “floc” in nutritionally poor medium conditions such as on the sea bottom where the bacterium was originally isolated. However, what gene regulates its flocculation or why this bacterium makes a floc under nutritionally stringent conditions is still unclear.

Objectives
Our objectives are to establish UV mutagenesis in Rhodovulum sulfidophilum and to obtain UV mutagenic clones showing aberrant flocculation behavior even in rich medium conditions where the bacterium usually doesn’t make an obvious floc.

Methods

Rhodovulum sulfidophilum DSM 1374T was exposed to UVC (254 nm) irradiation and UV mutagenesis was induced. After UVC irradiation, rifampicin-resistant mutants were collected and these flocculation behaviors were examined.

Conclusions
UV mutagenesis in Rhodovulum sulfidophilum DSM 1374T was established and 107 rifampicin-resistant mutants were obtained. Two out of the obtained clones showed strong flocculating behavior even in nutritionally rich medium conditions and these flocculation ratios were approximately 40% and 60%, respectively, whereas that of wild type was ca. 20%.
Background

*S. aureus* and *S. epidermidis* are two important nosocomial pathogens which can interact in biofilms (e.g. *S. epidermidis* inhibits *S. aureus* biofilm formation by producing proteases).

Objectives

To study the mutual effect of *S. epidermidis* and *S. aureus* on gene expression in dual species biofilms.

Methods

RNA-Seq was performed on single and dual species biofilms and the results were confirmed by the appropriate phenotypic tests.

Conclusions

In dual species biofilms, *S. epidermidis* genes encoding resistance to erythromycin, oxacillin and tobramycin were upregulated and when biofilms were treated with these antibiotics, we observed that more *S. epidermidis* cells survived oxacillin treatment in dual species biofilms than in single species biofilms.

Urease genes of *S. aureus* were downregulated in dual species biofilms and an urease assay confirmed that there was more urease activity in *S. aureus* single species biofilms compared to dual species biofilms. Literature data described a link between urease activity and pH and we also observed that the pH of *S. aureus* single species biofilms was lower than the pH of dual species and *S. epidermidis* single species biofilms. Metabolic activity results in acid production and RNA-Seq data showed there was less expression, in dual species biofilms, of *S. aureus* genes encoding proteins involved in metabolic processes. Viability assays showed that there were more metabolically active cells in *S. aureus* biofilms compared to dual species and *S. epidermidis* single species biofilms. Altogether, our results show that *S. epidermidis* has an inhibitory effect on the metabolic activity of *S. aureus* in dual species biofilms.
Background

Bacterial cells in biofilm are more resistant to wide range of disinfectants than free living cells, and finding an appropriate agent against them is demanding task.

Environment is certainly the source of many products which have anti-biofilm effects, but they have not been proven, yet.

Objectives
To investigate the anti-biofilm effect of 100% pure red raspberry fruit juice on Pseudomonas aeruginosa biofilm formation.

Methods
Minimal inhibitory concentration of red raspberry fruit juice was evaluated by dilution method, and its anti-biofilm effect was tested in microtiter plate using crystal violet staining according to modified technique by Stepanovic.

Conclusions
Out of 30 tested strong-adherent Pseudomonas aeruginosa isolates, after treating with red raspberry fruit juice, 19 (73.3%) isolates were poorly adherent, while 8 (26.7%) had no adherence ability.

This study confirmed the great potential of red raspberry fruit juice as anti-biofilm agent for preventing microbial colonization, but it is necessary to discover the exact biochemically active ingredient against biofilm formation.
Background

Many of the world’s most precious artworks are made of stone. Their irreversible deterioration due to biological attack is a worldwide concern. Microorganisms colonize outdoor lithic surfaces and develop into biofilms at the interface solid/air (subaerial biofilms, SABs), which, in turn might cause aesthetic, chemical and physical decay. Although it has been estimated that at least 99% of the world’s microbial biomass exists in biofilms, the role and behavior of microorganisms within the biofilm matrix and their complex interactions with the external environment is still unknown.

Objectives

This work provides a pioneering and multidisciplinary research to investigate the behavior of microorganisms within the biofilm matrix for sorting out time-spatial relationships and to elucidate microorganism-EPS, inter-organism, biofilm-atmosphere and biofilm-stone interactions.

Methods

This work spans sophisticated molecular, chemical, physical and data modeling techniques and it is approached from two complementary angles:
1- Lab-scale study to delineate specific transcriptional responses of mono- and multi-species biofilms as well as the biofilm-stone interactions under controlled environmental conditions.
2- Real heritage case studies to investigate the shifts in the microbial community structure and function under different environmental conditions. Through comparing phylogenetic and functional diversity under different environmental scenarios, we
provide evidence that any intuition gained from the lab-scale experiments is relevant to true environmental biofilms.

Conclusions

The findings obtained so far will contribute to better understand the complexity of all the interactions encountered within SAB communities, and how these interactions may influence the biofilm outcome and the biodeterioration of the stone materials under different environmental conditions.
Background

Bacterial collagen-like proteins (CLPs) have been identified in a broad range of pathogen bacteria and it is important to biofilm formation and bacterial adhesion to host cells. Some bacterial CLP-encoding genes (clps) have also been found in non-human pathogenic strains such as B. cereus and B. amyloliquefaciens, which are types of plant-growth promoting rhizobacteria (PGPR).

Objectives

To elucidate the role of CLPs playing in the interaction between PGPR with plant host, the biological functions and cellular location have to explore.

Methods

CLP-encoding genes ClpA, ClpB, ClpC, and ClpD in strain FZB42, were inactivated separately by Site-Directed Mutagenesis. Scanning electron microscopy (SEM) and hydrophobicity value detection were used to assess the morphologic bacterial cell shape and cell surface architecture. Immune gold labeling was used to indicate the location of CLPs on the cells. The bacterial flagella morphology observed by transmission electron microscopy (TEM).

Conclusions

Comparing with wild type strain, clps mutant strains showed differently on phenotypes of bacterial colony shape, cell autoaggregation, biofilm formation, as well as adhesion to surface of abiotic materials or the roots of Arabidopsis thaliana. Immune gold labeling shown CLPs located in the outer layer of the bacterial cell, including the cell wall, outer membrane, flagella, or other associated structures. In addition, the bacterial flagella appeared fewer, scattered and bent for clps mutant strains. Above
results indicate that the bacterial out layer located CLPs are closely involved in cell shape development, biofilm formation, and bacteria-plant interaction.
SYSTEMS MICROBIOLOGY OF ALGINATE-LIKE EXOPOLYSACCHARIDES
BIOSYNTHESIS IN GRANULAR SLUDGE BIOFILMS: A PILOT AND FULL-SCALE
STUDY WITH DOMESTIC WASTEWATER

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Background

Environmental biotechnology systems using aerobic granular sludge (AGS) are
becoming new standards for intensified wastewater treatment. AGS originates from
self-aggregation of activated sludge microorganisms into mobile biofilm structures
under operational selection pressures. Extracellular polymeric substances (EPS)
such as alginate-like exopolysaccharides (ALE) play an important structural role by
supporting microbial encapsulation in a gel matrix.

Objectives

The complex microbiome of AGS was studied using a systems biology approach.
Investigations targeted bacterial phylotypes carrying genetic and metabolic potential
for ALE biosynthesis. Such correlation analysis will enable to elucidate molecular
mechanisms underlying granular biofilm formation.

Methods

A 98-L pilot sequencing-batch reactor was operated for granulation and nitrogen
removal from domestic wastewater. Evolution of particle size distribution and EPS
polysaccharide and protein fractions were determined by laser diffraction, phenol-
sulfuric acid, and folin-phenol methods, respectively. AGS from full-scale system was
subjected to physicochemical characterization of ALE. Bacterial populations and
genetic signatures related to ALE biosynthesis are currently prospected from pilot
and full scales using phylogenetic and metagenomic profiling.

Conclusions
Over 3 months of operation granulation displayed gradual increase in the fraction and mean diameter of granular biofilms from 5 to 60% and 50 to 300 µm, respectively. The polysaccharide-to-protein ratio increased concomitantly from 0.4 (m/m) in activated sludge flocs to 0.8 and 1.0 in nitrifying and denitrifying granules, respectively. Lyophilized granules from full-scale plant contained 15% of ALE, i.e. twice higher than typically measured in flocs. Correlating these physicochemical gradients to phylogenetic structure and genomic signatures will allow for fundamental understanding of the granulation phenomenon and its temporal triggering.
Background
Biofilms pose a way to overcome toxic influences like antibiotics and metal ions. As the number of multiresistant bacteria increases (e.g. *Staphylococcus aureus*) steadily, metals (copper, silver) celebrate their renaissance/revival for many purposes in healthcare facilities. Not much is known about the tolerance against these heavy metals.

Objectives
The known positive effect of the heavy metal tolerance of formed biofilms implicates that there could be a regulatory effect of copper ions on the formation and maturation of biofilms. Previous studies on this matter present inconclusive data, pointing in both directions.

Methods
The influence of different CuCl$_2$ and CuSO$_4$ concentrations (0.5 – 5 mM) on the formation of biofilms of two *S. aureus* strains as well as on already formed biofilms was tested in a multi well plate assay. Protein and polysaccharide concentrations were quantified and biofilm growth was measured after 48h and 96h.

Conclusions
If grown in copper containing solutions the both strains used showed a different behavior. One strain was inhibited at higher copper concentrations (> 2.5 mM), the other strain presented an increased EPS production at the same level. Both, CuCl$_2$ and CuSO$_4$, did not show an inhibitory effect on already formed biofilms of both strains. Interestingly lower concentrations (<3 mM) revealed a biofilm promoting effect. Further studies with more strains are needed to be certain which influence copper has on biofilms and their formation as apparently the strains show different behavior.
Background

Biofilm formation is important for rhizobia to colonize several environmental niches. Within the microcolonies developed by *Rhizobium leguminosarum*, rhizobial cells interact tightly through lateral and polar connections forming compact aggregates. These microcolonies are surrounded by a biofilm matrix, whose main component is the acidic exopolysaccharide (EPS). In addition, the PrsDE secretion system has been implicated in biofilm formation. This system is responsible for the secretion of Rap proteins that share one or two Ra/CHDL (cadherin-like-) domains.

Objectives

We aimed to explore the role of Rap(s) and other surface factors in biofilm development.

Methods

RapA was analyzed by biophysical and biochemical approaches. Functional studies were assessed by mutagenesis and gene overexpression.

Conclusions

We showed that RapA, which consists of two Ra/CHDL domains, is a unipolar calcium dependent lectin that specifically binds the EPS. RapA overexpression enhanced biofilm formation and expanded the distance between cells. In line with these observations, we showed that high levels of RapA secretion increases capsular polysaccharide (CPS) formation, which is structurally and genetically related with the EPS. We propose that RapA and the other Rap proteins, which are predicted to harbor one or two EPS/CPS-binding domains, play some role remodeling the biofilm matrix structure. Exploring the role of another important surface polysaccharide, the lipopolysaccharide (LPS), we found that the O-chain core region of the LPS is crucial for cell-cell cohesion. Mutants defective in the O-chain core moiety developed
biofilms with an altered three-dimensional structure. Further studies are required to give insight into the interplay between the biofilm matrix components and Rap(s).
Biofilms

BIOFILM ACCUMULATION AT DIFFERENT WATER TREATMENT STAGES
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Background
Biofilms are microbial communities and highly relevant for water quality. In principle biofilms may affect human health in drinking water reservoirs and distribution systems hindering the efficient operation of these systems. Also they may also pose a health risk due the presence of pathogenic bacteria such as \textit{Legionella pneumophila} and \textit{Escherichia coli}.

Objectives
The aim of this study was to describe biofilms accumulating within controlled pilot plant after different treatment steps. The use of various methodological approaches should describe biofilms and the associated water phases in order to assess potential factors associated with human health risks.

Methods
In total 6 biofilm samples were collected from the PE-X pipes of the pilot drinking water treatment plant. The plant is designed for the drinking water treatment of a small rural community located in Upper-Austria (193 inhabitants in 2014; average daily water consumption: 20 m\textsuperscript{3}). The treatment has 4 successive steps: 1) oxygen enrichment (using pressurized air) 2) biological nitrification, 3) rapid sand filtration and 4) UV-disinfection. Biofilms were characterized by chemical (TOC, DOC, heavy metals), microbiological (HPC, flow cytometry) and molecular biological (16S rRNA gene sequencing) and microscopic methodologies.

Conclusions
The multi-parametric approach applied in this study has shown valuable results with good correlations among chemical and microbiological characterization. The activity of biofilms (viability) could be described by flow cytometry and fluorescent microscopy. Phylogenetic analysis of sequenced culturable fractions revealed differing diversities in the microbial communities.
Biofilms are diverse microbial communities, bound by a matrix of extracellular polymeric substances, that allow cells to survive a variety of stressful, potentially lethal, conditions. To successfully avoid exposure to such toxic conditions, cells must transition rapidly from planktonic to sessile state, requiring quick acting regulatory mechanisms that can both inhibit motility and promote adhesion. We previously showed that the evolutionarily conserved type IV pili (TFP) are crucial for surface adhesion and microcolony formation in the model archaeon Haloferax volcanii. We also demonstrated that six conserved H. volcanii pilus subunits, the pilins PilA1-6, are involved in the regulation of flagella-dependent motility. Additionally, a subset of these pilins promotes microcolony formation, while at least two of the remaining pilins inhibit it. Recent studies suggest that pilin glycosylation catalyzed by the glycosyltransferase, AglB, plays an important role in the regulation of microcolony formation. Consistent with the pilins required for microcolony formation (PilA5 and PilA6) not being glycosylated, and the remaining pilins requiring glycosylation to function properly, we have shown that a ΔaglB strain forms microcolonies. Stress adversely affects AglB-dependent glycosylation, which is also required for flagella-dependent motility, hence inhibiting motility and promoting microcolony formation. Consistent with this we found that under low salt conditions H. volcanii is not motile and readily forms microcolonies. This differential post-translational modification that allows H. volcanii cells to respond quickly to changes in the environment that result in conditions favorable to sessile cells and biofilm formation rather than motile planktonic cells may be broadly conserved across prokaryotes.
The potential of halophilic microorganisms to convert waste glycerol to valuable compounds: glycerol carbonate (GlyC) and glycidol (GlyD)

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Background: Waste glycerol is the most important by-product in the production of biodiesel from biomass. As a direct consequence, waste glycerol (unpurified) was produced in quantities which soon will exceed current market demands leading to serious environmental problem by its storage.

Objectives: This work enlarges the application area for waste glycerol, proposing its usage as raw material in the production of glycerol derivatives (e.g. GlyC, GlyD and polyglycidol). Thus, alternative solutions for the storage problem of waste glycerol that can generate negative effects on the environment are developed. GlyC, GlyD and polyglycidol represent products of interest for fine chemical industry. Thus, GlyC is used as solvent for cosmetics and in medicine, while GlyD and polyglycerols appear as synthesis intermediates of plastics and resins in the polymer industry, but also in the pharmaceutical and cosmetic industries.

Methods: The products of interest were revealed in culture media using GC-MS and GC-FID techniques. Also, the polyglycidol products were evaluated (Mw, Mn and PI) based on GPC technique.

Conclusions: Preliminary results revealed ability of various moderately halophilic bacteria isolated from saline and hypersaline environments from Romania to convert waste glycerol to GlyC and GlyD. Lipase enzyme is responsible for GlyC production, while lipase and decarboxilase mixture convert the glycerol to GlyD. Polymerization of GlyD is performed under basic conditions.
Production of antimicrobial and anticancer substances by Bacillus pumilus SG-32 and Bacillus firmus P1-1 isolated from oil reservoirs in Brazil.

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BACKGROUND: Thirty strains isolated from Brazilian oil reservoirs were tested against each other for the production of antimicrobial substances. Two strains of Bacillus, B. pumilus SG-32 and B. firmus P1-1 were selected due to their ability to inhibit more than 60% of the strains tested. These strains were also investigated for cytotoxic activity against human cancer cell lineages.

METHODS: The screening of antimicrobial substances was performed by the agar diffusion method described for Rosado and Seldin (1993). Thirty strains previously isolated from oil reservoirs in Campos Basin (RJ, Brazil) were tested against each other for antimicrobial production. Two selected strains were growth in Marine Broth for 72 hours at 40°C to increase cell biomass. Anticancer screening was performed against nine different types of human cancer cell lineages.

RESULTS: Among the thirty strains tested, three of them (two strains of Bacillus pumilus, SG-30 and SG-32, and one strain of Bacillus firmus, P1-1) were able to inhibit more than 60% of the bacterial strains under study. Strains P1-1 and SG-32 were chosen for subsequent cytotoxic activity assay against human cancer cells. Strain P1-1 presented better effect than the positive control doxorubicin against colon and breast cell cancer lineages. Strain SG-32 was not as efficient as the positive control; nevertheless, five types of cancer cell lineage were inhibited.

CONCLUSIONS: Three different Bacillus strains from Brazilian oil reservoirs were able to produce antimicrobial substances, two of them with significant activity against cancer cell lineages, showing potential for further clinical trials.
SAFETY EVALUATION OF SOME COMMERCIAL SKIN COSMETICS USING A GENETICALLY MODIFIED ACINETOBACTER BIOLUMINESCENT BIOREPORTER DF4/PUTK2


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Background
Human skin is suffering from continued use of cosmetics, which may leads to ecosystem disruptions of useful skin microbiota. Therefore, a need to ensure its safety-in-use is important. Despite, microbiological assays employed to determine the lowest cosmetic concentration gave the lowest microbial contamination level during usage; toxicity assay must be performed first.

Objectives
In the present work, the Bioreporter DF4/PUTK2 was employed to assess the toxicity of eight local commercial cosmetic products. DF4/PUTK2 is a genetically modified strain belongs to the genus Acinetobacter that widely spread in nature and skin microbiota. Previously, DF4/PUTk2 was successfully employed to assess toxicity of phenolics and heavy metals.

Methods
Present results indicated that among the eight tested cosmetics, the Bioreporter was directly sensitive in order of decreasing sensitivity to aftershave balm, skin-refining toner, moisturizing cream and purifying cream with bioluminescence inhibition percent (BI%) equal to 64%, 58%, 50% and 30% respectively, Followed by eye makeup remover and sun cream with (BI%) equal to about 15% but after exposure time 300 and 700 min respectively. However, no decrease for bioluminescence with eye color powder and color trend – fresh foundation was observed. Also, the effect of exposure time on bioluminescence was tested. It was noticed that the cosmetics that expected to remain for a long time on skin seems most toxics.

Conclusions
In conclusion, the Bioreporter DF4/PUTK2 can be used to assay the toxicity of cosmetics and it is candidate to be a prescreening and quality control tool to pre-select the most toxic cosmetic for further chemical analysis.
INCREASING SOLUBLE EXPRESSION OF ALPHA-LUFFIN RIBOSOME INACTIVATING PROTEIN BY ENBASE FED-BATCH CULTURE

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Background

Considering the undeniable advantages of natural toxins from RIP family to produce anticancer immune-conjugates, well optimized recombinant expression of this category of proteins is of importance regarding industrial aspects. One of the proposed cellular mechanisms involved in EnBase fed-batch mode of cultivation is the change in protein expression via ribosomal manipulation. However, this system has not been applied to cytotoxic proteins.

Objectives

For the first time here, the expression profile of α-luffin, a ribosome inactivation protein (RIP) with an innate toxicity, was optimized in EnBase system. In fed-batch fermentation mode, the early incubation time was preferable at 30° C culture temperature whereas at 25° C the extended protein synthesis period (12 and 24h post induction) resulted in higher amount of soluble recombinant protein.
Methods

Mature α-luffin encoding cDNA was synthesized and subcloned into pET28a under the control of T7 promoter fused to the 6-HIS tag. The E. coli expression procedure was compared in traditional batch and newly developed fed batch; EnBase® Flo system at 25°C and 30°C incubation temperatures.

Conclusions
As the first study investigating the efficacy of EnBase fed-batch mode for the production of a toxic protein (α-luffin) it is concluded that in spite of the efficacy of this system in producing higher soluble protein ratios compared to batch cultivation growth rate, incubation temperature and time need to be optimized so as to cover innate cytotoxicity. The optimized condition proposed here is promising in terms of large scale soluble production of α-luffin without the need for refolding.
ANALYSIS OF TRANSCUTANEOUS DELIVERY OF TAT-BONT/A(1-448) FUSION PROTEIN AS THERAPEUTIC AGENT

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Background

Botulinum neurotoxin type A (BoNT/A) composed of three domain. One catalytic domain and one binding domain which are linked together by a translocation domain. In recent years, BoNT/A was used as a therapeutic agent for treatment some abnormal muscle contractions such as strabismus, blepharospasm, spasmodic or face wrinkles caused by ageing process. The injection of this drug causes some undesirable side effects such as irritation, pain, bleeding at the point of injection. Cell penetrating cationic peptides (CPPs) are three to 30 peptide residues which freely pass through cell membranes. Tat peptide is a CPP which is a favorite way to transduction of biological macromolecules (peptides, proteins or nucleotides) into cells.

Objectives

In this study, Tat(47-57) peptide directly fused to catalytic domain of BoNT/A. The penetration of this recombinant protein was analyzed in vitro.

Methods

A genetic construct containing Tat gene sequence, residues 47-57, was fused with that of BoNT/A (residues 1-448) and cloned into a bacterial host for further expression. Obtained fusion protein, was purified and transferred into a 6 well plate seeded by HeLa cell line and was incubated at 37°C for 1 to 2 h. Transduction of protein was further analyzed by western blot.

Conclusions

Results showed that Tat(47-57) peptide can carry the BoNT/A(1-448) into HeLa cell line in comparison to BoNT/A(1-448) as control. Cell transduction was time dependent (protein band in western blot after 2 h was thicker than 1 h). CPPs are promising tools for
transferring different type of drugs directly into intact skin for therapeutic purposes through a non-invasive way.
CAN S. CEREVISIAE PRODUCE BIO-BASED POLYMERS FROM PENTOSE SUGARS?

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Background

A sustainable future necessitates an urgent transition from petroleum-based industries towards the production of bio-based bulk chemicals from renewable feedstock [1]. The bio-based polymers, such as polyhydroxyalkanoates (PHAs), can be produced from monomeric sugars. Poly-3-D-hydroxybutyrate (PHB) is one of the most studied PHAs and is a promising biopolymer produced naturally by several bacterial species [2]. In parallel, xylose is the main pentose sugar present in lignocellulosic biomass (LCB), which is considered the cheapest carbon source on earth [3]. Thus, LCB is a potential substrate for the bio-based production of bulk chemicals such as PHB.

Objectives

In this work, the capacity of recombinant Saccharomyces cerevisiae to produce PHB using pentose sugars (xylose) as the main carbon source was evaluated.

Methods

Engineered S. cerevisiae strains, capable of pentose utilization were transformed with the PHB pathway genes from the natural and well-characterized PHB-producer Cupriavidus necator. The two host strains carried genes for xylose utilization from the yeast Scheffersomyces stipitis, but two variants of the xylose reductase (XR) with different NADH/NADPH preference ratio were tested [4]. The resulting strains were evaluated for their PHB-production capacity.

Conclusions

The production of PHB from xylose-rich medium was achieved using S. cerevisiae. The highest PHB yield (g/g xylose) was achieved with the strains carrying the mutated S. stipitis XR with a higher NADH/NADPH preference, highlighting the importance of redox balancing for PHB production.

USE OF FUNGAL ROTATING DISK REACTOR (RDR) FOR BIODEGRADATION OF TEXTILE ANTHRAQUINONE DYES

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Background

Biodegradation potential of fungi for removal of dyes from textile wastewater has been established and applied in aerated reactors.

Objectives

The aim was to compare dye-degradation efficiency of a newly isolated Trametes suaveolens strain with different fungal species in RDR. Reactive RB19 (C.I.61200) and disperse DB3 (C.I.61505) dyes were used.

Methods

2-L RDR (6 disks, rotation speed 2 rpm, air flow 50L.h⁻¹) was used for dye decolorization (50mg.L⁻¹) in malt extract-glucose medium at 22°C. Decolorization was measured spectrophotometrically and the end-products by GC/MS. Bacterial luminiscence test, plant growth tests and Ames test were used to measure acute toxicity and genotoxicity.

Conclusions

Using batch mode, RB19 decolorization rates were: Irpex lacteus > T. suaveolens > Pleurotus ostreatus. When continuous mode was used, the dye degradation capacity of T. suaveolens was 2-fold compared to the batch mode. Majority of the degradation products were aromatic alcohols. No genotoxicity was detected after degradation. The acute toxicity measured with Vibrio fischeri, Sinapis alba and Lemna minor decreased 1.2-, 2.7- and 1.6-fold, respectively. The results demonstrated the efficiency of fungal RDR for removal of textile anthraquinone dyes.

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SULFATE-REDUCING BACTERIA AS WHOLE-CELL BIOCATALYSTS FOR HYDROGEN PRODUCTION

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Background
Sulfate-reducing bacteria (SRB) are a major group of environmental bacteria with important biotechnological applications in waste water treatment and bioremediation. These organisms have an extremely high hydrogenase activity and in natural habitats where sulfate is limited they produce hydrogen fermentatively and grow syntrophically with other organisms. However, their potential as \( \text{H}_2 \)-producers has been poorly investigated.

Objectives
Given the high number of hydrogenases present in SRB genomes [1] we explored the potential of the model strain *Desulfovibrio vulgaris* Hildenborough for \( \text{H}_2 \) production from three substrates (lactate, ethanol, formate).

Methods
Among the substrates tested the highest \( \text{H}_2 \) production was observed from formate [2]. Formate has emerged recently as an environmental friendly storage of \( \text{H}_2 \) that can be easily transported and has relatively low toxicity. Therefore a lab-scale \( \text{H}_2 \) production process with gas sparging was designed to evaluate the potential of SRB as biocatalysts for formate-driven \( \text{H}_2 \) production [3].

Conclusions
By optimizing the operation conditions of bioreactor such as temperature, gas flow rate and cell load, a high \( \text{H}_2 \) production rate was obtained. In addition, a high specific hydrogen production rate and 100% efficiency of substrate conversion were achieved, demonstrating the high potential of SRB for \( \text{H}_2 \) production from formate [3]. Our studies reveal that SRB can be used as an efficient biocatalyst for the conversion of formate to hydrogen.


INVESTIGATING MICROALGAE AS A BIOTECH PLATFORM FOR HIGH-VALUE OILS

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Background

Natural oils are found in a diverse range of products such as soaps, paints, cosmetics, antibacterial treatments and nutritional supplements (fish oils).

Objectives

As part of a consortium with the University of Cambridge, the University of Aberdeen and Rothamsted Research we aim to establish the microalga *Nannochloropsis gaditana* as a platform for the production of these commercially important oils. The advantages of microalgae as a production platform include low media costs, lack of toxins and fast growth rates; *N. gaditana* was chosen due to its naturally high lipid content.

Methods

The production of bespoke lipids will require a combination of metabolic modelling and nuclear genetic engineering to introduce or knock out specific elongase and desaturase genes. To test the genetic tractability of this species, two constructs were prepared that contained ble-gfp or ble-HA fusion genes; the ble marker encodes resistance to zeocin. *N. gaditana* was transformed by electroporation and colonies were selected on media containing zeocin. All of the colonies tested carried the expected transgene and accumulated Ble-GFP or Ble-HA fusion protein, as detected by western blotting.

Conclusions

The *N. gaditana* nuclear genome is amenable to genetic transformation by random integration, paving the way for our metabolic engineering studies over the next three years.
MALIC ACID PRODUCTION FROM RENEWABLE SOURCES BY ASPERGILLUS ORYZAE

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Background

L-malic acid is a C4 dicarboxylic organic acid and considered as a promising chemical building block. It can be applied as food preservative and acidulant, in rust removal because of its chelator properties and as polymerization starter unit due to its bifunctionality. Up to now it is produced chemically from crude oil via maleic anhydride. The mould Aspergillus oryzae produces malic acid in large quantities from glucose and other carbon sources. The microbial production of organic acids from renewable sources has the potential to be a sustainable alternative to petroleum and to reduce greenhouse gases as CO₂ fixation is involved in microbial biosynthesis.

Objectives

The potential of malic acid production from renewable resources is evaluated. Therefore, different carbon sources based on lignocellulosic biomass, e.g. fractions of pyrolysis oil and hemicellulosic sugars, are tested.

Methods

A. oryzae was cultivated in preculture medium in shaking flasks for 17 hours. Due to nitrogen limitation and an excess of glucose in the production medium the fungus started to produce malic acid. A bioreactor process using glucose as carbon source was already established¹ and used as basis for the evaluation of alternative carbon sources. Organic acid concentration was measured by HPLC.

Conclusions

A. oryzae proved to be a promising natural host for malic acid production with the potential to use several renewable carbon sources.

References:

EXPLORING NON-CONVENTIONAL YEAST SPECIES AS SOURCE FOR ENHANCING ANAEROBIC XYLOSE FERMENTATION IN METABOLIC ENGINEERED SACCHAROMYCES CEREVISIAE

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Background

Efficient conversion of xylose, a five-carbon sugar present in significant amounts in the hemicellulose fractions of a wide range of lignocellulosic biomass, is required for the development of second generation bioethanol. Xylose is not naturally consumed by the baker’s yeast, Saccharomyces cerevisiae, therefore heterologous pathways has to be introduced by genetic engineering. This includes introducing one of the two known pathways for the conversion of xylose to xylulose: xylose isomerase (XI) or xylose reductase and xylitol dehydrogenase (XR-XDH). In most xylose-consuming fungal species, XR is an enzyme that uses NADPH as the dominant co-factor for xylose reduction to xylitol. It has been observed that S. cerevisiae strains harbouring XR variants with an increased preference for NADH have more efficient xylose fermentation, therefore exploring the diversity of XRs with different co-factor affinity is essential.

Objectives

In this work we have evaluated the xylose utilisation under anaerobic conditions from two constructed S. cerevisiae strains carrying different varieties of xylose reductases.

Methods

The strain TMB3422, carrying a mutated version of the Scheffersomyces stipitis XR gene and a constructed strain, carrying a novel XR gene from a fermenting yeast species that was isolated from rotting-wood samples, were compared for characteristics under fermentative conditions.

Conclusions
The best xylose fermenting yeast strain analysed revealed significantly higher ethanol yield and productivity, as well as lower xylitol yield under anaerobic conditions, which represents an advance towards the efficient fermentation of lignocellulosic hydrolysates to ethanol.
SOLID WASTE FROM COFFEE PROCESS FOR β-GLUCOSIDASE PRODUCTION BY BACILLUS SUBTILIS

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Background
The coffee production for the Brazil is a second important commodity however the solid and liquid waste during the process is an environmental problem. Some works focus on use these waste to produce organic acids, enzymes and other products

Objectives
The aims were optimize the β-glicosidase production for Bacillus subtilis in submerged fermentation using coffee pulp as substrate and evaluated the optimize condition culture in bioreactor.

Methods
The coffee pulp was dried at 60 °C for 72 h until reaching constant weight. After that, the coffee pulp was ground in Willey mill (1.0 mm). The microorganism used in study was Bacillus subtilis CCMA0087 belongs to Culture Collection of Agricultural Microbiology (CCMA) of Federal University of Lavras (Brazil), isolated from Brazilian Cerrado fruit namely Marolo. The enzyme production was done by Response Surface methodology. The media culture was based in Kasana et al. (2008) with modification (% NaNO₃, 0.1 K₂HPO₄, 0.05 MgSO₄, 0.05 KCl, 0.02 peptone and coffee pulp (0.32 – 3.68) as a carbon source. The β-glucosidase activity as measured by p-nitrophenyl-β-D-glucopyranoside (PNPG) as substrate.

Conclusions
In bench scale, Bacillus subtilis UFLA BCEF1130 show high β-glucosidase production (22.59 UI/mL) at pH 3.64, 36.6 °C and concentration coffee pulp of 36.8 g/L. However, in bioreactor the enzyme production was 2.5 less than bench scale in the same experimental conditions. The work continues to understand the scale up effect over the β-glucosidase production.
FEMS-0666
Biotechnology and industrial microbiology

CHARACTERISTIC OF EXTRACELLULAR HYALURONIDASE FORM PSEUZOZYMA APHIDIS YEAST

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Background

Hyaluronidases (Hz) are enzymes that degrade predominantly hyaluronan (HA). Until recently, three groups of Hz were reported: hyaluronate 4-glycanohydrolases (found in Vertebrates and venom of some insects), endo-β-glucuronidases (leeches and crustaceans) and hyaluronan lyases (bacteria). Recently a new group of HA degrading enzymes from Micromycetes belonging to genus Penicillium was reported [Bakke M, Kamei J & Obata A., 2011, DOI: http://dx.doi.org/10.1016/j.febslet.2010.11.021]. These enzymes hydrolyze β-1,4 glycosidic bonds in HA, similar to Vertebrate Hz, however have different conserved domains. We report discovery of another producer of hyaluronate 4-glycanohydrolase from the Kingdom Fungi, the yeast Pseudozyma aphidis.

Objectives

To characterize P. aphidis extracellular Hz.

Methods

P. aphidis was cultivated in bioreactor Techfors-S with 20 L working volume (Infors AG, Switzerland). The enzyme was isolated from cultural broth filtrate and partially purified by diafiltration on Sartorius Hydra membrane cassette with 10 kDa cut-off and ion-exchange chromatography on DEAE-Sepharose Fast Flow (GE Healthcare). The products of HA enzymatic hydrolysis were analyzed by HPLC and LC-MS. Gel permeation chromatography, SDS-PAGE and Morgan-Elson colorimetric assay were used for enzyme characteristic.

Conclusions

P. aphidis produced extracellular enzyme with hyaluronate 4-glycanohydrolase activity. Temperature and pH optima of HA hydrolysis were 45 °C and pH 3.0 respectively. The enzyme didn't hydrolyze neither chondroitin sulphate nor β-(1.3;1.6)-glucan schizophyllan. P. aphidis Hz showed high thermal stability that makes advantageous its application for enzymatic production of hyaluronan oligomers.
FEMS-0678
Biotechnology and industrial microbiology

PENICILLIUM CHRYSOGENUM B13, A FERULOYL ESTERASE FACTORY
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Background
Feruloyl esterases (FAEs) represent a diverse group of hydrolases catalyzing the cleavage of ester bonds between plant cell wall polysaccharide and phenolic acid. These enzymes are key tools for degradation of plant cell wall and are widely distributed in plants and microorganisms. FAEs release ferulic acid and other aromatic acids from these polymeric structures and have received an increasing interest in biotechnological processes for industrial and medicinal applications.

Objectives
In the worldwide market there are not fully purified FAEs or enough enzyme stocks. Discovery of new FAEs with novel properties continues to be an important research area that has increased drastically since 1990. Following this research line (CTM2012-32026 project¹), genetic engineering techniques to improve the FAEs production in the filamentous fungus Penicillium chrysogenum have been applied.

Methods
Several transformants, whose production of FAEs is significantly improved with respect to the parental strain, have been obtained. Special interest deserves B13 transformant that multiplies the FAE activity more than 90 times when the substrate of the enzyme is methyl ferulate and over of 14 when the substrate is sugar beet pulp. Furthermore, this FAE activity from the B13 strain was compared with the activity that is present in some commercial enzymes versus synthetic and natural substrates.

Conclusions
The analysis of the results revealed that the B13 enzymatic extracts triplicate the FAE activity in comparison with the analyzed commercial enzymes.

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PARAMETERS AFFECTING THE PRODUCTION OF LIPASE ENZYME BY A NEW ISOLATE YEAST.

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**Background**

Lipases (E.C.3.1.1.3; tri glycerol acyl hydrolases) are hydrolizer enzymes which catalyze free fat into glycerol and triglycerid. Industrial lipase commonly is provided by external cellular activity from strains of microorganisms. Microbial lipases are produced in liquid phase and its activity is influenced by concentrations of carbon/nitrogen resources, pH, temperature and metal ions.

**Objectives**

In this study, fungi as a lipase source, isolated from the village which is located near Eskisehir city. In the production of lipase, environmental conditions and media components are investigated as important parameters.

**Methods**

Vegetable oils as carbon source, carbohydrate source and nitrogen sources are used for lipase activity investigation. To find the optimal condition, firstly between 3.0 – 9.0 pH and 10 – 40\(^\circ\)C several experiments are carried out to detect the effects of these varieants on lipase production.

**Conclusions**

As a result of several experiments for production of lipase from *Rhodotorula*, the optimal condition is pH 5.0, temperature 30\(^\circ\)C, 1% concentration of carbon from olive oil and the best source of nitrogen, the pepton has been considered. As a result of combination of carbon and nitrogen sources with olive oil, the activity of lipase was increased.
PRODUCTION OF LIPASE ENZYME BY A NOVEL FUNGAL SOURCE, TRICHODERMA SP.
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Background

The industrial use of enzymes obtained from animals, plants and microorganisms has developed recently and the most important source for enzyme production has been determined as microorganisms. Lipases has a important application areas in industry.

Objectives

In this research the lipase enzyme which has a industrial importance was obtained from new fungal source that was isolated from soil. Subsequently it was determinated as \textit{Trichoderma citrinoviride} by the analysis of 18S rRNA sequence.

Methods

Lipase activity was determined by titrimetric assay. Optimum conditions were detected for production of lipase.

Conclusions

The lipase enzyme by isolated fungi (\textit{Trichoderma citrinoviride}) was extracellular and showed its highest activity on the 4. day of the incubation period. In the production of lipase, culture conditions and media components are investigated as important parameters. Various carbon and nitrogen sources were used for highest lipase activity. Also we detected that pH 5.5 and temperature 30 °C are optimum conditions for lipase production.
Background

The project MySterI (Mycobacterial Steroids for Industry), supported by the ERA-IB 3rd joint call, aims at producing high value steroid precursors using a novel bioconversion strategy that will yield a greening process with lower production costs (@MySterI_ERA_IB; http://www.era-ib.net/mysteri-0). The bioconversion of phytosterols (low cost plant material similar to cholesterol) to desired steroid precursors is performed using engineered fast-growing mycobacteria in a single fermentation step.

Objectives

The targeted precursors are: androst-4-ene-3,17-dione (AD), 3β-hydroxyandrost-5-ene-17-one (DHEA) and then 11α-hydroxyandrost-4-ene-3,17-dione (11-α-OH-AD) and testosterone.

Methods

The key points tackled by the Consortium are: i) Genome sequencing and annotation of *Mycobacterium* sp. NRRL B-3805 (AD-producer) to identify key bioconversion genes and to enable ‘omics tools; ii) Understanding of phytosterol bioconversion by means of ‘omics technologies; iii) Development of the genetic engineering tools for *Mycobacterium* sp. NRRL B-3805; iv) Construction of mycobacterial strains capable of producing 11-α-OH-AD, DHEA and testosterone; v) Designing more efficient and eco-friendly methods of production and downstream processing for the three selected compounds.

Conclusions

The MySterI pipeline is a top to bottom process that brings together six interdisciplinary research groups from Universities, Research Centres and Industrial companies in 3 different EU member states [Spain (INBIOTEC; Gadea Biopharma), Germany (Technische Universität Dortmund) and UK (University of York)], as
well as Norway (Stiftelsen SINTEF) and Russia (Pharmins Ltd.). Therefore, the MySterI results in progress are: i) novel strains capable of producing three valuable C19-steroid precursors from phytosterol ii) knowledge of the biochemistry of steroid biotransformations iii) optimized fermentation and eco-friendly downstream processes for the single-step production.
GENOME SEQUENCING AND ‘OMICS DEVELOPMENT FOR THE STEROID PRECURSORS PRODUCER MYCOBACTERIUM SP. NRRL B-3805


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Background
Microbial biotransformation to generate new steroid drugs is a well-established application of industrial biotechnology. Thus, several worldwide enterprises are involved in steroid production (e.g., Gadea Biopharma SL, Sanofi, Pfizer). Biotransformation can begin with different raw materials, but the mixture of plant sterols (phytosterols) have been the research focus in recent decades. However, the studies to date show that improvements in phytosterols broth culture solubility, product recovery and mainly in microbial strains are needed.

Objectives
Optimization of the whole molecular biology methodologies valid for a non-well characterized microbial strain has been carried out on the androstenedione (AD)-producer strain Mycobacterium sp. NRRL B-3805. Thus, i) growth conditions; ii) phytosterols solubility analysis; iii) growth titration in dirty media; iv) steroids quantitation by HPLC; v) genomic DNA extraction suitable for genome sequencing; vi) high quality RNA for transcriptomics (RNAseq, microarrays) analyses; as well as vii) secreted protein purification for 2D-DIGE secretome methods have been improved.

Methods
An example of successful optimized process is the genome sequencing pipeline defined for Mycobacterium sp. NRRL B-3805. This genome sequencing and annotation, as well as the understanding of phytosterols bioconversion by means of ‘omics technologies (RNAseq, microarrays, 2D-DIGE) are capital for the ERA-IB 3rd joint call supported project MySterI (@MySterIERA_IB; http://www.era-ib.net/mysteri-0).

Conclusions
Thus, key bioconversion genes and target genes suitable for the development and use of genetic engineering tools have been defined. This genome was fully sequenced using Ion Torrent and Sanger technologies. It comprises 5,421,338 bp in a circular chromosome with a GC content of 67 %.
SOLID PHASE EXTRACTION BASED ON THE USE OF HELVELLA CRISPA AS A Fungal Biomass For The Preconcentrations Of Pb And Al By ICP-OES

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Background
The use of fungal biomass for the preconcentration and separation of traces of heavy metals and some organic materials at trace levels are popular due to the good adsorption properties such as high surface area and high adsorption capacity of the fungus.

Objectives
This study describes the biosorption potential of Helvella crispa as a sorbent for solid phase extraction (SPE) and preconcentration of Pb²⁺ and Al³⁺.

Methods
The model solution was passed through the prepared SPE column at an adjusted flow rate with a peristaltic pump. Then, 10.0 mL distilled water was passed through the column. The retained Pb²⁺ and Al³⁺ on the fungal biomass was then eluted with 5.0 mL of 1.0 mol L⁻¹ HCl. The concentration of Pb²⁺ and Al³⁺ in this solution was determined by ICP-OES.

Conclusions
The limit of detection (LOD) of Pb and Al was found to be 0.10 ng mL⁻¹ and 0.03 ng mL⁻¹, respectively. The sensitivity of ICP-OES improved 39.8-fold for Pb and 39.5-fold for Al. Linearity was obtained in the concentration range of 1.25–50.0 ng mL⁻¹ for Pb and 0.5–50.0 ng mL⁻¹ for Al. The relative standard deviation (RSD) of the method under optimum conditions was lower than 8.4% (n=5) for Pb and Al, which was validated through the analysis of certified reference tea and poplar leaves samples. The biosorption capacity of immobilized Helvella crispa for Pb and Al was found to be 31.2 mg g⁻¹ and 45.7 mg g⁻¹, respectively.
WHICH IS A BETTER APPROACH FOR ENZYME EXPLORATION?: CULTURE-DEPENDENT AND -INDEPENDENT (METAGENOMIC) APPROACHES TO THE SCREENING FOR AROMATIC-DEGRADING ENZYMES.

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Background
Although metagenomics is regarded as one of the best approaches for mining enzymes, its superiority over traditional culture-dependent approach is still controversial.

Objectives
Therefore, culture-dependent and -independent approaches were used for extensive retrieval of the extradiol dioxygenase (EDO) gene from the environment to investigate the relationship between the EDO genes from isolated bacteria and the metagenomic EDO genes from which they were isolated.

Methods
In our previous study, we identified 91 fosmid clones showing EDO enzyme activity using a metagenomic approach (Environ Microbiol 9, 2289-2297, 2007). In the present study, we newly isolated 88 phenol-utilizing bacteria from the same environmental sample and identified four EDO genes from them. Of these, two EDOs had amino acid sequences similar to those reported previously in aromatic-utilizing strains, and one EDO had a sequence almost identical to that of metagenomic EDOs identified in our previous study. Unexpectedly, one EDO showed no similarity to any class I EDOs and was categorized as class II, which has not been found in past any metagenomic approaches. Quantitative polymerase chain reaction (PCR) assay indicated that the low-abundance class II EDO gene can be enriched by culturing approaches.

Conclusions
We conclude that the combined use of the two approaches can explore the gene community more extensively than their individual use.

ENGINEERING STREPTOMYCES DIASTATOCHROMOGENES 1628 TO INCREASE THE PRODUCTION OF TOYOCAMYCIN

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Background
Toyocamycin (TM) is a member of the nucleoside antibiotics. It shows antifungal activity and might be used to treat plant diseases. TM is produced by *Streptomyces diastatochromogenes* 1628 but the production rate is always very low. Recently the gene *toyG* was cloned and its function in TM biosynthesis was reported. However, nothing was known about the influence of *toyG* on TM production.

Objectives
We studied the influence of three genes on TM production. *Vgb, frr*, and *toyG* were co-expressed, either in double- or triple-combinations.

Methods
Over-expression of these genes in double or triple leads to great significant increased production of TM in strain *S. diastatochromogenes* 1628. At the end, among all these different strains constructed in this work, the strain *S. diastatochromogenes* 1628-VGF, harboring *vgb, frr* and *toyG* that was placed under the control of *PermE*, respectively, exhibited the largest increase in TM production. Notably, the TM production of *S. diastatochromogenes* 1628-VGF reached the highest level at 831.2 mg/L at 72 h, while TM yield of original strain *S. diastatochromogenes* 1628 at 415.1 mg/L at 84 h.

Conclusions
Based on known knowledge, TM yield produced by engineered strain *S. diastatochromogenes* 1628-VGF was the highest as ever reported. The high-yield, energy-saving strain *S. diastatochromogenes* 1628-VGF constructed in this study shall be beneficial for the industrial production of TM.
EXPRESSION OF THE CELLULOSOMAL GENE CLUSTER OF CLOSTRIDIUM CELLULOVORANS IN BUTANOL-PRODUCING CLOSTRIDIUM BEIJERINCKII

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Background
Biofuel production from cellulosic biomass consists of pretreatment, saccharification, fermentation and distillation. However, production is hindered by the high cost compared with energy crops, such that innovative technologies are required. One approach is consolidated bioprocessing (CBP), in which pretreatment, saccharification, and fermentation are consolidated.

Objectives
Clostridium cellulovorans 743B, an anaerobic and mesophilic bacterium, produces a large extracellular enzyme complex (cellulosome) that efficiently hydrolyzes cellulosic biomass. The cellulosomal gene cluster of C. cellulovorans includes cbpA, which encodes a scaffold protein, and seven downstream glycoside hydrolase genes. Clostridium beijerinckii NCIMB 8052 cannot produce a cellulosome and does not degrade soft biomass, but instead produces butanol, acetone, and ethanol from mono- and disaccharides. The aim of this study was to express the cellulosome of C. cellulovorans in C. beijerinckii to obtain a bacterium able to carry out CBP.

Methods
The large cellulosomal gene cluster containing the cbpA gene was used to transform C. beijerinckii by electroporation. Protein-level expression of the gene cluster encoding the scaffold protein CbpA and mannanase A was demonstrated by western blotting. Cellulose and galactomannan activities on polysaccharide substrates were evaluated by Congo red staining.

Conclusions
The transformant was shown to be capable of hydrolyzing galactomannan and produced butanol directly. This butanol-producing bacterium provides the basis for the CBP of lignocellulose to biobutanol.
EFFECTS OF IRON AND MAGNESIUM IONS COMBINATION ON HYDROGEN PHOTOPRODUCTION BY RHODOBACTER SPHAEROIDES

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Background

Iron and magnesium are essential for growth of photosynthetic bacteria. Mg\(^{2+}\) participates in the structure of photosynthetic electron carriers and pigments. Fe\(^{2+}\) is known as the component of enzymes, involved in bio-hydrogen (H\(_2\)) production.

Objectives

In this work the effects of metal ions combinations on growth and H\(_2\) photoproduction by *Rhodobacter sphaeroides* MDC6522, isolated from Jermuk mineral springs in Armenia, in compare to single ions effects, were investigated.

Methods

*R. sphaeroides* was cultivated under anaerobic conditions upon illumination in the presence of Fe\(^{2+}\) and Mg\(^{2+}\) various concentrations. The H\(_2\) yield was calculated by the decrease of redox potential, measured by platinum electrode.

Conclusions

The results pointed out the concentration dependent effects of metal ions. *R. sphaeroides* was unable to grow well in the absence of Fe\(^{2+}\) and Mg\(^{2+}\), which indicates the importance of these ions for bacterial growth. The highest growth rates of *R. sphaeroides* cells were obtained in the presence of 0.08 mM Fe\(^{2+}\) and 5 mM Mg\(^{2+}\), in compare to control (0.04 mM Fe\(^{2+}\) and 1 mM Mg\(^{2+}\)). The simultaneous addition of metal ions stimulated bacterial growth rate ~1.2-fold in compare to the single ions. The highest H\(_2\) yields were obtained in the presence of 0.08 mM Fe\(^{2+}\) and 10 mM Mg\(^{2+}\), which were ~1.5-fold higher than control. However, the H\(_2\) production was increased ~2.5-fold, when two metals were added simultaneously in concentration 0.08 mM Fe\(^{2+}\) and 5 mM Mg\(^{2+}\). Thus, combinations of metal ions in appropriate concentrations are preferable for H\(_2\) production enhancement in *R. sphaeroides*. 
HIGH-LEVEL EXPRESSION, RAPID ON-COLUMN REFOLDING AND PURIFICATION OF RECOMBINANT HUMAN VASCULAR ENDOTHELIAL GROWTH FACTOR 165 IN ESCHERICHIA COLI

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Background
Vascular endothelial growth factors (VEGFs) are a family of proteins that promote angiogenesis and participate in a variety of physiological and pathological processes. VEGF165 is the major isoform induced by hypoxia, oncogene mutations, and cytokines such as IL-1, IL-8, and TNF-α and secretes by many cell types. It is a potent angiogenic factor and mitogen that stimulates proliferation, migration, and formation of endothelial cells. Cancerous cells are able to grow and metastasize by VEGF secretion.

Objectives
A recombinant VEGF165 might be used in order to produce an antibody against it and also it could be used as a growth factor in some biological mediums.

**Methods**
The RNA was extracted from Hela cell, and them used for cDNA synthesis. The gene encoding human VEGF isoform 165 (hVEGF165) was cloned into the expression vector pET32a (+) to construct a fusion expression plasmid that induced the thioredoxin (Trx) Gene. The expressed protein was purified by affinity chromatography using Ni-NTA resin. High concentration of the recombinant protein obtained from a single-step.

**Conclusions**
Fusion of VEGF to thioredoxin and careful codon optimization of the eukaryotic sequence could be improved to a high-level insoluble protein expression in comparison to an un-optimized, His-tagged construct. The thioredoxin-fused protein was successfully purified using an on-column Ni-NTA purification procedure. SDS-PAGE and Western Blotting analysis confirmed the desired expression. rhVEGF165 could readily and rapidly purify with on-column Ni-NTA system chromatography.
ADHESION OF ALKANOTROPHIC ACTINOBACTERIA TO SOLID SURFACES AND HYDROPHOBIC LIQUIDS

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Background
Mechanisms of adhesion of biotechnologically significant alkanotrophic actinobacteria are poorly studied.

Objectives
1. To study the adhesion of actinobacteria to a wide range of solid carriers and hydrophobic liquids under various growth conditions. 2. To determine physical–chemical and molecular factors of actinobacterial adhesion. 3. To define adhesion conditions providing the highest catalytic activity of actinobacteria.

Methods
100 strains from the IEGM Collection of Alkanotrophic Microorganisms (acronym IEGM, WDCM # 768, www.iegm.ru/iegmcol) were used in this study. The substrates for adhesion were polystyrene microplates, technical polymer tissue, industrial wastes (sawdust, poultry feathers, and leather), \( n \)-alkanes C10–C16, solvents, and different oils. Molecular genetic and biochemical mechanisms of adhesion were studied using targeted and \textit{in vivo} Tn5 mutagenesis, chemical analysis of cell wall, and specific inhibition. Cell morphology, nanomechanical properties, adhesion forces, and distribution of actinobacteria on solid carriers were studied by interference microscopy and combined confocal and laser scanning atomic force microscopy. Thermodynamic data were obtained using infrared thermography.

Conclusions
Adhesion was shown to be a key mechanism in hydrophobic compound oxidation by
actinobacteria. Involvement of glycolipid biosurfactants, proteins, and cell appendages in adhesion was revealed. Monolayer actinobacterial biofilms with high (115 mg·l\(^{-1}\)·h\(^{-1}\)) hydrocarbon-oxidizing activities were formed on solid surfaces hydrophobized with glycolipid biosurfactants (Figure). The results obtained could be used to develop biocatalysts for targeted biotransformations and pollutant degradation. The research was supported by the RFBR grant (14-04-96013), the RSF grant (14-14-00643), and the President of the Russian Federation for Leading Scientific Schools’ grant (4607.2014.4).
BIOTRANSFORMATION OF BETULIN USING RHODOCoccus ACTINOBACTERIA

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Background
Bacterial transformation of triterpenoids (in particular betulin) is an efficient method for single-stage production of biologically active compounds with high regio- and stereoselectivity.

Objectives
1. To demonstrate the possible use of Rhodococcus actinobacteria for selective biotransformation of betulin, a pentacyclic lupine-type triterpenoid. 2. To study the interaction mechanisms between rhodococci and betulin, and its biotransformation pathways. 3. To evaluate the possible application of products from bacterial betulin oxidation for subsequent synthesis of biologically active compounds using chemical methods.

Methods
The biological resources of the IEGM Collection of Alkanotrophic Microorganisms (acronym IEGM, WDCM # 768, www.iegm.ru/iegmcol) were used in this study. Conventional bacteriological methods, including betulin transformation by resting rhodococcal cells were employed. The mechanism of cell interaction with betulin was determined using methods of phase contrast, confocal lazer scanning, and atomic force microscopy. Rhodococcal viability was estimated using a LIVE/DEAD® BacLight™ Bacterial Viability Kit (Invitrogen, USA).

Conclusions
Biotransformation of betulin (0.5 and 3.0 g/l) using Rhodococcus actinobacteria was demonstrated for the first time. Betulone was identified as a major metabolite. The optimal conditions for betulin bioconversion by resting cells of R. rhodochrous IEGM 66 were selected. Further chemical modifications of betulone resulting from bacterial betulin transformation lead to the formation of 3,4-secobetulone with a marked cytotoxic activity (IC50 3.05 мкМ) against melanoma.

The research was supported by the RFBR grant (14-04-96017), and the President of the Russian Federation for Leading Scientific Schools’ grant (4607.2014.4).
COMMERICALLY IMPORTANT HIGH MOLECULAR WEIGHT DEXTRAN FROM INDIGENOUS WEISSELLA CIBARIA CMGDEX3

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Background
Extracellular polysaccharides (EPS) of microbial origin are associated with the cell surface or excreted in the extracellular environment. Several food grade microorganisms, including lactic acid bacteria (LAB), synthesize EPS. The synthesis of EPS by LAB has gained remarkable interest because of LAB categorization as safe organisms and vast utilization of synthesized EPS in markets from food to health and industry. Dextran is a polysaccharide composed of D-glucose units which is synthesized by various LAB under appropriate conditions and widely used in several industries including food, pharmaceutical and chemical.

Objectives
Present study was designed to isolate LAB capable to synthesize good quality dextran and characterize the synthesized dextran.

Methods
In this study five bacterial strains were isolated from different vegetables and fruit samples on the basis of slime production in selected agar medium with 15% sucrose. When those five strains were cultivated in medium for the production and extraction of EPS, the highest crude yield of EPS was obtained from CMGDEX3 which was isolated from cabbage and identified as Weissella cibaria by 16S rRNA gene sequencing. EPS extracted from W. cibaria CMGDEX3 was purified and analyzed for structure by FTIR, ¹H and ¹³C NMR spectroscopy. Molecular weight was determined by gel permeation chromatography.

Conclusions
Analysis of EPS demonstrated that W. cibaria CMGDEX3 synthesized commercially important linear dextran that predominately had α (1→6) glycosidic linkages with few (3.4%) α (1→3) linked branches. Molecular mass determination showed that it was a high molecular weight dextran of an average > 2,000,000 Daltons.
SCREENING AND SELECTION OF PROBIOTIC LACTIC ACID BACTERIA INHIBITING PATHOGENIC BACTERIA AND VIRUSES

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Background

Probiotic refers to viable, nonpathogenic microorganisms that, when ingested, are able to reach the intestines in sufficient numbers to confer health benefits to the host. Intake of probiotics reduces the amount of pathogens improving thus the intestinal microbial balance of the host and lowering the risk of gastro-intestinal diseases.

Objectives

To isolate potential probiotic bacterial strains inhibiting rota- and polio- viruses as well as pathogenic bacteria.

Methods

Three thousands bacterial strains isolated from various sources including human milk and fermented foods were screened for potential probiotics inhibiting rota- and polio-viruses as well as pathogenic bacteria such as enterotoxigenic Escherichia coli, Clostridium perfringens, Listeria monocytogenes, Staphylococcus aureus, Salmonella Typhimurium,. The antibacterial activity of each culture filtrate was determined by agar-diffusion method and the antiviral activity of heat–treated culture broth was examined by reduction of plaque forming unit in vitro culture system. The properties of the bacterial strains with particular functions such as adherent ability to enterocyte, tolerances towards gastric juice, bile, and tolerances to heat and cold-dry conditions were further investigated to select strains for probiotic use.

Conclusions

One potential lactic acid bacterium with desirable probiotic characteristics as well as the highest anti-viral and anti-bacterial activities was selected and identified as Lactobacillus plantarum according to its 16rRNA gene sequence.
OPTIMIZATION OF THE SECRETION OF BACTERIOCINS AND THE BACTERIAL CELL GROWTH OF GEOBACILLUS SP. 15 STRAIN

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Background

Bacteriocins are natural peptides secreted by variety of bacteria and provide the bacteria with a competitive advantage in their environment, eliminating competitors to gain resources. Bacterocin production may be strongly influenced by growth environment and medium composition. Many different studies have indicated the potential usefulness of bacteriocins in food preservation, veterinary, medicine or as phytosanitary measures for plant protection. To date, only few bacteriocin producing thermophilic bacterial strains of genus Geobacillus have been identified. Studies of factors affecting the production of bacteriocins by Geobacillus spp. bacteria are relatively scarce.

Objectives

This research was focused on the analysis effect of salts and to establish optimal conditions for bacteriocins production and bacterial cell growth for thermophilic bacteria Geobacillus sp. 15 strain.

Methods

Geobacillus sp. 15 strain was used as a bacteriocin producer. Geobacillus stearothermophilus NUB36187 (BGSC No. 9A11) was selected as the indicator strain for the bacteriocin assay. Nutrient broth (NB) liquid media were chosen for the strain growth and bacteriocin expression monitoring. It were supplemented with salts: KCl, ZnSO₄, MnSO₄, CuSO₄, MgCl₂, CaCl₂, NaNO₃, NaHCO₃. Bacteriocin activity was detected by agar well diffusion assay.

Conclusions

The greatest influence on bacterial growth was in NB medium with MnSO₄ or with KCl and MgSO₄. The best bacteriocin expression was achieved in media supplemented with two salts at a time: KCl and MgCl₂, MnSO₄ and NaHCO₃ or CaCl₂ and ZnSO₄.
Background
This study presents the development of consolidated bioprocessing for succinate production from microalgal biomass using engineered *Corynebacterium glutamicum*.

Objectives
Our objective in this study is to develop the *C. glutamicum* strain capable of utilizing starch as carbon source using the synthetic biology-platform and to apply the strain for production of succinate from microalgal biomass as sole carbon source.

Methods
We developed the starch-degrading and succinate-producing *C. glutamicum* strains, which produced succinate (0.16 g succinate/g total carbon source) from a mixture of starch and glucose as a model microalgal biomass. Subsequently, the engineered C. glutamicum strains were able to produce succinate (0.28 g succinate/g of total sugars) from pretreated microalgal biomass of CO$_2$-grown *Chlamydomonas reinhardtii*. For the first time, this work shows succinate production from CO$_2$ via sequential fermentations of CO$_2$-grown microalgae and engineered *C. glutamicum*.

Conclusions
In conclusions, consolidated bioprocessing based on microalgal biomass could be useful to promote variety of biorefineries.
AQUEOUS TWO PHASE EXTRACTION OF JONESIA DENITRIFICANS XYLANASE 6 IN PEG 1000/PHOSPHATE SYSTEM.

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Background
The impetus for research in the field of bioseparation has been sparked by the difficulty and complexity in the downstream processing of biological products. Indeed, 50% to 90% of the production cost for a typical biological products resides in the purification strategy. There is a need for efficient and economical large scale bioseparation techniques which will achieve high purity and high recovery, while maintaining the biological activity of the molecule. One such purification technique which meets these criteria involves the partitioning of biomolecules between two immiscible phases in an aqueous system (ATPS).

Objectives
The purification of xylanase from the strain Jonesia denitrificans BN13 using ATPS an efficient and economical bioseparation technique.

Methods
The Production of xylanases is carried out in 500ml of a liquid medium containing birchwood xylan. In each ATPS, PEG 1000 is added to a mixture consisting of dipotassium phosphate, sodium chloride and the culture medium inoculated with the strain Jonesia denitrificans. The concentration of PEG 1000 was varied : 8 to 16 % and the NaCl percentages are also varied from 2 to 4% while maintaining the other parameters constant, the xylanolytic activity was detected by zymogram coupled to SDS-PAGE.

Conclusions
The results showed that the best ATPS for purification of xylanases is composed of PEG 1000 at 8.33%, 13.14 % of K₂HPO₄, 1.62% NaCl at pH 7. We obtained a yield of 96.62 %, a partition coefficient of 86.66 and a purification factor of 2.9. The zymogram showed that the activity is mainly detected in the top phase.
Background

The filamentous actinobacterium Microbispora sp. ATCC-PTA-5024 produces the lantibiotic NAI-107 (Maffioli et al., 2014), which is effective against multidrug-resistant Gram-positive pathogens (Jabés et al., 2011).

In actinomycetes, the biosynthesis of antibiotics is generally elicited as a physiological response controlled by a complex regulatory network involving global regulators, playing pleiotropic roles, and pathway-specific regulators, which activate the biosynthesis of biologically active molecules (Bibb, 2005).

Objectives

The integration of bioinformatic tools and holistic technologies has allowed the development of consolidated strategies to manage huge amounts of complex molecular information on gene expression and biochemical capabilities deriving from 'omic' investigations, ultimately leading to novel approaches to explore microbial physiology. In the perspective of using Microbispora strains for the industrial synthesis of a NAI-107, insights on the molecular physiology of this actinobacterium would be beneficial to develop robust and economically-feasible production
An extensive investigation on proteomic changes associated with lantibiotic production was performed on the *Microbispora* sp. ATCC-PTA-5024 wild type strain at different growth stages by using combined two-dimensional difference in gel electrophoresis and mass spectrometry approaches. To evaluate the effect of NAI-107 on bacterial vitality, comparative proteomic experiments were also performed on a not producing strain following NAI-107 addition. The obtained results were then integrated with bioinformatics, fluorescence microscopy and molecular genetic experiments.

The results of this study elucidate the regulatory networks, the biochemical pathways and the molecular processes occurring during growth and lantibiotic production (Figure 1), thus providing the first functional picture of a member of the *Microbispora* genus.

**Figure 1.** Synoptic scheme summarizing the signalling systems and the regulatory cascades associated with the physiological differentiation processes in *Microbispora* sp.
OPTIMISATION OF THE BIOFLOCCULANT PRODUCED BY PANTOEA SP., A NOVEL BACTERIUM ISOLATED FROM SEDIMENT FROM THE BEACH AT MTUNZINI, KWAZULU-NATAL, SOUTH AFRICA

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Background
We screened microorganisms with potential for bioflocculation. An organism that secretes large amounts of bioflocculant was isolated from a sediment sample collected from the beach at Mtunzini on the coast of northern Kwazulu-Natal, South Africa.

Objectives
The primary objective of this study was to optimised the bioflocculant production by the Pantoea sp. Further studies on the characterisation and applications of the bioflocculant produced by this organism will be done to underscore the biotechnological importance of this species as a producer of secondary metabolite useful for water and wastewater treatment.

Methods
From detailed morphological and biochemical analysis, the isolate was identified as Pantoea sp. with 91% probability. Cultivation studies revealed that the isolate produced bioflocculant optimally with sucrose as a source of carbon (92.4% flocculating activity), inoculum size of 3% (v/v), at an initial pH of 6.0, with Ca²⁺ as an aid to coagulation, and a production time of 96 h.

Conclusions
We proposed that this organism has great potential as a producer of bioflocculant due to its ability of producing high flocculating activity when cultivation conditions are varied.
INFLUENCE OF THE MOLASSES AND AERATION ON THE PRODUCTION OF CELLULOSE BY GLUCONACETOBACTER XYLINUS

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Background
The bacterial cellulose (BC), produced by Gluconacetobacter xylinus, has been presented as a useful alternative to replace the vegetable cellulose as well as their use in many industrial applications. A carbon source and a system optimized for efficient aeration is important for a greater production of BC.

Objectives
This study has as its objective determine the effect of molasses in the production of BC by Gluconacetobacter xylinus IFO 13693 under static conditions and static with aeration.

Methods
The synthesis of bacterial cellulose (BC) by Gluconacetobacter xylinus was carried out in static culture discontinuous with and without aeration intermittent at room temperature, in the presence of molasses as the main carbon source at initial concentrations of 13.3 % (w/v). The concentration of BC concentrations, sucrose, glucose and fructose were determined each week. To determine the formation of cellulose and the coefficient of performance of the product was used the Microcal Origin software 6.0®.

Conclusions
By the fourth week, BC values found in the static and static and aeration experiments were around 103 g/L and for static 78.7 g/L, respectively. The kinetics for the hydrolysis of sucrose in the medium fixed the model of Michaelis-Menten, with a Vmax of 0.0041 moles/L/h and 0.0036 moles/L/h, and a Km of 0.021 M and 0.019 M for each culture condition, respectively. The yield coefficients have values of 0.013 to 8.7 g dry cellulose/g of glucose consumed. This shows the importance of the substrate and aeration on the synthesis of cellulose.
Background
Laccases are polyphenol oxidases with numerous industrial and bioremediation application, most laccase activity is demonstrated in fungal species.

Objectives
The study evaluated the potential of fungi isolated from selected agro waste for laccase production

Methods
Fungi cultures were screened for laccase production by plate test using 2, 2, azinobis (3 ethyl benthiazoline 6 sulphonate) (ABTS) and submerged culture using synthetic medium.

Conclusions
Utilisation of selected agro wastes (sawdust, planatian and banana peel) residue for laccase production was evaluated. Five out of twelve isolates were postive and identified as GEOTRICUM Spp,CEPHALOSPORIUM Spp, TRICHODERMA Spp, TRAMETES Spp and FUSARIUM Spp. Optimum enzyme activity was observed using TRICHODERMA Spp at 57.1U/ml followed by TRAMETES Spp with 51.99U/ml, FUSARIUM Spp had 29.2U/ml while GEOTRICUM Spp and CEPHALOSPORIUM Spp had 28.04U/ml and 9.72U/ml respectively. TRICHODERMA Spp was selected as the most potent in producing enzyme and therefore used for further studies. Effect of carbon source, inoculum sizes, pH and shaker speed was evaluated. Sawdust gave the highest yield with 151.17U/ml followed by plantain peel with 62.49U/ml,banana peel was lest with 54.94U/ml. The optimum pH for sawdust, plantain peel and banana peel was 5.95, 5.94 and 5.83 respectively. using shaker incubator @ 150rpm with sawdust as carbon source,Laccase yield of 310U/ml was obtained at pH of 6 and temperature of 25 degree celcius inoculum size of 1000000 spores per ml. Thus sawdust can serve as cheap substrate for laccase production
IDENTIFICATION OF CELLULOSOMAL ENZYMES FOR THE CONSTRUCTION OF CELLULOSOMES FROM CLOSTRIDIUM CELLULOVORANS

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Background

Clostridium cellulovorans 743B, an anaerobic mesophile, produces a multienzyme complex, called the cellulosome, for the efficient degradation of cellulosic biomass. The cellulosome consists of carbohydrate enzymes and scaffolding proteins. The scaffolding protein CbpA consists of nine hydrophobic repeat domains (cohesin domains) for the binding of enzymes containing dockerin domains (cellulosomal enzymes).

Objectives

The expression pattern of the cellulosomal enzymes in the presence of various carbon sources was previously shown by proteome analysis, but the constituent enzymes of the cellulosome and their arrangement as a complex have yet to be determined. In this study, we identified the cellulosomal enzymes involved in the cohesin-dockerin interaction.

Methods

C. cellulovorans was cultivated on cellobiose and its cellulosome was purified from cell extracts using Avicel affinity chromatography. Fourteen enzymes were found to make up the cellulosome. Nine glutathione-S-transferase (GST) fusion mini-scaffolding proteins, each containing a cohesin domains of CbpA, were expressed in Escherichia coli. Cellulosomal enzymes that specifically or nonspecifically bound cohesin domains were detected by GST pull-down assay, which was also used to qualitatively analyze the interaction between five types of dockerin domains and all cohesin domains.

Conclusions

The interaction was found to be significantly dependent on the type of dockerin domain and the particular cohesin domain. These results imply a specific arrangement of enzymes in the cellulosome.
Background

Microbial lipases catalyze hydrolysis of lipids, enabling bacteria to use vegetable and other oils as a carbon and energy source. Lipid hydrolysis share common intermediates with synthesis of valuable biopolymer polyhydroxyalkanoate (PHA). Bringing together these two processes in a single microorganism could further increase its potential for application in industrial biotechnology.

Objectives

The objective was to screen *Pseudomonas* strains for lipase activity and capability to accumulate PHA in order to identify or to genetically modify these microorganisms to use waste cooking oils for production of valuable biopolymers.

Methods

Seven *Pseudomonas* strains were tested for lipase activity on agar plates containing either Tween 80 or olive oil as carbon source. Bacterial capability to utilize reused fryer oil as a sole carbon source was tested in liquid medium. For selected *Pseudomonas* strains lipase activity was quantified using para-nitrophenyl palmitate enzyme assay and genetic determinants of lipase activity were amplified. In order to select appropriate host for the heterologous expression of lipase all *Pseudomonas* strains were tested for their ability to accumulate PHA.

Conclusions

Among seven tested *Pseudomonas* strains *Pseudomonas aeruginosa* PAO1 showed highest lipase activity, while both *P. aeruginosa* PAO1 and *Pseudomonas putida* KT2440 could accumulate PHA. Since *P. aeruginosa* PAO1 is known opportunistic pathogen, *P. putida* KT2440 was more appropriate host for the biopolymer
production. *P. aeruginosa* PAO1 lipase-encoding genes were amplified and cloned into *P. putida* KT2440, and heterologous expression was optimized. Further analyses should reveal if these constructs have potential for the industrial application.
MYSTERI: MYCOBACTERIAL STEROIDS FOR INDUSTRY

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Background

Steroid-based pharmaceuticals are extremely important for health, and conversion of cheap agricultural waste into highly valuable steroids is therefore an attractive prospect. Previous studies have indicated that in order to achieve this several areas need improvement, including the microbial strains used for bioconversion.

Objectives

By utilizing the bacterium Mycobacterium sp. NRRL B-3805 we aim to replace the current microbial and chemical multistep processes with a single-step production of several steroids including androstenedione (AD), 11α-hydroxy-androstenedione (11α-OH-AD), dehydroepiandrosterone (DHEA) and testosterone from a cheap phytosterol feedstock.

Methods

We have developed genetic engineering tools and procedures to manipulate Mycobacterium sp. NRRL B-3805 to improve the bioconversion of phytosterols. Mycobacterium sp. NRRL B-3805 metabolises steroids through the action of cholesterol oxidases, which modify the A-ring of the steroid core before cleaving the side chain through multiple enzymatic steps to produce AD. The genome of Mycobacterium sp. NRRL B-3805 was sequenced and was mined for potential cholesterol oxidases, hydroxysteroid dehydrogenases and other known enzymes involved in cholesterol catabolism. The genome mining enabled strain manipulation, as a means to optimise steroid production. Quantitation of steroid production was determined using GCMS.

Conclusions

Production of 11α-OH-AD is being investigated by amplifying a putative 11α-hydroxylase and the associated oxidoreductase from Aspergillus ochraceus cDNA and placing both coding sequences under bidirectional inducible promoters for expression in Saccharomyces cerevisiae and 11α-OH-AD production in media
containing AD. If active these enzymes will be expressed in *Mycobacterium* sp. NRRL B-3805.
Background

In the future, petrochemical industry may be replaced by biotechnological processes producing building block chemicals from renewable carbon and employing genetically engineered bacteria and other microorganisms. However, besides searching for suitable microbial production platforms, a major hurdle for a large-scale implementation of bioprocess-based technologies is the availability of economic and sustainable carbon sources.

Objectives

An alternative substrate not competing with food or fodder production might be methanol due to its high synthesis capacities and a potentially sustainable production from natural gas, agricultural waste materials and biogas. However, main weaknesses of this C1 compound are its high volatility and toxicity. Its use in fermentation processes therefore calls for an innovative real-time control strategy.

Methods

In this study, we decided on biocalorimetry as a basis for optimizing C1 substrate feed regimes in fermentations with the methylotrophic bacterium *Methylobacterium extorquens*. According to Hess’ law every metabolic flux is quantitatively related to the heat production rate. Thus, calorimetry provides real-time stoichiometric and kinetic information of the fermentation process and allows reacting fast and efficiently to any metabolic changes.

Conclusions

By comparing different calorimetry-based control strategies we established a procedure leading to the highest growth and product formation rates of *M. extorquens* on methanol. Additionally, we could show that this approach can be extended to other, even more toxic substrates such as formic acid. In conclusion, this study demonstrates that biocalorimetry is a very efficient tool for process control and can be applied to optimize any microbial product formation process.
PHYSICOCHEMICAL AND BIOLOGICAL PROPERTIES OF THE ANTIFUNGAL POLYENE ANTIBIOTIC ROSEOFUNGIN

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Background
Due to the upward trend of fungal infections and development of resistance to existing drugs, the requirement for efficient antifungal agents has increased.

Objectives
Strain Streptomyces roseoflavus var. roseofungini 1-68, a producer of the antibiotic roseofungin.

Methods
Methods for spectrometry, NMR- and mass spectroscopy, elemental analysis, ozonolysis, and biological testing were used.

Conclusions
Roseofungin is an original natural highly efficient antifungal antibiotic. Roseofungin is amorphous powder, it decomposes at t>130°C; fluoresces in UV, is quite soluble in pyridine, dimethylformamide, acetic acid, lower alcohols, soluble in water-saturated butanol and wateracetone. UV-spectrum has 2 absorption maxima: at 263 nm and 363 nm. Roseofungin in concentrations of 0.5-12.5µg/ml inhibits the pathogenic agents of superficial and deep mycoses-trichophytosis, microsporia, candidiasis, cryptococcosis, sporotrichosis, chromomycosis, aspergillosis, and others. High virus-inhibiting activity of roseofungin has been established against various strains of influenza virus, Sendai paramyxovirus and Newcastle disease virus.
DEVELOPMENT OF NEW BIOLOGICAL PREPARATIONS OF THE “RIZOVIT-AKS” SERIES BASED ON NODULE BACTERIA OF LEGUMINOUS PLANTS CULTIVATED IN KAZAKHSTAN

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Background
Development of biological preparations based on active strains of indigenous nodule bacteria of leguminous plants improves their productivity and soil capabilities.

Objectives
Strains of nodule bacteria isolated from the rhizosphere of leguminous plants:

Methods
Biotechnological and microbiological methods were used in the study.

Conclusions
Modification of the culture media containing bean broth or yeast extract has been carried out, and effect of carbon and inorganic salt sources on the growth and biomass accumulation of indigenous strains of nodule bacteria from soybean, alfalfa, peas, chickpeas, lentils studied. The best carbon source for most strains of root nodule bacteria is sucrose at a concentration of 4-10.0 g/L. A number of salts (CaCO₃, MnSO₄, ZnSO₄, CoCl₂) stimulated the biomass accumulation of nodule bacteria strains up to 10¹¹ CFU/ml.

Application of biological preparations of the “Rizovit-AKS” series to inoculate the seeds of leguminous crops reduces the time required for passing phenophases by legumes: on average 2 to 3 days for soybean; 2 to 5 days for peas; up to 2 days for lentils, 2 to 4 days for alfalfa. Biopreparations increase the plant growth, green mass and number of nodules under all tested soil and climatic conditions of Kazakhstan. With the use of biopreparations, an increase in the yields of leguminous crops: for soybeans by 3.2 t/ha (34.4%), lentils – by 3.3 t/ha (41.3%); peas – by 8.3 t/ha (140.7%).
Background
Our current society is confronted with a series of global geopolitical, economic, energetic and climate crises. While there is no common solution for all these issues, there is an obvious main connection, the need for a novel renewable and environmentally friendly energy carrier. Hydrogen gas is considered an attractive alternative to fossil fuels with certain methods of biohydrogen production being able to utilize various low-priced industrial and agricultural wastes, thus coupling organic waste treatment with renewable energy generation.

Objectives

By designing a novel hybrid dark fermentation-photoheterotrophic degradation biohydrogen production process using industrial wastewater as a substrate we aim to resolve the major scientific and technical bottle-necks preventing the biohydrogen production systems to become economically feasible, bringing thus the technology closer to the industry.

Methods

A laboratory-scale bioreactor experimental setup was developed in order to identify the most suited microbial and micro-algal consortia, as well as the key metabolic pathways involved. In addition, an advance statistical design of experiments approach was used to identify the influence and interactions of key micro-environmental factors.

Conclusions

The obtained preliminary results prove that this novel system has the potential to produce considerably more H₂ yields compared to traditional systems as well as generate valuable oils through the micro-algal metabolic activity. In addition, such a novel technology could capture the generated CO₂ produced during the dark fermentation step, and used it by the micro-algal species together with the dark fermentation effluent.
Background

Hyaluronan (hyaluronic acid, HA) is a linear polysaccharide composed of alternating D-glucuronic acid (GlcUA) and N-acetylglucosamine (GlcNAc) linked by $\beta(1,4)$ and $\beta(1,3)$ glycosidic bonds. This macromolecule is found in various tissues of vertebrates, where it has different biological functions. Industrially, HA has been manufactured by fermentation of group C streptococci.

This polysaccharide is produced by processive synthase from the activated UDP-sugars (UDP-GlcA and UDP-GlcNAc). A positive correlation between molecular weight (MW) and concentration of UDP-GlcNAc was previously reported [1].

Objectives

To increase hyaluronan yield or its molecular weight by strengthening UDP-precursors biosynthesis.

Methods

Fermentations were carried out in fermentor Sixfors (Infors HT, Switzerland) with working volume of 300 ml. HA yield was determined by isopropanol precipitation. HA Mw was analysed by SEC-MALLS.

Conclusions

In order to increase yield or molecular weight of hyaluronic acid in Streptococcus zooepidemicus fermentation, cultivation medium was supplemented with GlcUA and GlcNAc. These molecules are involved in hyaluronan biosynthetic pathway.

Extracellular GlcNAc significantly increased both HA yield and molecular weight (35% and 21% respectively). HA yield positively correlated with GlcNAc concentration up to 1g/l. Higher concentrations had detrimental effect on HA production. No significant
effect on HA yield was observed for GlcUA addition.
EVALUATION OF BACTERIAL ENZYMES IN DEGRADATION OF TEXTILE DYES

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Background

Because of the ease of their synthesis and the greatest variety of colors, azo dyes are the largest class of synthetic dyes. Generally they are very recalcitrant to biodegradation. Thus, in the current study the biodegradation of an azo dye namely Remazol Blue (RB) by wastewater-isolated bacteria (*Bacillus megaterium*, *Micrococcus luteus* and *Bacillus pumilus*) has been investigated in media containing molasses as a carbon and energy source.

Objectives

The goal of this work was to identify decolorization and degradation properties of Remazol Blue dye by bacterial strains at batch scale.

Methods

Enzyme activity: The reaction mixture contained 400 μl of 50 mM sodium phosphate buffer (pH 7.0), 200 μl of the sample and 200 μl of Remazol Blue. The reaction was initiated by the addition of 200 μl NADH (final concentration 2 mM) and followed photometrically.

Bioremoval assays: Experiments were carried out at different (from 28.7 to 97.9 mg/L) dye concentrations at pH 7 throughout 3 days.

Conclusions

*Bacillus*-derived azoreductases were responsible for biodegradation of RB. The maximum azoreductase activity was obtained as 39.9 U/ml for *B. pumilus* at 2nd day. The removal yields of *B. pumilus* were 71.0%, 74.3%, 67.2% and 69.9% for 28.7, 57.9, 78.3 and 97.9 mg/L dye, respectively. There were also high azoreductase activities for *B. megaterium* at the first and second days of incubation time such as 37.4 and 33.6 U/ml while *M. luteus* did not show any azoreductase activity.
Background

Among many heavy metals, Cr(VI), Ni(II) and Cu(II) are the ones having many usage fields, and therefore wastewaters including them have to be treated.

Objectives

This study aimed to isolate Ni(II), Cu(II), Cr(VI) resistant bacteria from Ankara Stream (Turkey) and determining EPS production and bioremoval capacities of them under different conditions.

Methods

Experiments were done with bacterial cultures in different media (active sludge: AS; active sludge with molasses: MAS) including Ni(II), Cu(II), Cr(VI) at varied pH values (6-9). Heavy metals were determined spectrophotometrically at 340 nm for Ni(II); 460 nm for Cu(II); 540 nm for Cr(VI). EPS amount was also investigated.

Conclusions

Mixed bacterial cultures had higher bioremoval than the pure cultures. Six heavy metal-resistant strains were purified. Strain 5 had the highest Cu(II) (69.1%) and Cr(VI) (43.1%) removal under 25 mg/l heavy metal at pH 7. The same strain was the efficient bacterium in bioremoval and EPS production. It had EPS amount as 1.05 g/l at 25 mg/l Cr(VI), and 0.74 g/l at similar Cu(II) concentration. Especially mixed and also pure bacteria could be used as a biosorbent in treatment of heavy metals.
Background

The depletion of fossil fuels and the concerns about environment have led scientists to find alternative renewable energy sources. Bioethanol is the most widely used liquid biofuel at present among these alternative fuels. Feedstock costs are the major part of the process. Thus, considerable work has been performed toward production of bioethanol using various kinds of feedstocks such as starch rich agricultural wastes and cellulosic biomass.

Objectives

To demonstrate the utilization capacity of agricultural wastes (apple and carrot pomaces) as feedstocks for bioethanol production.

Methods

Feedstock preparation: Appropriate amounts of apple and carrot pomaces were mixed with distilled water and homogenized to obtain desired concentration of feedstock solutions. Different pretreatment methods were performed to find the most effective way to obtain fermentable sugars in pomaces.

Fermentation conditions: The yeast cells were precultured in YPG medium at pH 6. 10% yeast suspension was aseptically transferred to anaerobic fermentation medium which is prepeared by pomace sugar containing distilled water.

Biomass, sugar and ethanol determination: Yeast biomass was determined by measuring optical density. The sugars were determined by phenol-sulphuric acid method. The bioethanol concentration was analyzed using gas chromatography.

Conclusions

Higher ethanol production values were obtained from apple pomace rather than carrot. The pretreatments performing with acid were more effective than the ones
performed with base. The highest ethanol production was obtained when the apple pomace was pretreated with 0.25% (v/v) H$_2$SO$_4$. 
BORON REMOVAL FROM AQUEOUS SOLUTIONS USING RHODOTORULA MUCILAGINOSA AS A BIOSORBENT

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Background

Boron (B) is a naturally occurring element founding in earth’s crust, rocks, soil, and water. Although B is an essential element and used at many industries, it could be toxic for organisms in excess concentrations.

Objectives

In the present study we aimed to remove B from aqueous media via biosorption by Rhodotorula mucilaginosa, which was isolated from B-contaminated wastewater.

Methods

Batch culture experiments including pH (4-7), increasing B (15-20 mg/L), biomass concentrations (1-4 g/L) and different biosorption methods (wet, dried and formaldehyde-treated cells) were carried out at 25±2 °C, at 100 rpm for 120 min. Boron concentrations were determined spectrophotometrically by measuring the absorbance at 585 nm with using carmine.

Conclusions

According to the data obtained, the optimum pH was found as 6 for the maximum B removal. Under 15.35 mg/L B, the highest biosorption was determined as 23.78%. Biosorption was increased with increasing biomass and the maximum yield (31.92%) was found at 4 g/L biomass. Wet cells removed B more effectively than formaldehyde-treated or dried cells and the yields were 23.78%, 19.31% and 12.07%, respectively. It can be concluded that R. mucilaginosa might be an effective biosorbent in B removal processes.
A NOVEL PATHWAY FOR ITACONIC ACID PRODUCTION IN USTILAGO MAYDIS

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Background

Itaconic acid is a promising bio-based platform chemical for the production of pharmaceuticals, adhesives and polymers. *Ustilago* is a promising fungal host for the production of itaconic acid. Contrary to established filamentous itaconate producers, *Ustilago* grows unicellulary. This has distinct process advantages relating to morphology control, viscosity and aeration. However, the itaconate production yield, titer and rate of *Ustilago* are below that of the commercial production hosts. In order to optimize itaconate production in *Ustilago*, detailed biochemical knowledge is needed.

Objectives

To characterize the itaconate production pathway in *Ustilago* in order to enable metabolic engineering.

Methods

An itaconate gene cluster was identified in *Ustilago*. The function of the cluster genes and their encoded proteins was characterized by knockout and overexpression. The activity of two essential catalytic proteins was determined *in vivo* by permeabilized cell assays and *in vitro* by *E. coli* expression and purification.

Conclusions

The genes *tad1* (trans-aconitate decarboxylase), *itp1* (Major Facilitator Superfamily extracellular itaconate transporter), *adi1* (aconitate-Δ-isomerase), *mtt1* (mitochondrial tricarboxylate transporter), and *ria1* (transcriptional itaconate regulator) are involved in the itaconate biosynthesis and possibly its further conversion.

In contrast to the known itaconate biosynthesis pathway of *Aspergillus terreus*, *Ustilago*’s itaconate production proceeds first via an isomerization from cis- to trans-aconitate, followed by decarboxylation of the trans-aconitate. First metabolic engineering attempts were successful, enhancing *U. maydis*’ itaconate production twofold by overexpression of *ria1* or *mtt1*. This work lays the foundation for further
optimization of *U. maydis*’ itaconate biosynthesis and is therefore a further step towards industrial application of this promising biocatalyst.
Background
Filamentous fungi are extensively studied due to their exoenzyme producing ability. Since lipases are effectively utilizable in various industrial processes, special attention is paid for these enzymes. Development of environmentally friendly technologies requires lipases able to catalyze the rapid synthesis and translocation of esters. Isolation and analysis of such enzymes contributes to the knowledge of these catalytic processes, and provides data on new lipases with potential biotechnological interest.

Objectives
A *Rhizopus stolonifer* isolate has been identified as high-yield lipase producer in our previous tests. Further studies revealed that the crude enzyme has significant transesterification activity. Therefore, our present work has focused on the purification and characterization of this *R. stolonifer* lipase.

Methods
Isolate was grown on liquid medium containing wheat bran. To purify the enzyme, after filtration and centrifugation, ammonium sulphate precipitation, size-exclusion and ion-exchange separations were performed. Biochemical characterization assays including substrate specificity, temperature and pH tolerance studies, and examination of the effect of some ions and organic solvents on the activity were carried out.

Conclusions
The optimal temperature and pH for the activity were about 50 °C and pH 5.0, respectively. The lipase has broad substrate specificity because it has effectively hydrolyzed the substrates containing fatty acids from C6 to C16. Enzyme activity was stimulated by 5% ethanol, butanol and propanol. Results showed that certain alkanes could stabilize the active conformation of the enzyme. The isolated *R. stolonifer* lipase exhibited high esterification and transesterification activity at 40-50 °C. This research was supported by the Hungarian Research Fund (OTKA PD 112234).
Background

Recent years, there is a fast-growing interest on the catalytical activities display by lipase enzymes, especially focusing on their synthetic activity in organic solvents. Translocation and synthesis of ester linkages result various aroma-, phenyl- and alkyl esters and different polymers. Many of these compounds have important role in the food-, pharmaceutical-, and oleochemical industries as well as in the biodiesel production. In these industries, development of modern environmentally friendly biotechnological technologies requires the exploration and utilisation of new microbial enzymes.

Objectives

Lipases can be produced by certain Mucoromycotina fungi as well, but only some of these enzymes have been tested for their synthetic activity in organic media. In this work, we evaluated the esterification and transesterification capacities of Rhizomucor, Rhizopus, Mucor and Mortierella lipase enzymes through alkyl ester synthesis, and we characterise of their fatty acid preference during these reactions.

Methods

Reactions were carried out in non-aqueous condition using lyophilised crude enzyme preparations. Transesterification was studied between p-nitrophenyl-palmitate and ethanol, while esterification reactions were examined with different chain length fatty acids and methanol. The effect of the reaction time on the ester production was also evaluated. Synthetic esters produced were detected by a GC-FID technique.

Conclusions

Each lipase catalysed the transesterification significantly faster than esterification. High ethyl-palmitate production was reached by lipases from Rhizopus stolonifer and Mucor corticola strains. During esterification reactions, the tested enzymes showed increased alkyl ester synthesis in the presence of medium-chain fatty acids. This
research was supported by the TÁMOP-4.1.1.C-12/1/KONV-2012-0012 and the Hungarian Research Fund (OTKA PD 112234).
NOVEL SMALL-SCALE AND ENZYMATIC TEST SYSTEMS TO ANALYSE FULL OPERATING BIOGAS PLANTS
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Background

One important renewable energy source is biomass from energy crops and organic waste that is used for the production of biogas. Biogas mainly consists of CH₄ and CO₂ and is generated by anaerobic fermentation of organic matter.

Objectives

For a more precise and targeted optimization it is crucial to understand the biological processes during biogas production.

Methods

Using authentic samples from full operating biogas plants it was possible to develop batch and continuous reactors mirroring the conditions and the performance of the full-scale biogas plant. With these test systems it was possible to observe effects of different substrates for syntrophic bacteria and for methanogenic archaea on biogas formation.

Furthermore, the development of novel sensors in biogas plants to quantify the activities of key enzymes is desirable. A potential candidate is the heterodisulfide reductase (Hdr). Methanogenesis finally results in the formation of the heterodisulfide, which is regenerated by the Hdr. In aceticlastic methanogens the Hdr is tightly bound to the cytoplasmic membrane but in hydrogenotrophic methanogens it is located in cytoplasm.

Conclusions

Most biogas sludge samples were stimulated by the addition of ethanol or acetate, which leads to increasing methane production by 35-126 %. These results indicate that syntrophic ethanol oxidation and aceticlastic methanogenesis were not rate limiting in routinely operating biogas plants.
Our test system allows a quantification of hydrogenotrophic/aceticlastic methanogens in biogas sludge based on Hdr activity. In summary 26% of total Hdr activity was found in the membranes while the cytoplasmic fraction contained 74% of total activity.
Background
White rot fungi are thought to degrade lignin using AA2 enzymes like lignin peroxidase, manganese peroxidase (MnP) and versatile peroxidase. It is unclear if S. commune is a white rot or brown rot fungus. This basidiomycete is a poor lignin degrader. No genes encoding AA2 enzymes are found in its genome, while it is enriched in AA9 genes. This would place S. commune in between brown and white rot fungi. We here introduced two A. bisporus mnp genes in S. commune.

Objectives
Introduction of A. bisporus MnPs in S. commune and its effect on the lignin degrading capacity of S. commune.

Methods
qPCR verified expression of these genes in S. commune. Decoloration of RBBR by mnp expressing S. commune strains indicated production of active MnP. This was confirmed using the MBTH/DMAB assay. Addition of hemin to standing cultures increased MnP activity 5-fold. Pyrolysis analysis indicated that a S. commune strain expressing both MnPs of A. bisporus degraded lignin in birchwood.

Conclusions
A. bisporus MnPs were succesfully introduced and expressed in S. commune. The G subunit of lignin seems to be preferentially degraded by S. commune MnP expressors. The ligninolytic pathway of litter degrading fungi can be reconstituted in S. commune.
MICHAEL-TYPE ADDITION USING 4-OXALOCROTONAT TAUOMERASE: BIOPROCESS IMPROVEMENT

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Background

Enzyme 4-oxalocrotonate tautomerase (4-OT) encoded by xylH gene is a part of aromatic compounds degradation pathway in Pseudomonas putida mt-2. Due to its promiscuous activity, 4-OT was described to catalyse Michael-type addition of acetaldehyde to β-nitrostyrene, the isomerisation of cis-nitrostyrene to trans-nitrostyrene as well as aldol condensation and dehydratation as free enzyme. Whole cell system based on recombinantly expressed 4-OT has been developed and shown to be effective biocatalyst of asymmetric Michael addition of acetaldehyde to β-nitrostyrene.

Objectives

The objective of this study was to improve biocatalytic production of 4-nitro-3-phenylbutanal employing two strategies: biocatalyst immobilization and product recovery using Amberlite XAD-2 polymeric adsorbent.

Methods

Whole cell biocatalyst was immobilized in 4% alginate gel. Capsules of 1 and 2 mm in diameter were made. Liquefaction of the capsules core was performed by addition of sodium citrate. Depletion of β-nitrostyrene was monitored by reduction of absorbance at 320 nm. Product was recovered from aqueous reaction buffer by addition of XAD-2 beads, followed by elution in minimal volume of ethylacetate.

Conclusions

Biocatalytic production of 4-nitro-3-phenylbutanal was improved by whole-cell immobilization in alginate capsules. Capsules were used in 12 reaction cycles, and retained 20% of the activity after two months of storage. Bioprocess was further improved by using XAD-2 beads for improved product recovery. With this
modification, the amount of organic solvent was reduced 20-fold in comparison to previously reported method making this process more environmentally friendly.
COMPARATIVE GENOMIC ANALYSIS OF SOLVENT PRODUCING CLOSTRIDIUM ACETOBUTYLICUM AND CLOSTRIDIUM BEIJERINCKII STRAINS

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Background
The clostridial acetone-butanol-ethanol (ABE) fermentation pathway is complex and represents one of the oldest industrial fermentations. Interest in the production of solvents from lignocellulosic sources has increased in recent years. However, low production yields, end-product inhibition and phage infections remain as disadvantages in the ABE fermentation.

Objectives
Genetic and metabolic engineering strategies leading to a stable, high-yielding process require prior genomic exploration of diverse solventogenic Clostridial strains.

Methods
Ten strains of C. acetobutylicum and C. beijerinckii were selected from batch fermentation experiments according to their solvent yields. Genomes and plasmids (when present) were sequenced using the Illumina HiSeq platform. Complete genomes were obtained through de novo and reference-guided assembly comparison with coverages of ~ 100x. Annotation was performed using RAST (http://rast.nmpdr.org). Key enzymes for solvent production where mapped in the genomes and/or plasmids to allow variant detection and comparison in future RNA-seq experiments. Raw Illumina reads were used as input for CRISPR structure detection using the Crass software (http://ctskennerton.github.io/crass/). Putative CRISPR fragments were assembled and queried against the CRISPR database (http://crispr.u-psud.fr/crispr/). Complete CRISPR structures were detected in 7 out of 10 genomes (average 2.5 CRISPR structures per strain), and repeat/spacer units were detected in all 10 sequenced strains. An alignment-based approach revealed the existence of uneven distribution of types (11 detected) across their genomes.

Conclusions
In combination, our results provide a sound basis for directed generation of engineered Clostridial solventogenic strains, with the potential to re-launch the ABE fermentation as a competitive industrial fermentation.
REROUTING OF CARBON FLUX TOWARDS GLYCEROL PRODUCTION IN YEAST SACCHAROMYCES CEREVISIAE

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Background
Currently there is an interest in development of microbial strains effectively converting cheap feedstocks to glycerol. Facultative anaerobic yeast Saccharomyces cerevisiae can be a good platform for development of recombinant strains overproducing glycerol under low-aeration conditions. In S. cerevisiae glycerol is synthesized from dihydroxyacetone phosphate by the action of glycerol-3-phosphate dehydrogenase (Gpd1) and glycerol-3-phosphate phosphatase (Gpp2). Other road of dihydroxyacetone phosphate catabolism occurs by its isomerization to glyceraldehyde-3-phosphate with triose phosphate isomerase (Tpi1), resulting to ethanol formation.

Objectives
We aimed to construct recombinant strains with reduced Tpi1 activity and increased Gpd1 and Gpp2 activities.

Methods
Homologous recombination was used for partial substitution of TPI1 gene promoter region with selective marker. Obtained strains contain 100, 50 or 25 base pairs of native TPI1 gene promoter before TPI1 ORF, and revealed corresponding sequential decreases in Tpi1 activity. Multicopy integration module was used for expression of hybrid GPD1-GPP2 ORF (encoding artificial fusion of both enzymes) under the control of strong constitutive promoter of the alcohol dehydrogenase gene.

Conclusions
Recombinant S. cerevisiae strains with 50 or 25 bp version of TPI1 promoter revealed up to 2 times increase in glycerol production as compared to the WT strain. Recombinant strains expressing GPD1-GPP2 fusion produced approximately 4 times more glycerol than that of WT strain. Combination of both approaches resulted to 5-fold increase of glycerol production as compared to the WT strain.
METHANOTROPHIC EXTREMOPHILES: FROM UNUSUAL PHYSIOLOGY TO NEXT GENERATION BIOCATALYSIS

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Background
The long-term goal of this research is to develop a robust microbial platform for conversion of greenhouse gases (CH\textsubscript{4} and CO\textsubscript{2}) into valuable chemicals as an alternate approach to widespread (bio)gas-flaring.

Objectives
The main objective of this project is to generate a comprehensive vision of methane oxidation in a model methanotroph (\textit{Methylomonas alcaliphila} 20Z) using modern systems biology approaches.

Methods
A multi-tiered systems-level approaches (genomics/mutagenesis, transcriptomics, metabolomics and fluxomics) were used to validate the metabolic network in \textit{M. alcaliphila} 20Z grown on methane and methanol during both, steady state and transitions from restricted to active utilization of C1-substrate.

Conclusions
Comparisons of different growth conditions highlighted functions important for methane oxidation and for the re-activation of methane oxidation machinery. Complete genome sequences of the methanotrophic strain enabled genome-wide metabolic reconstruction. The first stoichiometric flux balance model of methane utilization was constructed and validated. A new vision of the core methanotrophic functions will be presented. The majority of the data (methane oxidation parameters, growth yield, mutant phenotypes) suggest that \textit{direct coupling} between methanol and methane oxidation is the most compelling mode of methane activation. The reconstruction of spatial organization of the metabolic network coupled with the computational modeling enables a mechanistic understanding and systematic analysis of the interplay between core methane oxidation processes and overall performance of the biological system. New data were used to improve conversion of the C1-substate into values-added chemicals and fuels.
OPTIMIZATION OF LEVAN PRODUCTION BY BACILLUS LICHENIFORMIS ANALYSED BY THE RESPONSE SURFACE METHOD

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Background

Levan is a fructose based exopolysaccharide produced by microorganisms as an energy reserve and defense. It has present low viscosity, high solubility in water, biocompatibility and other properties that have industrial and medicine applications. Levan-type polymers have been reported as hypocholesterolemic, cell-proliferating, anti-inflammatory and antitumour agent, an immune modulator, thickener, stabilizer, flavors, and a aroma carrier [1].

Response surface methodology (RSM) is the most popular optimization method used in recent years because the application of statistical designs for experiments and its modeling defines the effect of various factors and its interaction [2].

Objectives

The aim of this study was to optimize fermentation medium composition for levan production by Bacillus licheniformis strain.

Methods

Basic medium for levan production contains sucrose as carbon source, yeast and beef extract as nitrogen source, and its pH level is important for excretion of polysaccharide. In this work it has been shown that high levan production by the studied strain could be achieved with ammonium chloride as a sole nitrogen source. Influence of three independent variables (pH, sucrose and ammonium-chloride concentration) was investigated using Box-Behnken statistic design.

Conclusions

Based on response surface model that we obtained, in medium that contain 200 g/L sucrose, 1.44 g/L ammonium-chloride and pH 7.5 maximum concentration of levan (41.29 g/L) is reached, which was in agreement with experimental data.

References:

MOLECULAR BREEDING OF YEAST CO-DISPLAYING ENDO- AND EXO-TYPE ALGINATE LYASES TO COMPLETELY UTILIZE MACROALGAE

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Background

Brown macroalgae is characterized by a large size and high productivity without requiring arable land, irrigation water, or fertilizer. It is a raw material suitable for productivity of bioenergy and biochemical productions. However, alginate, which is the main component of brown macroalgae, is an undegradable polysaccharide with poly (M)-, poly (G)-, and poly (MG)-blocks composed of α-l-guluronic acid (G) and its C5 epimer, β-d-mannuronic acid (M), as monomeric units.

Objectives

In order to degrade alginate polymers effectively, both endo- and exo-type alginate lyases are necessary. Therefore, we co-displayed endo- and exo-type alginate lyases from the alginate-degrading marine bacterium Saccharophagus degradans [1] on the cell surface of Saccharomyces cerevisiae with our original method, cell surface engineering technology [2].

Methods

The alginate lyases from S. degradans were produced as fusion proteins with cell wall anchoring domain of α-agglutinin in S. cerevisiae. The production and localization of alginate lyases on the yeast cell surface were confirmed by immunofluorescence microscopy. The enzyme activities of the displayed alginate lyases were measured by using the dinitrosalicylic acid method.

Conclusions

The results suggested that the engineered yeasts co-displaying endo- and exo-type alginate lyases could lead to efficient production of monomeric units from alginate polymers [3]. This system is a unique easily feasible to utilize any biomass.

References

SCREENING AND MOLECULAR CHARACTERIZATION OF FUNGI CAPABLE OF LACCASE PRODUCTION FROM DIFFERENT FOREST ECOSYSTEMS OF MICHOACÁN STATE, MEXICO.
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Background
Contrary to temperate regions, knowledge on laccase producing fungal species from ecosystems allocated at neotropical regions is still scarce. Search for ligninolytic fungi in these ecosystems can provide laccases with novel biotechnological potential.

Objectives
The goal of this study was to explore novel fungal species with laccase activity from different forest ecosystems.

Methods
Fungal species were isolated from soil and basidiocarp samples collected from five different geographic regions with dominant vegetation of Pinus spp., Abies spp., and Quercus spp. Extracellular laccase activity by fungal isolates were analyzed by guaiacol assay. Identification of laccase positive fungi was carried out by sequence comparison and phylogenetic analysis of internal transcribed region (ITS1F-5.8S-ITS4B) of rDNA gene with reference taxa.

Conclusions
A total of thirty (30) laccase producing fungi, representing twenty one (21) species of fifteen (15) genera, were purified. Fifteen strains (15) with high laccase activity of basidiomycetes were identified in thirteen genera i.e., Bjerkandera, Corticiaceae, Coriolopsis, Echinodontium, Ganoderma, Hexagonia, Irpex, Limonomyces, Psathyrella, Peniophora, Phlebiopsis, Trametes and Trichaptum. Seven (7) Trichoderma and only one Penicillium species were found as low laccase producer, isolated from different soil and basidocarp samples. Trichoderma tomentosum was recorded as most isolated laccase producing isolate from different soil samples.
followed by *T. atroviride* from different regions. However, *Penicillium pinophilum* was only laccase producing species of genus *Penicillium*, isolated from soil sample of *Pinus* spp., and *Quercus* spp., dominant area.
COMPLEX PROBIOTIC FEED SUPPLEMENT BASED ON CONSORTIUM OF MICROBIAL STRAINS FOR INDUSTRIAL POULTRY BREEDING

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Background
Large-scale poultry breeding often provokes outbreaks of various bacterial, viral and mixed infections leading to fowl loss, while shortage of vitamins and hardly digestible feed ingredients reduces nutritive value and downgrades commodity outlook for meat and egg products.

Objectives
Aim of this study – development of technology for production and application of complex probiotic feed additive possessing antimicrobial, antiviral and antioxidant action allowing to prevent and cure fowl diseases, to raise nutritive value and digestibility of fodder, to promote poultry resistance to stress factors.

Methods
Microbiological, biochemical, genetic engineering, physical-chemical methods were used in this research: engineering a strain-producer of chicken leukocyte alpha-interferon, submerged fermentation of microorganisms, lspectrometry, freeze-drying and etc.

Conclusions
To broaden the activity spectrum of probiotic feed supplement several bacterial species capable to complement each other were applied. Active principle of the feed additive — strain Bacillus subtilis produces surfactins, polyene and peptide compounds displaying antagonistic activity against strictly and facultatively pathogenic microbial species colonizing avian gastrointestinal tract. It was found that Bacillus subtilis bacteria generate cellulases, proteases and other enzymes facilitating assimilation of fodder. Enterobacteria Pantoea agglomerans synthesize carotenoid pigments – sources of antioxidant activity and precursors of vitamin A in the body. The protein of chicken leukocytic alpha-interferon synthesized by Escherichia coli strain possesses pronounced non-specific antiviral and immune-stimulating activities.

Technology of producing complex probiotic feed supplement in dry form for poultry was developed.

Biological trials in vitro and in vivo demonstrated its safety and high efficiency in feeding experiments with broilers and hens.
EXPLORING THE COMPLEX MICROBIAL COMMUNITY OF ACTIVATED SLUDGE SAMPLED FROM AN INDUSTRIAL TANNERY WASTEWATER TREATMENT PLANT BY ILLUMINA MISEQ.

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Background

Background: In the frame of the BIOSUR project (LIFE11 ENV/IT/075), a one-year monitoring of an industrial tannery wastewater treatment plant was performed, both from chemical-physical and biological points of view.

Objectives

Objectives: This study is conducted to investigate the evolution of the bacterial community of activated sludge in the same treatment plant during 2013.

Methods

Methods: Bacterial diversity is analyzed by the use of the Illumina MiSeq platform for sequencing hypervariable regions V3-V4 in the 16S rRNA genes. Bioinformatic analysis of the obtained 12,429,502 paired-ends reads is performed using the QIIME pipeline. The assembled 4,039,284 contigs are binned into 5,156 Operational Taxonomic Units (OTUs) that are assigned to taxonomic groups by the use of the Silva 111 database.

Conclusions

Conclusions: Preliminary results indicate that the bacterial diversity of active sludge is maintained at very high levels throughout the whole year. The dominant bacterial phylum is Proteobacteria, which accounts for 40%. Multivariate analysis shows that there is a discontinuity between two groups of samples. The first group, corresponding to samples collected from January to July 2013, is detached from the second group, corresponding to samples collected from September, after a period of drastic decrease of work in tanneries. The Shannon diversity index ($H$), used to calculate the diversity of bacterial communities, indicates a slight increase in the diversity level in the second period. Further analysis will better clarify the complexity
of the microbial community and the influence of specific environmental parameters on its structure and evolution.
IMPROVEMENT OF GLYCEROL PRODUCTION BY DELETION OF ADH1 GENE AND OVEREXPRESSION OF GPD1 AND GPP2 GENES IN SACCHAROMYCES CEREVISIAE

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Background

Glycerol is widely used in cosmetics, food, tobacco, pharmaceutical, leather and textile industries. In addition, it is considered as a cheap raw material for microbial fermentation. That is why the construction of yeast strains-producers of glycerol became an actual objective for modern metabolic engineering. Our strategy comprised the deletion of ADH1 gene, encoding alcohol dehydrogenase in S. cerevisiae and overexpression of both glycerol-3-phosphate dehydrogenase and glycerol-3-phosphate phosphatase genes (GPD1, GPP2) in order to redirect the carbon flux to glycerol instead of ethanol formation.

Objectives

Our goal was to design recombinant strains with deletion of ADH1 gene combined with increased Gpd1 and Gpp2 activities.

Methods

Homologous recombination was used for deletion of ADH1 gene with selective marker. The integrative module was used for expression of hybrid GPD1-GPP2 ORF (encoding artificial fusion of both enzymes) under the control of strong constitutive promoter of the alcohol dehydrogenase gene.

Conclusions

Glycerol concentration during the course of fermentation was approximately 4.7-fold enhanced in a Δadh1_GPD1-GPP2 recombinant strain if to compare with BY4742 (2.57 g/L), reaching 12.14 g/L (0.168 g/L/h) on the third day of fermentation. Furthermore, the glycerol production in a recombinant Δadh1_GPD1-GPP2 strains with deletion of ADH1 gene and overexpression of GPD1 and GPD2 genes was nearly 1.2-fold increase in comparison with Δadh1 strain, which contained only the disruption of ADH1 gene.
MODIFICATION OF METABOLIC PATHWAY FOR GLYCEROL PRODUCTION IN YEAST SACCHAROMYCES CEREVISIAE

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Background

Several processes are known for the microbial synthesis of glycerol from carbohydrates that use osmotolerant yeasts, algae and a number of bacteria. However, all these utilize aerobic conditions, so the glycerol production demands air or oxygen purging throughout the fermentation, which considerably increases production costs. Therefore the development of the \textit{Saccharomyces cerevisiae} strains capable of efficient glycerol production from glucose under anaerobic conditions is of great interest.

Objectives

We aimed to increase glycerol production by derepression of \textit{ILV2} in part with \textit{GPD1-GPP2} fused genes.

Methods

Homologous recombination was used for construction of the truncated version (deficient 5'–165 bp and lacking a mitochondrial targeting signal) of the yeast \textit{ILV2} gene, encoding for acetolactate synthase with a strong constitutive promoter. Obtained strains shown increased acetolactate synthase activity. Multicopy integration module was used for expression of hybrid \textit{GPD1-GPP2} ORF (encoding artificial fusion of glycerol-3-phosphate dehydrogenase and glycerol-3-phosphate phosphatase) under the control of strong constitutive promoter of the alcohol dehydrogenase gene.

Conclusions

Overexpression of the truncated version of \textit{ILV2} with presumably cytosolic localization, strongly activates glycerol production under anaerobic conditions. Preliminary results possessed that the glycerol production by recombinant strains co-expressing \textit{ILV2} and \textit{GPD1-GPP2} was increased as compared to the wild type strain and strain expressing solely \textit{ILV2}.
AN ANTIFUNGAL PEPTIDE FROM ACTINOBACTERIA (STREPTOMYCES SP. TKJ2) : ISOLATION AND PARTIAL CHARACTERIZATION

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Background
Actinobacteria are of special biotechnological interest since they are known to produce chemically diverse compounds with a wide range of biological activity. This distinct clade of Gram-positive bacteria include some of the key antibiotic producers and are also sources of several bioactive compounds.

Objectives
The aim of our study is the purification and the partial characterization of an antifungal protein produced by Streptomyces sp. TKJ2 isolated from Algerian soil.

Methods
A newly filamentous bacteria was recovered from Tikjda forest soil (Algeria) for its high antifungal activity against various pathogenic and phytopathogenic fungi. The nucleotide sequence of the 16S rRNA gene (1454 pb) of Streptomyces sp. TKJ2 exhibited close similarity (99 %) with other Streptomyces 16S rRNA genes. Antifungal metabolite production of Streptomyces sp TKJ2 was evaluated using six different fermentation media. The antifungal protein produced by Streptomyces sp. TKJ2 on PCA medium has been purified by ammonium sulfate precipitation, SPE column chromatography and high-performance liquid chromatography in a reverse-phase column.

Conclusions
The UV chromatograms of the active fractions obtained at 214 nm by NanoLC-ESI-MS/MS have different molecular weights. The F20 Peptidique fraction obtained from culture filtrat of Streptomyces sp. TKJ2 precipitated at 30% of ammonium sulfate was selected for analysis by infusion ESI-MS which yielded a singly charged ion mass of 437.17 Da.
Background

In recent years public emphasis has been focused on the problems of rapid arable land degradation: structural deterioration of ploughed layer, reduced humus content, rising acidity – all leading to loss of soil suppressive potential. Tillage quality downward trend requires urgent measures for preservation and remediation of agricultural resources. A vital method to promote soil fertility and harvests of cultivars envisages ploughing-in of straw from crop rotation precursor treated by phytoprotective microbial preparation accelerating decomposition of plant residues and regeneration of soil biocenoses.

Objectives

Aim of this study was to assemble a consortium of microbial species distinguished by phytoprotective and hydrolytic potential.

Methods

Various investigation methods were used: microbiological, biochemical, molecular-genetical.

Conclusions

500 microorganisms isolated from diverse natural sources were screened to yield 33 antagonists of crop soil pathogens. 3 antagonistic bacterial strains were found to show the highest levels of antimicrobial and cellulolytic activities. The selected variants were characterized by the following properties:

- growth suppression of phytopathogens belonging to genera *Fusarium*, *Botrytis*, *Colletotrichum*, *Chaetomium*, *Pseudomonas*, *Xanthomonas*;

- increased seedling/root size and crude weight;
- production of hydrolytic enzymes – endo-1,4-β-glucanase and xylanase;
- 10-12% reduction of straw fiber content by 1 month;
- upgrading soil cellulytic activity;
- stimulating effect on major ecologo-trophical microbial groups in soil (ammonifying, oligonitrophilic, cellulytic, micromycetes).

The obtained data evidence attractive application prospects of microbial consortium as active ingredient of biopreparation to trying down infectious background, to restitute soil microbial cenoses and to promote plant growth.
SULFUR OXIDIZING BACTERIA COMMUNITY: CHARACTERIZATION AND MONITORING IN A REACTOR FOR SULFIDE REMOVAL FROM INDUSTRIAL WASTE.

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Background

Tannery wastewater contains high concentrations of pollutants, mostly sulfides and sulfates, which are usually chemically removed by expensive and inefficient methods. An innovative oxidizing bioreactor for biological removal of sulfide from gaseous industrial waste streams is developed and set up in the frame of the BIOSUR project (LIFE11 ENV/IT/075).

Objectives

This is achieved thanks to the development of a specialized chemolithoautotrophic sulfur-oxidizer biomass, which has been studied in order to elucidate its evolution.

Methods

The involved microbial community is selected in a side reactor by incremental addition of sulfides in selective conditions: low pH (2-4) and a concentration of sulfides of 2-4 mg/l. Such a specialized biomass was characterized and monitored by means of T-RFLP fingerprinting, clone-library construction of 16S rRNA coding genes and isolation in pure culture.

Conclusions

Preliminary results show the evolution of a selected, specialized sulfur-oxidizer biomass from the onset of the reactor to the stationary phase. In this scenario, about 65% of screened clones are represented by bacteria whose 16S rRNA gene sequence shows a similarity higher than 97% with that of members of the genus Halothiobacillus, well-known as Sulfur Oxidizing Bacteria (SOB). The same results has been also confirmed by isolation in SOB selective medium. Finally, obtained data have been comparatively analyzed in order to monitor microbial community evolution from the start-up throughout the end of the experiment.
IMPROVEMENT OF GLUTATHIONE PRODUCTION IN METHYLOTROPHIC YEAST HANSENULA POLYMORPHA

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Background

Glutathione (γ-L-glutamyl-L-cysteinyl-glycine; GSH) is the most abundant non-protein thiol compound of the most living organisms that protects cells from environmental and oxidative stresses. This tripeptide is an important natural compound for medicine, biotechnology and food industry. Microbial GSH overproduction is limited by mechanisms of feedback inhibition of γ-glutamylcysteine synthetase (GCS), the first and rate-limiting enzyme of GSH biosynthesis, by the end product. In addition the expression of gene coding for GCS is repressed by GSH. Thermotolerant methylotrophic yeast H. polymorpha is considered as promising organism for genetic modification and design of competitive GSH producer.

Objectives

To obtain a competitive producer of GSH in methylotrophic yeast H. polymorpha by overexpression of modified GCS with eliminated repression-inhibition mechanisms normally exerted by GSH.

Methods

Standard methods of yeast molecular genetics, yeast biochemistry and physiology have been used. The modified versions of GSH2 gene were obtained by error prone PCR.

Conclusions

The modified versions of GSH2 gene obtained by error prone PCR were cloned under the control of strong constitutive promoter of glyceraldehyde-3-phosphate dehydrogenase in replicative plasmid pYT3. Selected transformants were analyzed for their resistance to different prooxidant agents (1,2,3-triazole, diethylmaleate, ethionine) as compared to strains carrying unmodified GSH2 gene. Selected strains possessed growth on medium supplemented with triazole. Strains providing more intensive growth on the selective medium revealed higher GSH accumulation as compared to strains carrying unmodified GSH2 gene, indicating the reduction of Gsh2 feedback inhibition. In this work the selection scheme providing generation of
Gsh2 insensitive to feedback inhibition was developed.
HOW TO INTERPRET GENOMIC UNITS INSTEAD OF COLONY FORMING UNITS: A CASE STUDY WITH LEGIONELLA PNEUMOPHILA

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Background
Legionella pneumophila has been proven to be the causative agent of the potential deadly Legionnaires’ disease since 1977. In most of the European countries, the reference method for enumeration of Legionella in water is culture (ISO 11731). Due to the long incubation time of this method, more and more laboratories use quantitative PCR (qPCR). Since 2012 the normalization for the Legionella qPCR increases considerably: the ISO 12869 was developed, there exist different interlaboratory tests and reference standards. Different reports showed that this technique is accurate enough to be implemented as a reference method. One remaining obstacle is the data interpretation: the culture is expressed in colony forming units (CFU) and the qPCR in genomic units (GU). Although there is a correlation between both units, there is no consensus on how GU can be translated to CFU.

Objectives
Based on a database of hot and cold sanitary water samples and a literature screening an action level was defined for the monitoring of Legionella in sanitary water.

Methods
200 water samples were analyzed by culture and qPCR following the ISO 11731 and ISO 12869 respectively. A correlation analysis was performed and the variance was calculated by the logarithmic differences between the culture and the qPCR results. Together with the results from literature, action levels for the Legionella monitoring are proposed.

Conclusions
Although the culture has been widely used for many years, one should consider the advantages of qPCR over culture and acknowledge its suitability as a reference method.
A novel beta-galactosidase from Arthrobacter sulfonivorans: purification, characterization and gene cloning

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Background

Beta-galactosidase (EC 3.2.1.23) also known as lactase is the enzyme that typically catalyzes hydrolysis of beta-1,4-D-galactosidic linkages in beta-D-galactosides, including disaccharide lactose, with glucose and galactose as end reaction products. This enzyme is able to catalyze synthesis of oligosaccharides, in particular galactooligosaccharides via galactosyl transfer reaction.

Arthrobacter sulfonivorans beta-galactosidase of unique for prokaryotes extracellular localization may find application in food industry for manufacturing lactose-free dairy products and in pharmacology as bioactive principle of medicines prescribed for patients suffering from lactase deficiency.

Objectives

The study was aimed at cloning of the gene encoding A. sulfonivorans beta-galactosidase, purification and characterization of the enzyme.

Methods

Fast protein liquid chromatography, electrophoretic analysis, enzyme assay, PCR, DNA sequencing.

Conclusions

A novel extracellular beta-galactosidase from A. sulfonivorans was recovered with an overall 207-fold purification, a 7.7% yield and specific activity 16 300 U·mg⁻¹ protein. The subunit molecular mass of the enzyme determined by SDS-PAGE analysis equalled 125 kDa. It was found that the enzyme displays pi 5.35, prefers ortho-nitrophenyl-beta-galactoside as substrate (Km 27 mM) and shows maximum activity at 40°C and at pH 7.5–9.5.

The beta-galactosidase gene was isolated from the genomic DNA library of A. sulfonivorans, sequenced, cloned and deposited in the GenBank database under
accession number KM277894.1. It was established that the gene carries an open reading frame consisting of 3132 bp (1043 amino acids) and encodes beta-galactosidase referred to Glycosyl Hydrolase Family 2 (CAZy database).
FUNCTIONALITY ANALYSIS OF STRUCTURAL DOMAINS FROM GD-95 LIPASE BY SITE SPECIFIC AND RANDOM MUTAGENESIS

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Background

The rapid evolution of bioconversion and ecotechnology strongly increases the focus on the enzymes which possess novel properties. One of the most interesting enzymes are lipases from *Geobacillus* bacteria, because they can be active at extreme conditions. The ability to manipulate these lipases depends on the knowledge about the importance on the functionality of their individual domains and amino acids.

Objectives

Determination of influence conservative amino acids located at C-terminal end on activity GD-95 lipase without 10 C-terminal amino acids (GD-95-10) and construction new GD-95 lipase variants using error-prone PCR.

Methods

Lipase from *Geobacillus* sp. 95 without 10 C-terminal acid was analyzed by Ala scanning mutagenesis. New variants of recombinant GD-95 lipase were created using several error-prone PCR (epPCR). Ala mutants and GD-95 lipase variants after epPCR were cloned into pTZ57R/T and pET-21c(+) vectors. The recombinant proteins were expressed in *Escherichia coli* BL21(DE3) and purified using affinity chromatography. The activity and thermostability analysis of new lipases was performed spectrophotometrically.

Conclusions

GD-95-10 lipase was analyzed in this work using Ala mutagenesis. New results confirmed, that higher influence on activity and functionality of GD-95-10 lipase make Asp371 and Tyr376. Also the new GD-95 lipase variants were created using epPCR. The analysis of lipases obtained after random mutagenesis was showed that only a few changes in amino acids sequence can result new physicochemical properties of *Geobacillus* lipases. Therefore our results are basis for further enzyme engineering experiments and for creation of new lipases for bioconversion and ecotechnology.
USE OF SIGMA FACTORS FROM BACILLUS SUBTILIS IN THE DEVELOPMENT OF AN ORTHOGONAL EXPRESSION SYSTEM IN ESCHERICHIA COLI.

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Background

Technological advances in synthetic biology, systems biology, and metabolic engineering have boosted applications of industrial biotechnology for an increasing number of complex and high added-value molecules. In general, the transfer of multi-gene or poorly understood heterologous pathways into the production host leads to imbalances due to lack of adequate regulatory mechanisms. Hence, fine-tuning expression of synthesis pathways in specific conditions is mandatory.

Objectives

Here we develop a new genetic circuit for regulated expression specifically in stationary phase due to clear advantages during this period (reduction of toxicity, competition).

Methods

This circuit consists of a heterologous sigma factor (σ) recognizing specific promoter sequences, which are not recognised by the native σ factors of E. coli and is expressed upon entering the stationary phase. First, several σ factors of B. subtilis were tested for their orthogonality in E. coli on the level of promoter recognition, by using a red-fluorescent reporter system. Secondly, the potential of σ factors of B. subtilis to work together with the E. coli core RNA polymerase was tested, by expressing these proteins together with their promoters. Based on the results a specific factor will be chosen for further optimisation and the corresponding gene can be cloned in the σS factor operon of E. coli, which is most abundantly expressed in stationary conditions.

Conclusions
Combining all these elements should allow us to create an orthogonal genetic circuit that is able to transcribe specific genes under stationary phase with a limited influence on the host cell’s metabolism.
Background
Polyethylene terephthalate (PET) is a synthetic aromatic polyester that is degraded by several hydrolases from actinomycetes. The heterogeneous enzymatic PET hydrolysis occurs at mild temperature and pH conditions and offers an environmentally friendly alternative to chemical plastic recycling processes. LC cutinase, TfCut2 and Cut190 are homologous polyester hydrolases produced by actinomycetes obtained from a compost metagenome, from Thermobifida fusca KW3 and Saccharomonospora viridis AHK 190, respectively.

Objectives
The enzymatic degradation of PET films by the three polyester hydrolases was compared at various reaction conditions to evaluate the effects of buffer composition, pH, ionic strength and metal ions on their enzymatic activity.

Methods
Genes coding for LC cutinase, TfCut2 and Cut190 were cloned into pET-20b(+) vector for heterologous protein expression in Escherichia coli BL21(DE3). Purified enzymes were used to degrade PET films at 60 to 70 °C for 1 to 50 h in different buffer systems as well as in the presence of Ca$^{2+}$ and Mg$^{2+}$. The released products were analyzed by reversed phase high performance liquid chromatography. The weight losses of PET films following an enzymatic hydrolysis by the different enzymes were determined gravimetrically.

Conclusions
The LC cutinase from the compost metagenome and TfCut2 from T. fusca showed significantly higher PET hydrolytic activity compared to Cut190 from S. viridis. The addition of both Ca$^{2+}$ and Mg$^{2+}$ stabilized the enzymes and thus facilitated an effective PET degradation. The enzymatic activity of the polyester hydrolases against PET films was strongly depending on the type and ionic strength of the selected buffer.
A DISRUPTIVE INNOVATION FOR AMBIENT TEMPERATURE PRESERVATION OF MICROORGANISMS

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Background
Because of the increase in the number of samples of microorganisms to be stored and shipped, classical preservation in freezers, subjected to risks of technical failure, is becoming more and more costly in space, energy and maintenance. Freeze-drying offers an alternative to cold preservation. However, it is time-consuming and requires precise cycle adjustments. Moreover, glass vials are subjected to breakage and can become leaky resulting in decrease in bacterial viability over time. Imagene innovation aims at preserving dehydrated bacteria at ambient temperature (AT) in hermetic minicapsules that maintain an anoxic and a low-controlled humidity environment. The minicapsules are resistant to impacts and have a unique, engraved datamatrix code allowing a tamper-proof traceability of the samples.

Objectives
This study aims at demonstrating the effectiveness of a novel and rapid process for a reliable long term AT storage of microbial strains.

Methods
Two bacteria, Pantoea dispersa and Aeromonas Salmonicida, a freeze-drying sensitive strain, were desiccated by evapo-concentration in presence of different stabilization solutions and encapsulated in minicapsules, which were stored either at AT or 37 °C (long-term storage simulation). Bacteria were recovered by simple rehydration.

Conclusions
The residual viability rates post-dessication were \(~80\%\) for P. dispersa and \(~60\%\) for A. salmonicida. Both strains kept a residual viability rate \(>50\%\) after 4 weeks at 37 °C and at AT in Imagene minicapsules. Viability rates following Imagene process were comparable to that of freeze-drying. In conclusion, our work demonstrated that this new process is suited to AT preservation of bacteria.
Background
Antrodia cinnamomea grows only on Cinnamomun kanehirai Hay (host specificity) and it is a unique fungus in Taiwan. In recent studies, the major components of Antrodia cinnamomea are triterpenoids whose functions were anti-inflammation and anti-cancer. So it is a valuable fungus in pharmaceutical potential. However, Antrodia cinnamomea grows slowly and does not obtain the fruiting body form easily under artificial cultivation.

Objectives
Based on this consideration, this study modified the plant factory as the new fungus fermenter to increase the triterpenoids products.

Methods
We isolated the new Antrodia cinnamomea strain (CUST-AC901) from the trees in the mountainous areas of Hualien in Taiwan. It was fermented on this new established fermenter without light, which is one type of closed plant production systems (CPPS) consisting of a thermally insulated, air conditioners, a CO2 and nutrient solution supply unit. The triterpenoids were analyzed by HPLC.

Conclusions
We established a new high-efficient solid-state cultivation with CUST-AC901 strain and obtained the optimal conditions (temperature at 20°C, CO2 at 10%, humidity at 85%, and with 1% peptone in whole grains medium). We also obtained compositions on fruiting bodies similar to the wild ones. It only took two months and reduced the producing cost. With HPLC analysis, we confirmed that total triterpenoids products were increased by four-folds than they were under traditional culture method. To our knowledge, this is the first report on omni-functional solid-state fermenter with high triterpenoids product efficiency and low cost as well as reaching to automatic mass production without expensive automatic production equipment.
THE IMPACT OF CYTOSOLIC AND PEROXISOME LOCALIZED TRANSALDOLASE AND TRANSKETOLASE ON XYLOSE METABOLISM AND ALCOHOLIC FERMENTATION IN METHYLOTROPHIC YEAST HANSENULA POLYMORPHA

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Background

Lignocellulose is currently considered as the most promising renewable feedstock for production of liquid biofuels. Xylose is the second most abundant sugar in nature, being a component of hemicellulose, which is in turn a part of lignocellulose. Yeasts and most of mycelial fungi can metabolize xylose by the so called xylose reductase – xylitol dehydrogenase pathway including xylose reduction to xylitol and subsequent xylitol oxidation to xylulose which is then phosphorylated by a specific kinase and enters pentose phosphate pathway (PPP). A simultaneous improvement of xylose conversion to xylulose and in the activities of PPP enzymes is required for efficient xylose utilization in different microorganisms. Xylose fermenting yeast Hansenula polymorpha contains in addition to cytosolic transaldolase (gene TAL1) and transketolase (gene TKL1) also peroxisomal transketolase (also known as dihydroxyacetone synthase, gene DAS1) and putative peroxisomal transaldolase (gene designated by us as TAL2).

Objectives

To investigate the role of cytoplasmic and peroxisomal transaldolase and transketolase in xylose metabolism and alcoholic fermentation in H. polymorpha.

Methods

Molecular-biology techniques

Conclusions

In the wild type strain of H. polymorpha overexpression of DAS1 and TAL2 turned out to be beneficial for xylose alcoholic fermentation. Moreover, mutants with knock out of at least one of these genes were impaired in xylose fermentation as compared to the
wild-type strain. To investigate the role of cytosolic transaldolase and transketolase in xylose metabolism of *H. polymorpha* recombinant strains overexpressing *TAL1* and *TKL1* genes were constructed. However, such strains revealed no substantial difference in ethanol production as compared to the wild-type strain.
Background

Wood is a direct food contact material used in the food industry since ancestral times. In Europe, wooden food contact surfaces are subject to the European regulation no. 1935/2004, which specifies that food contact material must not interfere with foodstuff characteristics (Anonymous, 2004).

Today, no standard recovery method was defined for wooden food contact surface. Therefore, it might be important to provide an efficient method to quantify the microbial load on the wooden surfaces in direct contact with food.

Objectives

The aim of this study was to compare three methods of recovering microorganisms from wooden surfaces: grinding, brushing and planning (Ismaïl et al., 2014).

Methods

Three microorganisms, well-known as risk along the food chain, were tested: *Listeria monocytogenes*, *Escherichia coli* and *Penicillium expansum*. We chose three wooden species - pine, poplar and spruce - which are mainly used to manufacture wooden packaging. We analyzed the influence of wooden moisture content, contact time and wood timbers on microbial recovery rates.

Conclusions

We identified that factors cited above influenced the microbial recovery rates from wooden surfaces. Grinding was the most reliable method with the best recovery yield: 30.1% for *Listeria monocytogenes* on spruce and *Escherichia coli* on poplar, and 30.4% for *Penicillium expansum* on poplar. Planing was chosen to be applied to thicker wooden as cheese ripening shelves and tested, then, for analyzing the microbial load of 54 various wooden ripening shelves. We did not found common
pathogens known as risks for dairy products. Our results suggest that wood is suitable for food contact.
Background
GRAS status, the availability of different types of mutants and the ability to perform eukaryotic post-translational modifications, including complex glycosylation and protease processing, make Saccharomyces cerevisiae an ideal host for the expression of recombinant proteins. Among recombinant enzymes, lipases are used in many different industrial applications including pharmaceutical synthesis, biodiesel production, detergent formulation and food industry.

Objectives
In the present work, lipase A from Bacillus subtilis was expressed in the S. cerevisiae CEN.PK113-5D strain as a fusion protein with the yeast cell-wall mannoprotein Pir4. In order to compare the effect of different promoters on the levels of lipase activity secreted into the growth medium, three different YEplac195 based constructions were created, differing only in the promoter controlling the expression of the gene fusion; the original weak and constitutive PIR4 promoter, the TEF1 promoter, strong and constitutive, and the inducible GAL1 promoter.

Methods
The resulting three strains were grown in rich non-selective medium and the levels of secreted lipase activity and plasmid loss were determined after 48 hours of growth.

Conclusions
Our results show that the use of a very strong constitutive promoter is a poor strategy, since plasmid loss occurs in over 80% of cells and consequently total activity levels remain low in comparison with the strain transformed with the construction including a weaker promoter, where plasmid loss occurs in around 20% of cells. Finally, maximum levels of activity were achieved using the GAL1 promoter in conditions of induction for 24 hours after 24 hours growth in non-inducing conditions.
ENGINEERED SALMONELLA ENTERICA TYPHIMURIUM CONTROLS TUMOR GROWTH IN MURINE MELANOMA MODEL

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Background
The cancer is the second most frequent cause of death. Melanoma, a malignancy that arises from melanocytes, accounts for approximately 10% of skin tumors and in recent years its incidence has increased. The mortality rate currently stands at 80%, while 5-year survival rate is less than 5%. The factors that affect cancer outcomes, such as disease recurrence, risk of second malignant neoplasms, and the late effects of cancer treatments, becomes more important. For almost 200 years has been known that bacteria have the ability to colonize solid tumors and induce tumor shrinkage. The bacteria Salmonella enterica Typhimurium are facultative anaerobes and has a particular promise as a cancer therapeutic because it can be manipulated and has been shown to preferentially accumulate in tumors compared with other organs after systemic injection. S. enterica strains can have the anti-tumor activity enhanced by genetic manipulation.

Objectives
This study aims to evaluate the anti-tumor potential of attenuated new mutants of S. Typhimurium in murine models.

Methods
Six –week-old female C57Bl/6 mice were subcutaneously injected with 5x10⁵ B16F10 (murine melanoma). After 12 days mice were injected intratumoral with 10⁷ CFU/mL of S. enterica LGBM127.

Conclusions
Our results showed a reduction in tumor growth and an increased survival rate in mice treated with S. enterica when compared to the control group (PBS). There were no signs of prostration after LGBM127 inoculation. More tests are needed, but our preliminary data demonstrate a high potential for the use of the attenuated strain of S. Typhimurium as antitumor agents.
Background

The enzymatic saccharification of lignocellulosic biomass and biobleaching paper pulp are attractive strategies for the bioprocess industry where xylanases have been heavily applied. These processes require enzymes that tolerate high stress environmental conditions such as extreme temperatures. The wild xylanase from *T. aurantiacus* (xynA) has been described as a potential enzyme for rational protein engineering studies. In the first step of this study, two no silent mutations were detected in the xynA sequence, cloned in *S. cerevisiae* strain, and it may explain the reduced expression and activity.

Objectives

New strategy was proposed to overcome the expression issues and enzyme improvement. First, xynA gene was rebuilt and it was used as template for constructions of mutants by site-directed mutagenesis (SDM) to increase thermal tolerance and activity.

Methods

For this, the original gene was amplified and cloning into pET28a. For the SDM reactions were used the Q5® SDM Kit. Primers were constructed according to the desired mutations and each one used for its changing. In the total, 13 reactions were prepared, in which only the first two were done to the reconstruction of xynA. All the mutagenic xylanases were characterized in pH (3.5-10) and Temperature (40-80°C) by enzymatic assay.
Conclusions

Regarding the optimum temperature, the enzymes are active at 65°C, and pH 5. About the stability temperature, the most of them are stable from 40°C to 65°C, and four mutants until 75°C, both in wide range of pH. These results helped the selection of the best mutagenic xylanases for future assays.
NOVEL BIOINSECTICIDES BASED IN BACILLUS THURINGIENSIS SPORES TO CONTROL PLAGUE DISEASES IN PLANTS

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Background

The massive use of synthetic chemical agents of biocontrol (i.e. insecticides) with potential toxicity to the users and the environment pushes for the development of alternative (ecological and natural) compounds. To this respect, the use of natural and human-friendly bacteria with insecticide (i.e. larvicide) activities (biocontrol agents) represents an interesting alternative.

Objectives

Bacillus thuringiensis, is a ubiquitous Gram-positive, spore-forming bacterium that forms soluble Vip toxins a parasporal crystal (Cry toxin) during its growth cycle. These B. thuringiensis toxins are efficient for the control of certain plant-detrimental insect species among the orders Lepidoptera, Diptera, and Coleoptera.

Methods

One hundred nineteen B. thuringiensis strains isolated from different agricultural regions of Argentina were characterized for Vip and crystal protein production and evaluation of toxicity against Lepidopteran insect's larvae. Finally the genomic sequencing of the 16S ribosomal subunit genes plus the PCR identification of cry and vip subclasses of selected isolates were performed.

Conclusions

Several of the novel isolates were able to simultaneously control Spodoptera frugiperda and Rachiplusia ñu larvae under laboratory and open field conditions. Interestingly, many of the isolates harbored both insecticidal genes (vip, cry). The role of the master transcription regulator Spo0A on insecticidal protein production is presented. The understanding of the genetic network under Spo0A command open new avenues for the development of intelligent biocontrol agents in the scope of integrated pest management for sustainable agriculture.
NOVEL BACTERIAL INOCULANTS, BASED IN BACILLUS SPORES, TO INCREASE AND PRESERVE THE NUTRITIONAL QUALITY OF ALFALFA IN LIVESTOCK FODDER

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Background

Many problems associated with haymaking would be solved by reducing hay-drying time or if hay could be stored at high moisture without fungal spoilage. Bacterial inoculants are an ecological alternative for improving hay (and silage) preservation.

Objectives

The aim of this study was to identify, isolate and select the best spore-forming Bacillus strains capable of inhibiting the growth of undesirable fungus and accelerate the dehydration of forage in order to conserve its original nutritional value.

Methods

Seventy Bacillus strains were isolated from alfalfa fields of Argentina. Small bales and open field assays were prepared and treated with each Bacillus strain in absence and presence of Fusarium verticilliodes. These bales were daily monitored throughout the trial for evidence of mold growth, dehydration of alfalfa and bale organoleptic quality. Germination rates / outgrowth and biofilm / spreading were evaluated in AFGK-buffer and LBY medium, respectively. Genomic identification of selected isolates was performed by sequencing of the gene coding for the 16S ribosomal RNA. Five isolates (initially identified as B. subtilis, B. pumilus and B. amyloliquefaciens) showed a broad spectrum of fungal control and an accelerated dehydrating capacity of alfalfa that improved the retention of its nutritional values in the forage. Viable spores were showed to be needed for these effects and the capacity of alfalfa colonization by mechanisms of biofilm formation and spreading were identified.

Conclusions

The novel Bacilli presented in this work, and the understanding of the displayed molecular mechanisms, are suitable for the production of novel inoculants to conserve forages of high nutritional value.
GENE DISRUPTION ANALYSIS FOR ETHANOL PRODUCTION IN GLYCEROL FERMENTATION IN GLYCEROL-ASSIMILATING KLEBSIELLA VARIICOLA AND ITS METABOLIC CHANGES

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Background

The gram-negative enteric bacterial genus *Klebsiella* can utilize glycerol as a sole carbon source and is known to produce some valuable metabolites, such as ethanol and 1,3-propanediol (1,3-PD). Previous studies, we have obtained streptomycin-resistance *K. variicola* mutant, strain TB-83D, as an ethanol producer from glycerol, and have demonstrated that it showed higher ethanol production than parent strain (Suzuki et al. Bioresour. Technol. 2015.). To further increase ethanol productivity, metabolic changes and inhibition of byproduct production by gene disruption were required.

Objectives

To further improve ethanol production, the aim of this study is to increase gene induction efficiency for gene disruption and to suppress byproduct production by gene disruption.

Methods

Gene disruption was performed by using modified Red recombination method. Metabolite concentrations were measured by HPLC (Waters alliance 2695) and F-kit D-/L-lactic acid (Boehringer Mannheim).

Conclusions

Since strain TB-83D produced mainly D-lactate under optimum conditions, we considered that suppression of D-lactate production is effective in improving ethanol production. To suppress D-lactate production, we disrupted lactate dehydrogenase gene (*ldhA*) by Red recombination. Gene disruption efficiency of *Klebsiella* species is lower than that of other *Enterobacteriaceae* family. Therefore, we modified the conventional Red recombination, and succeeded in improvement of gene disruption efficiency. Although *ldhA* disruptant did not produce lactate in glycerol fermentation, ethanol production increased than TB-83D. Metabolic flux analysis and metabolic changes are in progress to optimize the ethanol production process, the results of which will be described in detail in due course.
DESIGN OF SACCHAROMYCES CEREVISIAE PRODUCING CELLULOSE-BASED CELLULOLYTIC ENZYME COMPLEXES FOR LIGNOCELLULOSIC BIOMASS UTILIZATION
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Background
Heterogeneous structure of lignin imparts plants with structural rigidity and also serves to protect cellulose and hemicellulose from degradation. Thus, prior to fermentative production of ethanol from the cellulose by yeast strains, the materials are degraded and hydrolyzed to release monomeric sugars.

Objectives
In this study, designer cellulosome was assembled in yeast Saccharomyces cerevisiae for utilizing of cellulose as the substrate. For utilizing of cellulose part in lignocellulosic biomass by simultaneous saccharification and fermentation, a recombinant scaffolding protein from Clostridium cellulosivorans and a chimeric endoglucanase E from Clostridium thermocellum were assembled as complex system. Compared to the results for single subunit, assembly of cellulosome-based enzyme complexes caused a noticeable increase such as 2.1-fold-higher in the level of enzyme activity. The resulting strain was able to ferment cellulose part in pretreated barley straw into ethanol with the aid of beta-glucosidase A from C. thermocellum. In the fermentation assay at 10 g/L initial substrate, approximately 2.1-folds higher ethanol than that of wild type was produced.

Methods
The use of complexed enzyme systems is one of the strategies for effective lignocellulosic biomass hydrolysis. Enzyme complexes were formed via the interaction of a dockerin domain with cohesin modules in the scaffolding protein.

Conclusions
Accelerating the biological degradation of lignocellulosic materials will benefit from the development of useful recombinant enzymes with hydrolysis ability. In future research, construction of designer enzyme complexes containing other lignin degrading enzymes could be used to develop biocatalysts that can completely degrade lignocellulose into single sugars.
Background:

The purple nonsulfur photosynthetic bacteria, *Rhodobacter sphaeroides*, are capable of growing under a variety of environmental conditions. Particularly this microorganism can use carbon dioxide as a carbon source to grow. In addition, this bacterium also has the capability to produce all kinds of the polyphenol, nucleic acid material, vitamin, bio-active substances and polyhydroxybutyrate (PHB). Therefore, this study focused on analyzing the properties of organic compounds and reduction of CO$_2$ in *R. sphaeroides* mutant strains.

Objectives:

In this study, relying on the ability to produce physiologically active compounds (PACs) and reduce CO$_2$, four different chemically mutated strains: *R. sphaeroides* KCTC 1434, MBTLJ-8, MBTLJ-13 and MBTLJ-20 were screened in order to select the most optimal strain.

Methods:

*R. sphaeroides* KCTC 1434, MBTLJ-8, MBTLJ-13 and MBTLJ-20 that are all chemically mutated strains, were used in this study. To be more precise, the comparison among these strains in terms of their ability to produce physiologically active compounds (PACs) and reduce CO$_2$ was carried out.

Conclusions:

The result showed that the mutant strain MBTLJ-8 performed the best ability of CO$_2$ fixation. In addition, it also showed that the CO$_2$ reduction and cell growth of mutant strains enhanced under the blue colored light emitting diode (LED) conditions. Therefore, our data indicated that the wave length of special light probably affected the cell growth, CO$_2$ reduction and organic compounds production of mutant strains.
Acknowledgments: This work was carried out with the support of “Cooperative Research Program for Agriculture Science & Technology Development (Project No: PJ01051502)” Rural Development Administration, Republic of Korea.
EFFECT AND ANALYSIS OF RHODOBACTER SPHAEROIDES EXTRACT UNDER VARIOUS CULTURE CONDITIONS

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Background:

*Rhodobacter sphaeroides* is a gram negative purple nonsulfur photosynthetic bacterium, facultative photoheterotrophic bacterium capable of growing phototrophically or chemotrophically as either a heterotroph or lithotroph in the presence or absence of O₂, depending on the energy source. In addition, it owns capability to produce all kinds of amino acids, nucleic acid materials, vitamins, bioactive substances, polyphenol and polyhydroxybutyrate (PHB).

Objectives:

In this study we focus on analyzing the properties of organic compounds and reduction of CO₂ in *R. sphaeroides* under various culturing conditions. Moreover, it is another goal of this study to confirm the effect on promoting growth of other microorganisms caused by the complement of *R. sphaeroides* extract in various culture conditions.

Methods:

*R. sphaeroides* KCTC1434 were used in this study. More precisely, the analysis of growth promoting effect of the extract of *R. sphaeroides* was conducted under various culture conditions.

Conclusions:

Our data indicated that addition of the *R. sphaerodes* extract into culture media accelerated the growth of microorganisms. Therefore, there was a strong evidence showing that the *R. sphaerodes* extract has multifunctional potential to increase the physiological activity of other organisms. It is, thus, determined as an enhanced material which is able to force to activate the growth promotion of other organisms.

Acknowledgments: This work was carried out with the support of “Cooperative
Research Program for Agriculture Science & Technology Development (Project No: PJ01051502) Rural Development Administration, Republic of Korea.
CLONING AND HETEROLOGOUS EXPRESSION OF ACID AND THERMOSTABLE $\beta$-GLUCOSIDASE IN PICHIA PASTORIS X-33 FROM THERMOASCUS AURANTIACUS RCKK

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Background
Thermostable cellulases offer several advantages like higher rates of substrate hydrolysis, lowered risk of contamination and increased flexibility with respect to process design. The production of cellulases from a thermophilic fungus identified as Thermoascus aurantiacus RCKK (Acc. No. JN676149) has been attempted.

Objectives
In order to increase production level of cellulases and eventually economize the process, heterologous expression of these genes in Pichia pastoris was carried out.

Methods
$\beta$-glucosidase gene from T. aurantiacus RCKK was cloned and overexpressed in P. pastoris X-33.

Conclusions
A 2.5 fold higher production of recombinant $\beta$-glucosidase was achieved at shake flask level in comparison to natively expressed enzyme. Expression of $\beta$-glucosidase of about 96kDa was confirmed by SDS PAGE, western blot and zymogram analysis. The purified recombinant $\beta$-glucosidase was found to be stable in wide range of pH (3.0-8.0), temperature (up to 50% activity after 3 h at 80°C) and tolerant to the presence of ionic liquid (1-ethyl-3-methylimidazolium acetate [C2mim][OAc]. These properties make the recombinant enzyme an important catalyst for carrying out an efficient hydrolysis of cellulose to sugars and enzymatic deinking of recycled paper pulp. This will eventually open up potential of this enzyme.
Background

The main obstacle to lignocellulose bioconversion is the presence of lignin and crystalline cellulose in plant cell wall. However, major factors have not been definitely determined between lignin and crystalline cellulose in enzymatic lignocellulose recalcitrance. The correlative information promotes a deeper understanding of lignocellulose bioconversion characteristics.

Objectives

Our aim is to determine the major rate-limiting factor in the hydrolysis of popping pre-treated rice straw (PPRS) by examining cellulase adsorption to lignin and cellulose, re-hydrolysis, and amorphogenesis.

Methods

Lignin was prepared from PPRS by enzyme treatment until complete hydrolysis of carbohydrates and used to measure enzyme adsorption. Re-hydrolysis experiments were repeatedly performed with 15.75 or 31.50 FPU g\(^{-1}\)-biomass cellulase for 1 or 3 hour until complete hydrolysis of 1 or 2% PPRS. Amorphogenesis of PPRS was carried out with 80% phosphoric acid on ice for 1 hour.

Conclusions

Lignin isolated from PPRS adsorbed 20% of enzyme loading from 3.15-31.50 FPU g\(^{-1}\)-biomass, adsorbing low levels of exoglucanases, endoglucanase and xylanase. Marked inhibition by cellulose structural effect was occurred during re-hydrolysis steps. Amorphogenesis of PPRS resulted in twofold higher cellulase adsorption and increased the yield of the first re-hydrolysis step from 13% to 46%. The total yield was increased to 84% in 3 h. These results provide strong evidence that cellulose structure, not lignin, has a major effect on the enzymatic hydrolysis of lignocellulose.
BIOLOGICAL RECYCLING OF POLY(L-LACTIDE) (PLA) BY MICROBIAL ENZYME

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Background

Poly(L-lactide) (PLA) is a biodegradable polymer. It is synthesized from lactic acid that is produced by microbes from agricultural products. It has much attention since it plays an important role to resolve the global warming problem. In general, PLA is polymerized commercially by chemical processes. However, these processes need severe conditions, i.e. high temperature, high solvent and catalyst loading. Recently, biological polymerization methods have been presented using lipases for the biocatalytic reaction.

Objectives

This work we aimed to apply the biological process for recycle of PLA.

Methods

Using a protease produced by Actinomadura keratinilytica strain T16-1. 1 g/L PLA powder was incubated at 77 °C for 24 h using an enzyme concentration of 100 mg/L. As a result, 750 mg/L lactic acid was obtained and used as substrate for re-polymerization of PLA. This was done by using commercial lipase under 0.1 vvm nitrogen atmosphere at 60 °C for 6 h.

Conclusions

Oligomers with molecular weight of 450 (n=4) were obtained. This is the first report to demonstrate the recycling of PLA wastes biologically.
OPTIMIZATION OF EXTRACELLULAR LIPASES PRODUCED BY NEWLY ISOLATED BACILLUS SAFENSIS STRAIN PSR

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Background

Lipases are an important enzyme for many biotechnological applications due to they are biocatalysts which have an activity at mild condition. Recently, bacterial lipase was very interesting because it can be used for re-polymerization of biodegradable plastics such as poly(L-lactide).

Objectives

The present work was aimed to screen lipase-producing bacteria from oil-contaminated soil in South of Thailand and study the optimization of medium composition for lipase-production.

Methods

Lipase-producing bacteria with 662 strains were isolated from 100 soil samples according to opaque zone formation on Tween 80 agar. Strain PSR exhibited the highest lipase activities in both agar plate and production medium. It was identified as Bacillus safensis strain PSR based on 16S rRNA gene sequencing. Moreover, the factors influencing lipase production were investigated in shake flask such as carbon and nitrogen sources, pH, temperature and incubation time. The maximum enzyme activity 63.86 U/ml was obtained by using 0.5% (w/v) rice bran oil and 0.12 % (w/v) yeast extract as carbon and nitrogen sources, respectively, at pH 6.85, 37°C and 60 hours cultivation.

Conclusions

The study provides a high bacterial lipase production which is a good candidate to apply for recycle of biodegradable plastic waste in the near future.
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RECONSTRUCTION OF A CORE METABOLIC NETWORK RELATED TO MEDIUM-CHAIN-LENGTH POLYHYROXYALKANOATE PRODUCTION BY PSEUDOMONAS SP. LFM046 FROM CARBOHYDRATES.
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Background
Medium-chain-length polyhydroxyalkanoate (PHA\textsubscript{MCL}) are biodegradable elastomeric polymers produced from renewable resources. \textit{Pseudomonas} sp. LFM046 is a PHA\textsubscript{MCL} producer from carbohydrates sources, presenting a better performance than the most studied \textit{Pseudomonas} \textit{putida} KT2440.

Objectives
In this work, based on 16S rDNA sequences, \textit{Pseudomonas} species phylogenetically close to \textit{Pseudomonas} sp. LFM046 were identified and their sequenced genomes were searched for genes involved in carbohydrates catabolism and PHA biosynthesis, allowing reconstructed a core metabolic network. To confirm this network, the whole genomic DNA of \textit{Pseudomonas} sp. LFM046 was sequenced.

Methods
For this, genomic DNA was extracted and fragmented using Covaris. Paired-end sequencing library was constructed using TruSeq DNA PCR-Free LT Sample Preparation Kit. The library quantification was performed using QUBIT, Agilent Bioanalyzer and real time PCR. The library was sequenced on an Illumina MiSeq sequencer.

Conclusions
\textit{De novo} assembly generated 34 contigs using 3,032,982 reads, and the average size of 50% of the contigs (N50) was 640,128 bp, with the maximum size of 1,038,386 bp. The genome size was estimated at 5,970,318 bp. All genes involved in carbohydrate metabolism and PHA biosynthesis previously detected in other \textit{Pseudomonas} species were also identified and annotated in the genome of \textit{Pseudomonas} sp. LFM046 using RAST software. It includes genes from Entner-Doudoroff (ED) and pentose phosphate (PPP) pathways, tricarboxylic acid (TCA) and glyoxyl acid cycles (GAC), \textit{de novo} fatty acids (FAB) and PHA biosynthesis. Thus, a core metabolic network of \textit{Pseudomonas} sp. LFM046 was reconstructed and is going to be used to determine flux distribution based on exchange flux measurements.

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Background

Modern waste water treatment plants utilize activated sludge systems to reduce soluble compounds in waste water. The diversity of the biocenosis is primarily affected by the composition of the treated waste water. Especially the distribution of carbon and nitrogen leads to a characteristic diversity. Information about the specific abundance of essential microorganisms can be used to improve the plant operation and stability of the biological degradation.

Objectives

While available quantitative methods like MPN (most probable number) and qPCR (quantitative polymerase chain reaction) are time consuming, this work led to the development of a rapid and quantitative variation of the fluorescence in situ hybridisation (qrFISH). The main objective was to design a method for a daily industrial application.

Methods
Fluorescence in situ hybridisation was carried out according to Nielsen (2009). Fluorescence particles were added to the sludge samples and used as a reference for quantification. Thereby the unknown loss of biomass throughout the FISH process could be compensated.

Conclusions

The current results showed a strong correlation between cell counts before and after the hybridisation process. Figures 1 shows the comparison of quantification results by fluorescence microscopy and conventional counting in a Thoma chamber. The experiments were conducted with enriched cultures of bacterial cells. The primary limitation of quantitative FISH was overcome, and the results were achieved in the targeted time frame of 8 hours.
CHARACTERIZATION OF RECOMBINANT LACCASE (BPCOTA) FROM BACILLUS PUMILUS MK001 AND ITS POTENTIAL FOR PHENOLICS DEGRADATION

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Background

Bacterial laccases are very less explored despite their superior properties such as high thermostability and wide pH stability. In the present study, Putative laccase (CotA) from Bacillus pumilus MK001 has been cloned and expressed in E. coli BL21(DE3).

Objectives

Hyper-production of recombinant bacterial laccase was carried out to utilize its potential in bioconversion of phenolics.

Methods

BPCotA was expressed in E. coli BL21(DE3) and purified to homogeneity. Purified enzyme was characterized for its thermal and pH stability. The 3D model of BPCotA was constructed based on the crystal structure of B. subtilis CotA (PDB Id: 1GSK) by homology modelling (Modeller 9.11). Modeled structure was refined and validated for stereochemical quality using PROCHECK, ERRAT, Verify 3D and PROSA servers. In silico interaction between enzyme and phenolics was studied out to predict in vitro phenolics degradation ability of the recombinant enzyme.

Conclusions

The recombinant BPCotA was purified with ~82% recovery exhibiting ~230 IU/mg specific activity. The recombinant BPCotA retained more than 50 % of the original activity after incubating at 80°C and 90°C for 90 min and 30 min, respectively, and more than 80% of its activity in the wide pH range 5.0-9.0 for 240 h. Increase in temperature from 30-90°C resulted in significant changes in secondary structure as predicted by far UV thermal CD spectra. Docking analysis showed that both ferulic acid and vanillin bind in the vicinity of predictive active site of recombinant protein. Further, in vitro action of the enzyme on phenolics resulted in bioconversion of both ferulic acid and vanillin.
THE EFFECT OF METHANE DIGESTED SLURRY FOR ETHANOL PRODUCTION FROM GLYcerOL BY KLEBSIELLA VARIICOLA STRAIN TB-83D.

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Background
Biodiesel fuel (BDF) waste contains large amounts of crude glycerol as a by-product, and its treatment is an environmental problem. In this study, it was attempted to produce bio-ethanol from crude glycerol. In previous study, we isolated Klebsiella variicola strain TB-83D that could produce ethanol from glycerol. This strain has high potential of ethanol production, and its production was 32 g/L by using yeast extract (YE) as a nutrient. However, YE is very expensive, so it is not suitable for an industrial use.

Objectives
The objective of this study is to decrease its culture cost. And we attempted to use methane fermentation digested slurry (MFDS). MFDS is residue of methane fermentation and its cost is very low.

Methods
The experiment was conducted by 1 L fermenter at 25°C and the culture pH was maintained at around 8.0 with 6 N NaOH. Basal medium consist of 50 g/L of glycerol and 600 g/L of MFDS is used for ethanol production by strain TB-83D.

Conclusions
The ethanol production was decreased to 17 g/L by replacing YE with sterilized MFDS. However, we think that this result seems to be a promising start to establish a low cost ethanol production from BDF waste. Next, we used non-sterilized MFDS to reduce sterilizing cost. Unfortunately, ethanol production was decreased but 1,3-propanediol and acetate production were observed. It may be caused by other bacteria survived in the MFDS. Now we are trying to optimize the culture condition that strain TB-83D is dominant when using non-sterilized MFDS.
CAT8 GENE IS INVOLVED IN REGULATION OF XYLOSE ALCOHOLIC FERMENTATION IN THE THERMOTOLERANT YEAST HANSENULA POLYMORPHA

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Background

Efficient alcoholic fermentation of hexose and pentose sugars by yeast is essential to achieve maximal ethanol yield from lignocellulosic hydrolyzates. It is known that CAT8 encodes carbon source-responsive transcriptional activator that controls expression of genes involved in gluconeogenesis. Deletion of CAT8 in the non-conventional yeast Pichia guilliermondii activated alcoholic fermentation.

Objectives

The aim of this work was to isolate mutants of H. polymorpha with knock out of the gene CAT8 and study their alcoholic fermentation. The strains were planned to be constructed in the backgrounds of the wild-type strain and of the mutant with elevated ethanol production from xylose due to overexpression of the genes encoded the key enzymes involved in xylose metabolism (xylose reductase, xylitol dehydrogenase, xylulokinase) and resistance to 3-bromopyruvic acid (glycolysis inhibitor).

Methods

The cassettes for CAT8 gene deletion was constructed by used of homologous recombination method with gene HphNT1 conferring resistance to hygromycin B as the selective marker. Fermentation was carried out at a temperature of 45°C with limited aeration. Concentrations of ethanol in the medium were determined using alcohol oxidase/peroxidase-based enzymatic kit ‘Alcotest”.

Conclusions

Deletion of CAT8 gene did not have effect on glucose alcoholic fermentation. At the same time, constructed knock out strains showed an improved xylose alcoholic fermentation relative to the parental strains. The Δcat8 strains isolated from the best available ethanol producer accumulated up to 12.5 g of ethanol/L at 45°C after 3 day fermentation without correction for evaporation. This is the 30 fold increase as compared to the wild-type strain.
CHIMERIC ROTAVIRUS AND HEPATITIS A VIRUS-LIKE PARTICLES INDUCE VIRUS-SPECIFIC HUMORAL AND CELLULAR IMMUNE RESPONSES IN MICE

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Background
Rotavirus and hepatitis A virus (HAV) are transmitted by the fecal-oral route and continues to be endemic throughout the world. Virus-like particles (VLPs) is safe and represents an economical approach to combined multivalent vaccine development.

Objectives
D2/VP7 VLPs demonstrated excellent combined vaccine candidate and may lead to the development of a commercialized vaccine worldwide for preventing both rotavirus and HAV.

Methods
In the current study, we evaluated the humoral and cellular immune responses in mice vaccinated with hepatitis A virus/rotavirus VLPs vaccine manufactured in methylotrophic yeast Pichia pastoris strain GS115. The VLPs were comprised of the hepatitis A virus D2 epitope and rotavirus ΔVP7 proteins. BALB/c mice vaccinated with chimeric VLPs.

Conclusions
VLPs produced high levels of neutralizing antibodies in sera against rotavirus and HAV. Induction of the secretion cytokines were detected from anti sera. In addition, splenic CD4+ and CD8+ T cell accessory molecules increased and produced cytokines after vaccination.
Background

Biopolymers are becoming more and more attractive for the industry due to their biological/natural origin, sustainability, biodegradability and low toxicity. This results in an increased demand for novel biopolymers, including exopolysaccharides (EPS) for different biotechnological purposes. It is worth to mention a broad spectrum for applications ranging from human health, over food and fodder production to chemical industry and environmental applications.

The ability to synthetize EPS is quite common among bacteria. Thus, as described exopolysaccharides differ in their physical and chemical properties, it may be surprising that only few of those products were successfully commercialized.

Objectives

In our studies we focused on Acidobacteria, highly divers and environmentally widespread but not well studied bacteria phylum, as a source of valuable compounds. The ability to synthetize EPS has been previously reported for some members of this phylum. However, its chemical composition and properties were not investigated up to date.

Taking into consideration a total lack of information on acidobacterial EPS, the aim of the present work was first of all to gain insight into the nature of those polymers and second to explore some of the characteristics potentially novel and superior for industrial applications.

Methods

We thus, focused on isolation, purification and physico-chemical characterization of EPS. Additionally, the stability of emulsions formed with selected oils and hydrocarbons.

Conclusions
The study proved different composition of EPS produced by closely related bacteria translating into differences in its properties but also a high stability of emulsions suggesting potential of EPS as stabilizing agents and/or bio-emulsifiers.
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DETERMINATION AND CHARACTERIZATION OF PHYTASE FROM A NEWLY ISOLATED BACILLUS SP.
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Background
Phytate is the primary storage of phosphate in plants, and abundant in legumes, cereals and nuts. However, phytate make a polyanionic chelating agent which reacts with proteins, amino acids and divalent cations in humans and animals. Resulting in malnutrition and several health problems. Phytase is used as additive for degradation of antinutritional phytate and the enzyme is desired to be highly thermostable for it to withstand feed formulation conditions.

Objectives
The production of phytase is aimed from a new microbial source that was isolated from natural sources.

Methods
12 different soil samples taken from barn and coop surroundings were used for isolation of microorganism. To identify microorganism, we used 16srRNA analysis. For phytase production, strains were inoculated to Phytase-Screening Medium for 3d at 30°C. After phytase activity detection, enzyme was purified by ammonium-sulphate precipitation and DEAE-Sepharose. Samples were analyzed by SDS–PAGE.

Conclusions
Isolate (K12) was found Bacillus subtilis according to 16S rRNA sequencing. Several production parameters were optimized in order to ensure high enzyme production. In production medium, glucose was found as the most suitable C-source and ammonium nitrate (NH₄NO₃) was found to be most suitable N-source. In the presence of glucose, NH₄NO₃, sodium phytate and stirred speed (150 rpm), 72 hours incubation at 30°C were the most suitable conditions for phytase production. Phytase obtained from optimized production conditions, at pH 6.5 at 60°C and enzyme substrate ratio of 1:9 has the highest activity. MgCl₂, CuCl₂, CoCl₂, ZnCl₂ solutions and EDTA causing a loss of phytase activity whereas CaCl₂ solution has been found to be effective as an activity enhancer.
Background

In recent years, the availability of explosive whole-genome information has thoroughly reshaped the strain reconstruction studies for potential industrial applications. Consequently, the concept of minimum genome factories (MGFs) was proposed, which can be defined as recombinant strains whose metabolism have been streamlined to the optimal minimal subset applicable to specific utilities. The construction of MGF generally involves multiple large-scale genomic manipulations leading to a highly reduced genome. To this end, a wide range of genome engineering systems have been developed and applied in various microorganisms. One new issue raised thereafter is the fitness assessment of numerous resulting genome-reduced cells.

Objectives

Establish a simple and efficient system to evaluate the genome-reduced *Saccharomyces cerevisiae* chassis as platform for the production of value-added secondary metabolites.

Methods

A set of *S. cerevisiae* cells harboring large-scale deletions of different chromosomal regions was obtained using the recently developed mazF-based scarless genome engineering system. On the other hand, the coding sequence of tyrosinase *PPO2* from *Agaricus bisporus* was amplified and cloned into yeast expression vector pLC12. The recombinant plasmid was transformed into all deletion cells by lithium acetate method. The product of *PPO2* can convert intracellular tyrosine into colored substance melanin, which can be measured through spectral.
Conclusions
The tyrosine formation of yeast deletant cells can be readily deduced from the formation of melanin and this will be helpful in strain optimization for biosynthesis of valuable chemical based on tyrosine.
Background

Four varieties of *Aureobasidium pullulans* (Dothideales, Ascomycota), a cosmopolitan melanised yeast-like fungus with good stress tolerance, were recently redefined as four separate species: *A. pullulans*, *A. subglaciale*, *A. melanogenum*, and *A. namibiae*. These species inhabit various environments, including those with temperature and pH extremes, high UV radiation and high salt concentrations – and are thus considered to be polyextremotolerant. *Aureobasidium* spp. are biotechnologically important organisms, mainly due to their production of the extracellular polysaccharide pullulan, antimycotic aureobasidin A, siderophores, and various extracellular enzymes. *Aureobasidium pullulans* is also used in agriculture as a biocontrol agent while *A. melanogenum* is implicated in human disease. The recent sequencing of the genomes of the four above listed *Aureobasidium* species opened new possibilities for numerous novel applications in biotechnology, medicine, pharmacy, food industry, biofuel production, and in other fields.

Objectives

The aim of the study was to develop a yeast strain with good amylolytic activity under normal and stressful (industrially relevant) conditions.

Methods

In this study, we have cloned all alpha-amylase and glucoamylase encoding genes from the *Aureobasidium* species, used in the genome sequencing, into *S. cerevisiae*. The amylolytic performance of the transformants at different temperatures, low water activity and other relevant conditions was then evaluated. The best amylases were expressed in an industrial *S. cerevisiae* strain.

Conclusions

Yeast strains with good amylolytic activity under the conditions of increased salt concentration, low water activity, low temperatures and other relevant conditions were developed.
IMPROVING THE PRODUCTION OF HYDROPHOBIN HFBI FROM TRICHODERMA REESEI ON AN INNOVATIVE BIOFILM SOLID PACKING SYSTEM FERMENTOR

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Background
Hydrophobins are amphipatic proteins with high surface activity produced by ascomycetes and basidiomycetes. Their interfacial properties make them very interesting to use them as emulsifiers and stabilizers for the food industry. However, a major drawback is the low production of protein without the use of genetic techniques. Here we report the use of an innovative biofilm reactor which uses a metallic packing system as a substrate to grow on.

Objectives
- Increase the production of hydrophobin HFBI from Trichoderma reesei using a biofilm reactor system

Methods
- Design of a reproducible, easy to scale up and relatively cheap method to increase the production of a fungal hydrophobin from Trichoderma reesei.

- Fungal strain and culture medium: Trichoderma reesei MUCL 44908 were
inoculated in 20L of culture medium supplemented with glucose.

-Bioreactor set up: Along with the fermentation under biofilm system a submerged fermentation was also carried out in order to compare the increase in the production. In the case of the biofilm, the reactor was filled with a metal structured packing (Sulzer, Chemtech) where a peristaltic pump recirculated the medium evenly over the top of the packing system, this cycle was continued for the total time of the fermentation.

Conclusions
- The protein production was 2.16 times higher that when submerged culture was used.
- The total weight of the fungal biomass was 33% higher that in submerged culture.
- The separation process between medium culture and fungal biomass was greatly enhanced, more than 99% of the biomass was attached to the biofilm.
MICROBES AS TOOLS FOR MYCOTOXIN DEGRADATION

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Background

Mycotoxins are secondary metabolites produced by several fungi. These toxic chemicals play a major role as feed and food contaminants. They can severely impact animal health, which is associated with high costs in livestock industry. Approximately 25% of all commodities worldwide are polluted by mycotoxins. The most prominent mycotoxins in contaminated food are Aflatoxins (hepatotoxic and carcinogenic), Deoxynivalenol (immunosuppression, protein synthesis inhibition), Zearalenone (infertility, abortion), Fumonisins (hepatotoxicity, nephrotoxicity) and Ochratoxin A (Neurotoxicity and carcinogenicity).

Objectives

To counteract this problem, several strategies are applied, which involve prevention of fungal contamination during production and storage, followed by physical, chemical, and biological methods to reduce mycotoxin levels.

Methods

Biological methods such as the use of microbial detoxifiers have the highest potential for feed decontamination. Bacteria capable of metabolizing mycotoxins allow decontamination and preserve nutritional value at the same time. To identify microbial detoxifiers, contaminated soil samples are cultured with a mycotoxin of interest to enrich, potential degrading strains. This eases the screening of single colonies for detoxification, which is a time consuming procedure and associated with extensive labor. Once the organism responsible for detoxification has been isolated and identified, further experiments are necessary to ensure the organism is fermentable, non-pathogenic and can be applied as direct feed additive.

Conclusions

To date several detoxifying microbes have been found and characterized. An example for such a detoxifier is the strain BBSH 797 which degrades trichothecenes and is the only detoxifying feed additive on the market.
Background
Laccases are oxidoreductases enzymes that use molecular oxygen to oxidize phenolic and non-phenolic compounds. Laccases’ applications are related to bioremediation of hazards compounds, biobleaching and mill water treatment on pulp/paper industry, as component for phenolic detection, and for the synthesis of new “eco-friendly” polymers, antibiotics and cosmetics. Although its importance, it is still lacking ways to produce high amounts of laccases. Therefore, the search of new laccases and new producers are highly important for its commercial application. The marine-derived basidiomycete Peniophora sp. CBMAI 1063, isolated from the sponge Anphimedon viridis (Brazilian coast) is able to produce significant amounts of laccase. Previous studies showed that the laccase produced by this fungus has a good pH/thermal activity range and metallic ions resistance.

Objectives
To evaluate the laccase production by Peniophora sp. CBMAI 1063 under the presence of CuSO\(_4\) and saline conditions.

Methods
The fungus was cultivated in 50 mL of optimized medium under different concentrations of CuSO\(_4\) and artificial seawater (ASW) during 7 days at 140 rpm and 28 °C. Laccase was quantified spectrophotometrically using ABTS as substrate.

Conclusions
The highest laccase activity (around 2000 U L\(^{-1}\)), was obtained in the presence of 2 mM of CuSO\(_4\) and 65% (v/v) of ASW. There was no laccase activity in the absence of ASW, suggesting its importance in fungal metabolic pathway. Peniophora sp. CBMAI 1063 has shown a great potential to be a laccase producer under saline conditions. The produced laccase are being characterized and its genes are being studied for heterologous expression.
**Background**

Microbial exopolysaccharides (EPS) have wide range of applications in food, pharmaceutical and other industries due to their stabilizing, gelling and texturizing properties. Various EPSs from different microorganisms including yeast and bacteria have been determined. They can be either homopolysaccharides (e.g. cellulose, dextran, pullulan, levan, curdlan) or heteropolysaccharides (e.g. xantan, gellan).

**Objectives**

In this study, exopolysaccharide production of a novel Bacillus strain ZBP4 on molasses was investigated. Effects of some environmental conditions on the production were also determined.

**Methods**

The bacterial growth was carried out in Nutrient Broth at 35 °C for 24 h. The medium for EPS production contained 60 g molasses, 5 g yeast extract, 5 g peptone, 1 g K$_2$HPO$_4$, and 0.1 g MgSO$_4$.7H$_2$O in one liter of distilled water. Effects of temperature (30 to 45 °C), pH (4.0- to 9.0), substrate concentration (10, 20, 30, 40, and 60 g/L molasses), nitrogen sources (ammonium sulfate, peptone, tryptone, and yeast extract), and aeration (80, 100, 150, and 200 rpm) on the EPS production were determined. For the isolation of EPS, cells were removed by centrifugation and supernatant was boiled and proteins were precipitated with trichloroacetic acid. Protein free supernatant was mixed with cold ethanol and centrifuged to precipitate EPS. Amount of EPS was determined with phenol-sulfuric acid method using glucose as standard.

**Conclusions**
The strain was produced highest amount of EPS at 60 g/L molasses concentration, 33 ºC, pH 5.0. Tryptone was the best nitrogen source for EPS production. At 100 rpm agitation speed EPS production was the highest.
ASSEMBLY AND GRAFTING OF PATHWAYS FOR SYNTHESIS OF BERRY HIGH-VALUE PHENOLICS IN LACTOCOCCUS LACTIS.

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Background

Plant phenolics demonstrate promising effects in the combat of cardiovascular disease, certain types of cancer, neurodegenerative diseases, allergies, diabetes and inflammation. Accordingly, the food industry actively searches for natural alternatives, such as anthocyanins, to replace synthetic dyes. The increasing consumers demand for plant-derived natural products with health-promoting (eg. nutraceuticals) and/or biotechnological applications (eg. food colorants) requires the implementation of innovative solutions to the large-scale production of such compounds. One such opportunity is offered by the exploitation of microbial hosts as cell factories.

Objectives

Our aim was to reconstruct the pathway for plant anthocyanin pelargonidin-O-glucoside production from flavanone naringenin in *Lactococcus lactis*.

Methods

In order to assemble a functional pathway, native genes from various plants and synthetic codon-optimized genes were cloned in *L. lactis*, and their expression was tested in this bacterium. Then, highly expressed genes were assembled in one anthocyanin production cluster in *L. lactis*.

Conclusions

In this study, we present the construction of an artificial gene cluster consisting of four native and codon-optimized plant genes for synthesis of anthocyanin pelargonidin-O-glucoside from naringenin in *L. lactis*. 
INHIBITORY EFFECTS OF VARIOUS ENVIRONMENTAL CONDITIONS ON METHANOGENIC CONSORTIA

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Background
In our previous work three methanogenic consortia were selected from environments with high concentration of methane with biological origin: (i) anaerobic digester biogas plant (AD), raw sewage sludge waste (RSS), cattle slurry (CS). Those consortia are highly specialized in stable and effective biogas production during mesophilic anaerobic digestion from maize silage.

Objectives
The aim of this study was verification of: (i) the ability to effective production of biogas from maize silage by the selected microbial consortia under various temperature and pH conditions, and (ii) the stability of microbial consortia community under various temperature and pH conditions.

Methods
Lab-scale anaerobic digestion process. Biogas yield: daily biogas production, methane content were analyzed by gas chromatography GC-MS. Physical and chemical analysis: pH, volatile fatty acids (VFA), chemical oxygen demand (COD), ammonium, total solids (TS), volatile solids (VS). Determination of total cell counts by fluorescence microscope at a magnification of 600, using the filter set WU for DAPI detection. DNA isolation and purification. PCR amplification. Denaturing gradient gel electrophoresis (DGGE) of archaeal and bacterial 16S rRNA.

Conclusions
The obtained results showed that the selected consortia are able to produce biogas in broad range of environmental conditions. However, the process is running with a lower methane yield in psychrophilic and acidogenic condition. These results also demonstrated that selected consortia have broad spectrum of activity and could be used as a “biostarter” in start-up trial operation in different biogas plant technology process.
COMPARATIVE EFFICIENCY OF ADDING ETHANOL OR POTASSIUM METABISULPHITE IN FUEL ALCOHOL FERMENTATIONS CONTAMINATED WITH DEKKERA BRUXELLENSES: EFFECT ON THE CELL NUMBER AND ETHANOL YIELD


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Background
Fuel alcoholic fermentations contaminated with Dekkera bruxellensis result in decreased ethanol yield.

Objectives
The aim of this work is to compare the efficiency of two cell treatments carried out prior to the fermentation to control the growth of D. bruxellensis without affecting the starter yeast Saccharomyces cerevisiae and the ethanol yield, in pure and mixed cultures.

Methods
The treatments consisted in the addition of 13% (v/v) ethanol or 250mg/l potassium metabisulphite (PMB) to the acidic solution (pH 2.0), for 2 hours at 30°C, under shaking, in which the cells were introduced. After treatment, the cells were washed twice with sterile water, centrifuged and inoculated in sugar cane juice for 12 hours. The cell number was estimated by plating the samples in WLN medium with and without actidione to count the CFU number of D. bruxellensis and S. cerevisiae, respectively. Alcohol production was also determined.

Conclusions
In the mixed culture, the addition of ethanol to the acidic solution had similar effect over S. cerevisiae and D. bruxellensis, however the cell number increased after the fermentation time only for S. cerevisiae. The ethanol production was reduced by 6%. When PMB was used, the ethanol production was decreased considerably (74%) although the number of S. cerevisiae was not affected. D. bruxellensis was not affected at all. The interaction between PMB and S. cerevisiae to inactivate the toxic effect of this substance may be the reason why the fermentation yield was affected considerably. The addition of 13% ethanol in the cell treatment was the most efficient and the least harmful. Support: FAPESP
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LIGNINOLYTIC ENZYME ACTIVITIES AND CORN STRAW DELIGNIFICATION BY THE FUNGAL STRAIN CMU-196 OF PARACONIOTHYRIUM SP.

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Background
Filamentous fungi degrade lignin by producing extracellular oxidative enzymes such as laccase (LAC), lignin peroxidase (LiP) and manganese peroxidase (MnP). These enzymes have diverse biotechnological applications, including straw delignification intended for biofuels production or ruminant feed. Paraconiothyrium is an ascomycete genus recently described whose species can produce ligninolytic enzymes; however, until now only few strains within the genus have been analyzed for this purpose.

Objectives
The aim of this work is to analyze the production of the extracellular LAC, LiP and MnP in the strain CMU-196 of Paraconiothyrium sp., and to evaluate the corn straw delignification by this strain.

Methods
Basal and induced enzyme activities were determined by incubating the strain CMU-196 in potato dextrose broth (PDB) at 24 °C and 128 rpm. Induced media were supplemented with ground corn straw (2% w/v), an aqueous extract of it (10% v/v) and CuSO₄ (150 µM). Corn straw delignification in a solid state fermentation was followed by scanning electron microscopy (SOM).

Conclusions
Maximum activity for MnP (83.72 U/ml) and LAC (103.64 U/ml) was shown at 5th day of incubation in PDB supplemented with ground corn straw. Maximum activity of LiP (25.66 U/ml) was observed at 6th day of incubation in basal medium. SOM images reveal an increase in delignification of vascular tissue of the corn straw inoculated with the fungus since week 4th until the end of treatment at week 12th. The strain CMU-196 produce high levels of the three extracellular ligninolytic enzymes and has potential for straw delignification.
Background

The contamination of fuel alcohol fermentations with *Dekkera bruxellensis* brings about decrease in ethanol productivity and yield.

Objectives

Growth and fermentation profiles of two strains of *D. bruxellensis* and one strain of *S. cerevisiae* were evaluated under varying conditions of shaking in synthetic medium with glucose or sucrose as carbon source in different concentrations. We aimed to verify if the conditions to obtain higher alcohol production are similar for both yeast species.

Methods

Erlenmeyer flasks containing yeast cells in 200 ml of synthetic medium containing glucose or sucrose (50, 100 and 150g/l) were maintained at 30°C for 96 h under 0, 150 or 250 rpm. Growth was monitored by optical density at 600 nm and alcohol production was determined by density.

Conclusions

The strains of *D. bruxellensis* displayed similar results concerning growth: it was higher under shaking conditions regardless the carbon source type and concentration and was higher in sucrose than in glucose. The best conditions to achieve higher alcohol production were not similar and it was dependent on the shaking and the carbon source concentration. For *S. cerevisiae*, the growth was higher with shaking and no effect of the carbon source type and concentration was detected. Alcohol
production was substantially higher in static cultures regardless the carbon source. Sucrose at 150g/L, which is the basic fermentation medium in static cultures, favors alcohol production by S.cerevisiae but it does not for D.bruxellensis, whose growth is stimulated by sucrose. These results confirm the role of this yeast as contaminant and shows the variability exhibited among strains. Support: FAPESP
Background
Membrane proteins (MPs) play a crucial role in cell biology of any living organism. Nearly 30% of all open reading frames encode MPs and more than 50% of currently available pharmaceuticals target MPs. Despite their importance, the function of most MPs has not been assigned, yet.

Objectives
Since production of MPs is commonly limited by their hydrophobic nature, it is inevitable to generate novel expression systems that allow the high-level production of this class of enzymes. Here, we present an alternative expression system based on the bacterium Rhodobacter capsulatus, which is – due to its phototrophic nature – especially suited for the synthesis of MPs.

Methods
The facultative anaerobic purple bacterium R. capsulatus is able to grow either chemotrophically or phototrophically. Under latter conditions, it forms an intra-cytoplasmic membrane system (ICM) and vesicles, harboring the proteins of the host’s photosystem and heterologously produced MPs. To characterize the expression properties of this system, several genes encoding monoxygenases, diagnostic-relevant MPs as well as GPCRs were cloned into a comprehensive set of novel broad host range expression plasmids of the pRho series. Subsequent accumulation, localization and activity studies were conducted to evaluate the alternative expression system.

Conclusions
With R. capsulatus and the pRho plasmids we have developed an alternative, powerful expression system for the synthesis of heterologous MPs. Our system is particularly suited for the production of monoxygenases, diagnostic-relevant proteins and GPCR-(like) proteins with yields of more than 10 mg per liter culture.
INNOVATION IN THE CONJUGATION PROCESS OF THE CAPSULAR
POLYSACCHARIDE PRODUCED BY HAEMOPHILUS INFLUENZAE TYPE B
LINKED TO TETANUS TOXOID

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Background

Haemophilus influenzae type b is a Gram-negative bacterium causing pneumonia and meningitis. Its capsular polysaccharide - PRP is the main virulence factor. Vaccines based on PRP conjugated to protein are very efficient. The low yield of the conjugate has motivated the development of conjugation process to attend the demand of low-income countries. DMT-MM (4-[4,6-dimethoxy-1,3,5-triazin-2-yl]-4-methylmorpholinium chloride) used as activation molecule is an alternative to improve the conjugation reaction and the anti-PRP IgG level was statistically similar to commercial vaccine.

Objectives

To improve the conjugation process to make feasible the production of conjugate vaccine in large scale (g).

Methods

PRP (2.3g at 10g/L) was oxidized by NaIO₄, quenched with glycerol and purified by tangential ultrafiltration resulting PRP-Oxi. PRP-Oxi (1.9g at 8g/L) was mixed with adipic acid dihydrazide (ADH) followed by addition of NaBH₄CN. After 4 hours, NaBH₄ was added and submitted to ultrafiltration resulting PRP-AH. PRP-AH (15g/L) was reacted with TT (15g/L) in the presence of DMT-MM (0.1M) and the conjugate product was purified by size exclusion chromatography.

Conclusions

In large scale some adjustment were required in order to achieve the desired yield. For instance, glycerol amount was increased to stop the oxidation (83%). In the next step the amount of ADH was raised to avoid intermolecular reaction with 94% of PRP-AH. These yields are in agreement with those obtained in small scale (mg). The last step (mg scale) yielded 46% and now we are working on gram scale. Further studies considering immunological aspects will be performed.
Supported by BNDES (Nº 11.2.0322.1/2012)
Background
The development of bio-based production leads to an increasing demand for rapid engineering of multiple complex phenotypes into a single production host. The approach of genome shuffling is providing a powerful platform for improving multiple complex phenotypes in ill-characterized hosts.

Objectives
The main objectives of this study were to improve acid tolerance and lactic acid production from Lactobacillus rhamnosus through genome shuffling.

Methods
In the present study, the genome shuffling was used to improve the acid tolerance of while simultaneously enhancing lactic acid production. A total of 10 yoghurt samples were randomly collected in sterilized glass bottles from farmers directly and were processed immediately for isolation of the lactic acid bacteria. The isolated 5 strains of Lactobacillus rhamnosus treated for adaptation of Low pH. Adapted wild-type strain in pH 4.0 medium was then used as the parental, or starter, for genome shuffling. All the isolated Lactobacillus rhamnosus strains were mutagenized using nitrosoguanidine (NTG) while genome shuffling was carried out using standard method. From the results, it was observed that, the mutants showed a small increase, from 6.1 to 11.2 g/l, in production of l-lactic acid in comparison with the Wild type strain.

Conclusions
In conclusion, genome shuffling successfully improved the tolerance of L. rhamnosus towards acid. The research here demonstrated that genome shuffling could greatly accelerate the improvement of important phenotypes of microorganisms by developing their ability to circumvent the extreme process condition.
Background

Filamentous fungi in general and *Aspergillus* in particular are amongst the most important cell factories in biotechnology, as they have an intrinsic potential to produce a variety of organic acids such as citrate, gluconate and itaconate in high quantities. In order to obtain high yields of citric acid by *A. niger*, the elimination of Mn (II) ions from the medium is a prerequisite. Mycelia grown under manganese deficiency form dense and very small pellets. In this study, *A. terreus* NRRL 1960, an itaconic acid producer strain was used to study the effects of external manganese (II) ion concentration on fungal morphology.

Objectives

Our primary objective was to monitor time-course changes in the morphology of *A. terreus*, grown under different external Mn$^{2+}$ ion concentrations.

Methods

Well-controlled, strongly aerated lab-scale batch fermentations were performed. Changes in fungal morphology was monitored by image analysis. External Mn (II) ion concentrations were determined by ICP-MS.

Conclusions

The overwhelming mycelial morphology of *A. terreus* NRRL 1960 progressively turned to pellet-like below 50 µgL$^{-1}$ Mn$^{2+}$ ion concentration. At 5 µgL$^{-1}$ Mn$^{2+}$ ion concentration and below, morphology was completely dominated by yeast-like, swollen cells.
ANTIBIOTIC RESISTANCE AND VIRULENCE PROFILE OF BACILLUS PUMILUS
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Background
Bacillus pumilus present a biotechnological and pharmaceutical relevance highlighted by their ability to produce compounds with different biological activities. Moreover, they can be also found as troublesome contaminants in industrial settings and also be associated with food poisoning. Recently, phylogenetic analysis supported on different biomarkers determined the reassignment of most \textit{B. pumilus} as \textit{Bacillus safensis} or \textit{Bacillus invictae}. The absence of a resistance profile to antibiotics of human and veterinary importance and also of toxin encoding genes are pre-requisites specified by EFSA when their use is intended in the food sector.

Objectives
To evaluate the antibiotic susceptibility, and the presence of antimicrobial resistance or entero- and emetic-toxins encoding genes in a clonally diverse \textit{B. pumilus} group collection.

Methods
Minimum inhibitory concentrations to different antibiotics were determined and interpreted according to clinical breakpoints and microbiological cut-offs, in a collection comprising \textit{B. safensis} (n=27), \textit{B. invictae} (n=9) and \textit{B. pumilus} (n=4). Additionally, acquired genes encoding resistance to relevant human and veterinary antibiotics (tetracylines, macrolides, aminoglycosides, glycopeptides, phenicols and oxazolidinones) and also entero- and emetic-toxins (cytotoxin K, haemolysin, non-hemolytic and emetic toxins) were screened by PCR.

Conclusions
Susceptibility to all antibiotics included in interpretative criteria for their testing on \textit{Bacillus} spp. was observed. Nevertheless, high MIC values (8-64 mg/mL) were observed for cefotaxime, a therapeutically relevant antibiotic. All \textit{B. pumilus} group species revealed the absence of acquired antibiotic resistance and virulence genes, suggesting that there is no recognized risk of their application in the food sector or in other relevant biotechnological uses.
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BIOTECHNOLOGICAL POTENTIAL OF BURKHOLDERIA GLUMAE PG1

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Background

Microorganisms represent prolific sources for biotechnological relevant compounds which often need to be produced in heterologous hosts like Escherichia coli. Here, we present the Gram-negative bacterium Burkholderia glumae PG1 as a promising alternative production strain.

Objectives

B. glumae PG1 is known to secrete the lipase LipA which is used by the BASF SE for the production of enantiopure building blocks. A detailed genome comparison of B. glumae PG1 [1] with other representatives of this genus revealed that this phytopathogenic strain lacks the gene cluster encoding the biosynthesis of the important virulence factor toxoflavin.

Methods

Furthermore, we identified 25 clusters including PKS and NRPS for the production of secondary metabolites as well as genes encoding rhamnolipid biosynthesis enzymes resulting in the production of 20 mg/L rhamnolipids under non-optimized conditions.

We analyzed the genome of the industrial lipase production strain B. glumae LU8093 which was generated by random mutagenesis and identified 72 single nucleotide polymorphisms compared to the wild-type. Two of them were associated with the lipAB operon and affect its transcription as well as the efficiency of lipase secretion.

Conclusions
The identification of the putative SGNH hydrolase LipG and the construction and functional analysis of a T7 polymerase based expression strain further underline the biotechnological potential of *B. glumae* PG1.

OPTIMIZATION AND KINETICS OF THE PHYTOHORMONE PRODUCTION BY INONOTUS HISPIDUS IN STIRRED TANK REACTOR SCALE

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Background
In this study, firstly, the ability of plant growth hormones (PGH) production capacity of more than a hundred Basidiomycetes isolates was screened. Objectives
Plant growth regulators are commonly used by agricultural manufacturers in worldwide. In our study, we have tried producing some of these plant growth regulators from a macro fungi, Inonotus hispidus.

Methods
Inonotus hispidus was firstly recorded as a PGH producer species and was selected for following experiments based on its 1033.65, 125.11, and 10.91 mg/L gibberellic acid (GA3), absisic acid (ABA) and indole acetic acid (IAA) production values, respectively. Secondly, PGH producing capacity of Inonotus hispidus isolate was determined while the growth of the mushroom on different culture types (static, submerged and solid state). On 15th day of submerged fermentation, maximum plant growth hormones production was determined as 2478.85, 273.26, and 30.67 mg/L for GA3, ABA and IAA, respectively. Then, Plackett-Burman and response surface experimental designs were used for optimization of PGH production by Inonotus hispidus. Finally, the determined optimum conditions was investigated in stirred tank reactor scale to produce PGH in higher volume and reached results were, GA; 5441.54, ABA; 390.39 and IAA; 44.79 mg/L. PGH production by free and immobilized cells was also compared.

Conclusions
Consequently, production amounts of GA3, ABA and IAA by Inonotus hispidus in reactor scale were increased respectively 5.26, 3.12 and 4.1 fold according to screening step. The kinetic parameters of PGH production and growth of Inonotus hispidus (m, r, Qs, rp, Qp, Yp/s, Yp/x) were also determined in optimize and non-optimize conditions in Erlenmeyer flask and stirred tank reactor scales.
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DNA APTAMERS BLOCKING ACTIVITY OF ANTHRAX LETHAL TOXIN

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Background

Anthrax is major biosecurity threat. Apart from natural outbreaks, lethality of anthrax and recalcitrance of B. anthracis spores to environmental factors and disinfectants made it the most feared bioterrorist weapon.

Anthrax lethal toxin (LeTx) consisting of receptor-binding (protective antigen), and protease (lethal factor) subunits is main target for development of anthrax diagnostic and therapeutic tools. Aptamers represent important platform for discovery of target-specific affinity reagents. Particularly, aptamers known to be the source of efficient allosteric protease inhibitors [1]. Immuno-aptameric PCR possesses the highest known detection sensitivity [2].

Objectives

The purpose of the present study was to develop new technique aimed at rapid isolation of high-affinity aptamers suitable as diagnostic and therapeutic candidates against LeTx.

Methods

In the aptamer library screening, key elements to isolate high-affinity binders are: prevention of carryover of target-specific aptamers by nonspecific oligonucleotides, and early elimination of unbound and weakly bound species. Using LeTx as the target, we developed efficient, cost and time saving protocol of enabling isolation of high-affinity aptamers by using on-rate selection, tandem affinity purification of target-specific aptamers by double proteolytic elution that leaves tightest bound aptamers associated with the eluted target, whereas the majority of binders with intermediate affinity already dissociated and removed during the competitor-based off-rate selection.

Conclusions

New aptamer selection technique is useful in isolation of ultra-high affinity aptamers targeting bacterial toxins.

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References

SELECTION OF NANNOCHLOROPSIS OCEANICA CELLS WITH HIGHER INTRACELLULAR LIPIDS CONTENT BY FLUORESCENCE ACTIVATED CELL SORTING.

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Background
Microalgae are now the focus of interest due to their ability to produce an intracellular lipid feedstock that is suited to conversion in biodiesel. Nannochloropsis oceanica (N.O.) is a unicellular green alga intensely investigated because of the natural ability to form intracellular lipid bodies that contain triacylglycerol.
As a way to select the cells with spontaneous higher amount of intracellular lipids were separated those cells from a heterologous population using the technique of fluorescence activated cell sorting (FACS).

Objectives
Visualization of intracellular lipid droplets of N.O. by epifluorescence microscopy.
Visualization and selection of cells with high intracellular lipids content from heterologous populations of N.O. cultures by FACS.
Quantification of total lipids in selected cellular populations of N.O. under different culture conditions

Methods
The microalgal cultures where suspended in DMSO and stained with Bodipy 505/515. Flow Cytometer (BD Influx™ cell sorter) was used to detect the fluorescence signal of Bodipy. Neutral lipids stained showed a particular emission peak at 530 nm.
Total lipid fatty acid profiles were determined by GC of methyl esters.

Conclusions
Intracellular lipid droplets stained with Bodipy 505/515 was easily visualized by epifluorescence microscopy in N.O. cells.
The FACS technique allowed the visualization and separation of cellular populations enriched in intracellular lipids from liquid cultures of N.O.
Analysis of lipids quantification confirmed the obtaining of cellular populations with higher amounts of total lipids and fatty acids reaching 46% of total lipids by mg of dry weight.
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Fungicidal activity of exo- and endo-polysaccharides from Amazonian fungi

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Background:
Endophyte fungi are usually found in Amazon Rainforest with high biodiversity, and biological and chemical potential. They can be found as endophytes, phytopathogens or saprophytes. Candida is a fungal pathogen responsible for opportunistic infections in humans.

Objectives:
To produce exo- and endo-polysaccharide extracts from Pestalotiopsis sp. and Talaromyces sp., and assay them against Candida albicans.

Material and Methods:
Two fungi from the GEMMA group fungal collection were cultivated for 18 days on PDY medium. Then the mycelium was separated from the medium. After add to culture medium from each fungus 1L of methanol, the mixture was left for 48 hours at 4 °C. The precipitate formed was separated by centrifugation and the solvent in the supernatant was evaporated in rotary evaporator. The aqueous phase was subjected to liquid-liquid extraction with ethyl acetate and the acetate extract was concentrated by rotary concentration.

Conclusions:
Three samples were selected for candidical assays: (1) Talaromyce inorganic partitioned, (2) Pestalotiopsis mycelium, and (3) Talaromyce culture aqueous. All extracts had revealed candidical activity and were subsequently tested at the concentrations of 1000, 750, 500 and 250 μg.mL⁻¹ to determine the minimum inhibitory concentrations. The sample (1) and (3) showed no satisfactory inhibition while (2) showed inhibition at 500 μg/ml.
EXPRESSION, PURIFICATION AND CHARACTERIZATION OF A BIFUNCTIONAL 99-KDA PEPTIDOGLYCAN HYDROLASE FROM PEDIOCOCCUS ACIDILACTICI ATCC 8042

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Background
Lactic acid bacteria inhibit the growth of pathogens and food deleterious microorganisms because they produce substances with antibacterial activity, such as peptidoglycan hydrolases (PGH). *Pediococcus acidilactici* ATCC 8042 has been reported to inhibit pathogenic microorganisms such as *Staphylococcus aureus* through the production of two proteins with lytic activity, which are approximately 110 and 99 kDa. The 99-kDa PGH has high homology to a putative protein reported in *Pediococcus acidilactici* 7_4, where two different lytics domains have been identified but not characterized.

Objectives
The aims of this work were cloning, expression, purification and biochemical characterization this bifunctional enzyme.

Methods
The 99-kDa PGH was cloned and expressed successfully and showed activity against *Micrococcus lysodeikticus*. The protein was then purified using gel filtration chromatography. For the characterization 4-Nitrophenyl-N-acetyl-β-D-glucosamine was used as substrate.

Conclusions
The pure protein showed an optimal pH for antibacterial activity and stability of 6.0 and 5.0 to 7.0, respectively. The optimal temperature for activity was 60°C, and it lost all activity after incubation at 70°C for 1 h. The number of strains susceptible to the recombinant 99-kDa enzyme was lower to those lysed by the mixture of the 110- and 99-kDa PGHs of *P. acidilactici*, which indicates that there might by synergy and interactions between these two enzymes. This is the first peptidoglycan hydrolase from LAB that has been shown to possess two lytic sites. The results of this study will aid in the design of new antibacterial agents from natural origins that can combat foodborne disease and improve hygienic practices in industrial sector.
BIOCHEMICAL CHARACTERIZATION OF A SPORE LACCASE FROM A HALOTOLERANT BACTERIUM, BACILLUS SP.

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Background

Laccases, one of polyphenol oxidases, have many applications in different industries and can be used in bioremediation, organic synthesis, and biosensors.

Objectives

In present study, different bacterial strains were screened to find laccase producing strain. Biochemical characterization and Crystal violet decolorization effect of the spore laccase from a spore producing strain was examined.

Methods

As total 50 bacterial strains isolated from a soil sample of a chromite mine in Iran were screened for laccase production on nutrient agar medium supplemented with 3% (w/v) NaCl and 0.02% (v/v) guaiacol. Spores from a spore forming Gram-positive strain showed laccase activity. The strain was identified and pH and temperature optima for the enzyme activity were determined. Effects of inhibitors and metal ions on the enzyme activity were studied. Decolorization of Crystal violet was investigated by spore laccase.

Conclusions

The closest relative of the laccase producing strain was Bacillus safensis FO-036b(T). The spore laccase showed optimal activity at pH 5 and 35°C, had a good tolerance to metal ions, and retained 104% of its initial activity against 1 mM of NaN₃. The enzyme decolorized 64% of 10 mg/L Crystal violet, so it could be a good candidate to bioremediation applications.
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TO LOW COST FERMENTATION PROCESSES
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Background

Fermentation processes have been used for over 45 centuries, especially for the production of food, with ethanol and lactic acid as the main biogenic compound. These products have a high added value, because they are used as stimulant and/or because of their food preserving nature. This high added value allows significant costs for the fermentation process.

During the last 30 years fermentation technology has also been applied for the production of chemicals and fuels. These products have to compete with low-cost petrochemicals, putting restrictions on the costs of the fermentation processes. The current state-of-art fermentation technology is often too expensive for competitive production of chemicals and fuels.

In the same period the instruments to modify microorganisms have evolved considerably. The ability to build whole genomes, the application of –omic techniques and genome scale modelling give unprecedented possibilities to design microorganisms for optimal product formation.

Objectives

Instead of adapting the process to the microorganisms to reach the best results, it is now time to develop strategies which adapt the microorganisms to the process.

Methods

Key drivers for process improvement are the reduction of oxygen requirement by redox engineering, the increase of product tolerance and the separation of the product from the aqueous broth by phase separation.

Conclusions
In this paper/presentation we describe our activities to realize this and estimate the cost reduction that can be realized. Itaconic acid and fatty acids will be shown as examples. A cost reduction to a value below that of contemporary ethanol production (800 €/kg) seems feasible.
REACTIVE OXYGEN SPECIES REGULATE LOVASTATIN BIOSYNTHESIS IN ASPERGILLUS TERREUS

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Background

Lovastatin (LOV) is a secondary metabolite, produced by Aspergillus terreus. It has great commercial importance since it lowers cholesterol levels in blood. Studies from our group, showed a link between reactive oxygen species (ROS) and LOV biosynthesis in submerged (SmF) and solid-state fermentation (SSF). Our results showed that sod1 gene (oxidative stress defense enzyme) was intensely expressed during rapid growth phase (or trophophase) in LOV fermentations, but it was down regulated in production phase or idiophase. Probably because of this, in that moment ROS levels increased, generating an oxidative state during idiophase (Miranda et al 2013).

Objectives

The objective of the present work was finding evidence that ROS contribute to the regulation of lovastatin biosynthesis.

Methods

ROS were manipulated during LOV fermentations. Exogenous antioxidants were used to eliminate or reduce ROS accumulation during lovastatin fermentations. Its effect on lovastatin production was determined. Also, its effect on the expression of LOV genes (lovE and lovF), and on genes yap1 and srrA (encoding oxidative stress defense transcription factors) was determined by Northern analysis.

Similar (complementary) experiments, increasing ROS by H2O2 addition to LOV fermentations, were also performed.

Conclusions

1. ROS accumulation in idiophase is not only necessary for the normal production of lovastatin, but contributes to the transcriptional regulation of the biosynthetic genes, including the specific regulatory gene lovE.
2. ROS regulation could be mediated by oxidative-stress-response transcription factors: Yap1 might be acting as a negative regulator; while SrrA could play an important role in positively controlling lovastatin production.
Background
Endophytes are microbes that colonize living, internal tissues of plant without causing any immediate of negative effect. Various studies have shown that endophytes are endowed with great potential for human welfare like having potential in the form of antibiotics, antiviral, anti asthamatic, antioxidant and anticancer compounds. Endophytes from medicinal plants have become a hot topic for metabolite discovery. There is still a lot of potential of endophytes which need to be discovered for pharmaceutical applications. Keeping these things in consideration, the aim of the present study was to isolate fungal endophytes from different accessions of medicinal plant *Withania somnifera* and to screen the endophytes for the production of various bioactive molecules.

Objectives
To isolate the endophytic microorganisms from *Withania somnifera* (Ashwagandha) & screen for antimicrobial activity against a wide range of pathogenic microorganisms

Methods
Isolation of endophytes using the standard methodology published earlier

Conclusions
Out of 7 endophytes, four isolates were identified by ITS amplification and sequencing. All the four isolates showed significant similarity with *Alternaria alternata, Alternaria tenuissima strain, Alternaria sp.* and *Alternaria compacta.*

The isolated endophytes were also screened for antimicrobial activity against a wide range of pathogenic microorganisms and after analyzing the results some of the endophytes showed promising antimicrobial activities.

Our results confirm that the isolated endophytes could serve as an alternative source of various bioactive molecules of novel structure and function as endophytes are still largely unexplored.
SECRETION AND FUNCTIONAL DISPLAY OF FUSION PROTEINS THROUGH THE CURLI BIOGENESIS PATHWAY

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Background
Curli are functional amyloids expressed as fibres on the surface of Enterobacteriaceae. Contrary to the protein misfolding events associated with pathogenic amyloidosis, curli are the result of a dedicated biosynthetic pathway. A specialized transporter in the outer membrane, CsgG, operates in conjunction with the two accessory proteins CsgE and CsgF to secrete curlin subunits to the extracellular surface, where they nucleate into cross-beta strand fibres.

Objectives
Here we investigate the substrate tolerance of the CsgG transporter and the capability of heterologous sequences to be built into curli fibres.

Methods
We genetically fused different proteins to the C-terminus of csgA.

Conclusions
Non-native polypeptides ranging up to at least 260 residues were exported when fused to the curli subunit CsgA. Secretion efficiency depended on the folding properties of the passenger sequences, with substrates exceeding an approximately 2 nm transverse diameter blocking passage through the transport channel. Secretion of smaller passengers was compatible with prior DsbA-mediated disulphide bridge formation in the fusion partner, indicating that CsgG is capable of translocating non-linear polypeptide stretches. Using fusions we further demonstrate the exported or secreted heterologous passenger proteins can attain their native, active fold, establishing curli biogenesis pathway as a platform for the secretion and surface display of small heterologous proteins.
Background

In the recent years, the applicability of actinomycete strains as probiotics for plants, serving as agents of biological control of plant diseases, has been actively investigated.

Objectives

415 strains of extremophile actinomycete strains from the unusual ecosystems of Kazakhstan.

Methods

Antifungal properties of extremophilic actinomycetes were studied against the fungal pathogens of grain cultures (genera Fusarium and Aspergillus) by agar diffusion method.

Conclusions

104 actinomycetes strains from the extreme ecosystems of Kazakhstan (solonchak, takirs and takir-type soils, solonetz and solod) showed antifungal activity against the fungal pathogens of grain cultures (Fusarium sp., Aspergillus niger). 20 strains of extremophilic actinomycetes have been selected, capable of growing at a low content of nutrients in the medium, in the presence of high salt concentrations (≥ 5%) and pH level (≥ 9.0). They have the ability to inhibit the growth of phytopathogenic fungi (diameter of inhibition zone 20-30 mm) in the different environmental conditions - neutral, in the presence of high NaCl concentrations (5% or more), and high pH values (9-11). This practically valuable feature is especially important for the plant cultivation since many biopreparations developed for neutral ecological niches become useless after salinization or alkalinization of the soil. Scientific novelty of the research results lies in the use of biotechnological potential of extremophilic actinomycetes in the development of biological preparations for plant cultivation under various environmental conditions.
Background

A major challenge in the sustainable production of biofuels and biochemicals is efficient enzymatic conversion of plant biomass into monomeric sugars. Most enzyme mixtures are currently produced by a small selection of fungal species (e.g. *Trichoderma reesei*, *Aspergillus niger*). However, the fungal kingdom holds many more fungal species which produce enzyme mixtures with beneficial characteristics such as high (hemi-)cellulase activity and high thermostability. *Sordariales* is one of the few fungal orders with thermophilic isolates, of which many have been associated with the production of thermostable enzymes.

Objectives

The aim of our study is to assess the diversity within *Sordariales* for efficient plant biomass degradation.

Methods

Phylogenetic analysis and growth analysis revealed that optimal growth temperature is a polyphyletic trait within *Sordariales* with separate mesophilic, thermotolerant and thermophilic clades. Four thermophilic clades were clearly distinguished: *Myceliophthora* species, *Thielavia terrestris*, *Chaetomium thermophilum*, and *Mycothermus thermophilus*. Thermophiles within *Myceliophthora* showed the most potential as efficient plant biomass degraders. Especially *M. heterothallica* had good growth on a large range of substrate and was able to produce offspring with a large physiological and genetic variety. Crossing and selection strategies were used to further improve *M. heterothallica* in degrading biomasses such as sugarbeet pulp and spruce. The mechanisms behind improved biomass saccharification were understood by proteomics and genome analysis.

Conclusions
This study showed the strategic strength of combining fungal diversity with specific selection strategies to find enzyme mixtures with interesting industrial properties.
PREPARATION OF ACIDIC AMINOPHOSPHONATES USING FUNGI

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Background

Biotransformation is substrate transformation to desired product by using suitable kind of biocatalysts. Nowadays, it is an alternative tool for the development sustainable technologies for the production of chemicals and drugs. Aminophosphonates are analogues of amino acids in which a carboxylic moiety is replaced by phosphonic functionality or related groups. They are compounds with stable carbon to phosphorus bond and have wide range of biological activities and variety of applications in industry [Żymańczyk-Duda, Phosphorus, Sulfur and Silicon, 2008, 183; Mucha A., J. Med.Chem. 2011, 54]. Biosynthesis of chiral, organophosphorous compounds is still not fully explored field of science.

Objectives

The aim of work was selection of the whole-cell biocatalysts with hydrolytic activity for preparation of phosphonate analogues of aspartic acid. Thus, biocatalytic methods, which allowed obtaining derivatives of aminophosphonic acids via biohydrolysis of lactam ring were elaborated.

Methods

To obtain phosphonic analogue of aspartic acid via biohydrolysis of O,O-dimethyl-4-oxoazetidin-2-ylphosphonate, whole cells of P. minioluteum were used as a biocatalyst. NMR spectra interpretation allowed postulating possible mechanism of presented microbial conversion as amide bond hydrolysis. Biohydrolysis was only achieved in deionized water as bioconversion medium. Application of another reaction media caused partial splitting of the substrate during biotransformation process. The separation process required reverse phase chromatography to obtain product from biotransformation mixture.

Conclusions

Presented method is simple and quite efficient. Application of P.minioluteum as a biocatalyst resulted in production of desired product with moderate yield. It means, that process is a good starting point for further scaling up the process.
SELECT ANTIMICROBIAL PEPTIDES (AMPS) AS BACTERICIDAL AGENTS FOR EX VIVO STORED HUMAN PLATELETS

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Background
Bacterial contamination of ex vivo stored human platelets and the transmittance of sepsis by platelet transfusion represent the highest infectious disease risk in transfusion medicine today.

Objectives
An alternative approach to detection of contaminated units is pathogen reduction technology that directly inactivates or prevents the replication of pathogens present in platelet products. We have developed a novel approach to pathogen reduction, one that promises to be more specific and less damaging to the platelet product and at the same time effective against 6 bacterial species common to transfusion of platelets.

Methods
The methodology is based on treatment of platelets with small synthetic peptides named PD1-PD4 derived from the thrombin-induced human platelet-associated antimicrobial proteins, and repeats of Arg-Trp (RW1-RW5), and their selected combinations. This treatment demonstrated microbicidal activity against Staphylococcus aureus, Staphylococcus epidermidis, Escherichia coli, Pseudomonas aeruginosa, Klebsiella pneumoniae and Bacillus cereus in the spiked platelet samples. We also demonstrated that AMP treated platelets maintain their in vitro properties during 7 days of storage. Preclinical evaluation of peptides PD3, PD4, and RW2-RW5 demonstrated that these AMPs do not adversely affect the human platelet recovery and survival in a SCID mouse model. We have further shown that the peptides are not immunogenic in rabbits, and do not affect platelet performance in in vitro tests.

Conclusions
The body of work presented here provides a proof of a concept that AMPs could be incorporated into a novel platelet-sparing pathogen reduction treatment and warrants further study.
STABLE VECTORS FOR EFFICIENT GENE EXPRESSION IN RALSTONIA EUTROPHA H16

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Background
The Gram-negative β-proteobacterium Ralstonia eutropha H16 is primarily known for polyhydroxybutyrate (PHB) production and its ability to grow chemolithoautotrophically by using CO₂ and H₂ as sole carbon and energy sources. R. eutropha H16 has also attracted significant interest for its ability to metabolize heavy metals, to degrade a variety of chloroaromatic compounds or chemically related pollutants.

Objectives
The use of R. eutropha H16 as a production organism under chemolithoautotrophic growth conditions is favored. However, inducible expression systems are yet required for gene expression in R. eutropha H16. The aim of this work was to identify inducible expression systems for the use in R. eutropha H16 facilitating gene expression.

Methods
A family of efficient and highly stable plasmid expression vectors was created for the use in R. eutropha H16 by applying various biosynthetic methods.

Conclusions
The use of promoters derived from bacteriophage T5 was described in this work and hereby the J5 promoter proved to be the strongest promoter yet to be applied in R. eutropha H16. Moreover, the implementation of the RP4 partition sequence in plasmid design increased plasmid stability significantly and enables fermentations with marginal plasmid loss of recombinant R. eutropha H16 for at least 96 h. The utility of the new vector family in R. eutropha H16 is demonstrated by providing expression data with different model proteins and consequently further raise the value of this organism as cell factory for biotechnological applications including protein and metabolite production.
SCREENING FOR CELLULOLYTIC ACTIVE NOVEL ANAEROBIC FUNGI  
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Background

Lignocellulosic plant material is degraded by anaerobic fungi (AF) with their rhizoid and various free and cellulosome-bound lignocellulolytic enzymes. AF could thus be used to improve biogas production from fiber-rich residues. AF were isolated from faeces of animals living on a high fiber diet. Screening them for the cellulolytic most active AF with conventional tests are labor intensive and time consuming, a quantitative Real-Time PCR (qPCR) routine for glycosyl hydrolase family 5 (GH5) endoglucanase should be developed as a suitable alternative.

Objectives

New anaerobic fungal isolates should be checked and compared for their cellulolytic potential by parallel determination of GH5 mRNA production and conventional parameters. Thus anaerobic fungi with outstanding cellulolytic abilities shall be identified.

Methods

To identify the best moment for mRNA extraction, growth curves of the isolates were recorded, and mRNA was extracted at several points. Additionally gas pressure was measured to see if mRNA production, growth and gas pressure developed accordingly. Reverse-transcribed cDNA was quantified in qPCR with primer system AF Endo designed to specifically detect GH5 endoglucanase of AF.

Conclusions

Novel AF isolates were achieved from a Cameroon sheep, a Kiang, an alpine goat and a domestic yak. In experiments carried out with two of the isolates the GH5 RT-qPCR system tested successfully. GH5 mRNA production was maximal after 18h with similarly developing growth, gas pressure and mRNA level. Further AF species are being tested for their cellulolytic activity. A future study shall compare gained results with the real life cellulolytic ability of tested AF.
INVESTIGATION OF SYNTHETIC LIGNIN BIOMODIFICATION BY PHANEROCHAETE CHRYSOSPORIUM: ANALYSIS OF STRUCTURAL CHANGES OF SYNTHETIC LIGNIN AND EFFECT OF MICROBIAL ENZYMES

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Background
These days, there has been a great interest in lignin as one potential renewable source for conversion into chemicals and materials. Lignin is complex and amorphous polymers, therefore, it needs to be modified to meet properties required for industrial uses.

White rot fungi secreting ligninolytic enzymes are promising for enzymatic modification, because they can degrade lignin selectively and more rapidly than other microbes. Therefore, comprehending of biomodification process by white rot fungi is necessary for industrial application of microbial treated lignin.

Objectives
The biomodification of synthetic lignin by white rot fungus, Phanerochaete chrysosporium, was investigated in this study. The fungal secretomes and structural changes of the fungal treated synthetic lignin were examined.

Methods
Synthetic lignin used in this study was synthesized from coniferyl and sinapyl alcohol. This polymer, dehydrogenative polymer (DHP) was used as substrate to elucidate the mechanism of microbial modification of lignin.

Detailed structural analysis of lignin were conducted as follows: phenolic hydroxyl group of DHP was analyzed by aminolysis reaction, and the molecular weight and changes of ether bonds in DHP was analyzed by gel permeation chromatography and nitrobenzene oxidation method, respectively. For examining the effects of fungal secretomes in biomodification of synthetic lignin, protein contents by using protein chip and activities of ligninolytic enzymes were measured, respectively.

Conclusions
Consequently, this study is expected to help to understand the biomodification mechanism of synthetic lignin by P. chrysosporium, and to evaluate availability of P. chrysosporium in biomodification of lignin for value added application of lignin.
HOMOLOGOUS AND HETEROLOGOUS EXPRESSION OF DEHYDROGENASES AND OXIDOREDUCTASES OF RALSTONIA EUTROPHA H16

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Background

*Ralstonia eutropha* is a Gram-negative, strictly respiratory facultative chemolithoautotrophic bacterium which can use H₂ and CO₂ as sole sources of energy and carbon in the absence of organic substrates. It has attracted great interest for its ability to degrade a large list of chloroaromatic compounds and chemically related pollutants. Furthermore it was already applied for the production of biodegradable polymer polyhydroxyalkanoates on an industrial scale. *R. eutropha* serves as a model organism for the mechanisms involved in the control of autotrophic carbon dioxide fixation, hydrogen oxidation and denitrification.

Objectives

We are interested in establishing specialized *R. eutropha* based cell factories by genetic engineering. The particular interest is constructing cells efficiently performing oxidoreductase reactions by overexpression of homologous and/or heterologous enzymes. One of the main types of oxidoreductase reactions is performed by dehydrogenases, which have a wide range of possible biotechnological applications.

Methods

A selection of alcohol dehydrogenases as well as short chain dehydrogenases of *R. eutropha* H16 was cloned and expressed in native versions in *E. coli* as well as in *R. eutropha* H16 in order to study basic functional capability of these enzymes. For each of the enzymes cofactor specificity and the substrate range, both for oxidation and reduction way was analysed in a photometric assay by the detection of NAD and thereby specific activity units were calculated.

Conclusions

Two of the tested enzymes showed high selectivity toward s-enantiomers for all tested secondary alcohols with NADP⁺ as preferable cofactor. Furthermore a difference in enzyme activity depending on the expression host could be observed.
METHANOTROPH-CATALYZED BIOCONVERSION OF METHANE TO METHANOL

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Background

Methane has attracted much attention as the alternative chemical feedstock and conversion of methane to methanol has attracted enormous interest. Indirect chemical conversion of methane to methanol is associated with a number of problems including low energy efficiency and high capital cost.

Objectives

We investigated a batch methane-to-methanol conversion in high yield using Methylosinus trichosporium OB3b whole cells. Cultivation and conversion relevant parameters including air/methane ratio and copper concentration were optimized.

Methods

M. trichosporium OB3b was cultured in a 600 mL joint flask with rubber septum containing 200 mL NMS medium with supply of 7:3 air/methane ratio at 30 °C. Various chemical inhibitors for methanol dehydrogenase were screened and added to accumulate methanol. Sodium formate was added as a reducing power source to maintain methane monoxygenase activity.

Conclusions

In the presence of 100 mM potassium phosphate as MDH inhibitor, 0.4 g/L methanol was accumulated and more than 70% (mol methanol/mol methane) of methane was converted to methanol. Volumetric productivity of 0.049g/L/h was obtained in the batch reaction with M. trichosporium OB3b as the biocatalyst. Biocatalytic conversion using methanotrophic microorganism will offer an energy-efficient method in commercialization of methane-to-methanol.

References

Background

Increased amino acid requirement of malignant cells is exploited in metabolic antitumor therapy, e.g., enzymotherapies based on asparagine or glutamine deprivation. L-Asparaginase (L-Asparagine amidohydrolase; EC 3.5.1.1) which catalyzes the conversion of L-Asparagine to aspartic acid and ammonia is one of promising enzyme in pharmaceutical industries. A wide variety of microorganisms have L-asparaginase. Halophilic bacteria may contain L-asparaginase with novel immunological properties that can be used in hypersensitive patients because of modified surface structures.

Objectives

The main objective of this study was to optimize the culture conditions for L-asparaginase production by selected halophilic strain.

Methods

The effect of different concentrations of NaCl and glucose ranging from 0-10% and 0-1% respectively and the influence of different carbon sources such as saccharose, maltose, glucose and fructose on the enzyme production was evaluated by varying one factor at a time. The effect of pH (5–9) and temperature (25-35-45 °C) were also studied. After 72h incubation, the samples were collected by centrifugation for the determination of L-asparaginase activity by the nesslerization method.

Conclusions
A halophilic bacterium *Vibrio* sp. Strain GBFx3 which was isolated from Gomishan Lake at Golestan province of Iran was screened for L-asparaginase activity. The highest amount of L-asparaginase production was observed in 0.5% glucose or 0.5% saccharose as sole carbon source, 2.5% NaCl, 34°C and pH 8 which were 0.63, 0.5, 0.4 and 0.43 IU/ml respectively.
DEVELOPMENT OF A POLYFUNCTIONING PROLONGED-ACTION BIOLOGICAL PREPARATION FOR CROP PRODUCTION BASED ON THE RHIZOSPHERE BACTERIAL STRAINS WITH ANTIFUNGAL AND PLANT-GROWTH STIMULATING PROPERTIES

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Background
Associations of the rhizosphere bacterial strains belonging to the genera Pseudomonas and Azotobacter simultaneously possess both antifungal and plant-growth stimulating properties and are the basis for new trends in the development of biological preparations for crop production.

Objectives
Strains Pseudomonas R-7 and Azotobacter A-6 isolated from the rhizosphere of tomato crops cultivated in Kazakhstan.

Methods
Antagonistic properties of the bacteria were studied using agar diffusion method against the pathogens of fungal diseases in tomatoes of the genera Phytophthora, Fusarium, Alternaria, Cladosporium. Growth-stimulating properties of bacteria were studied by inoculation of tomato seeds with a culture liquid of bacteria with a cell titer of n x10⁹.

Conclusions
Strain Pseudomonas R-7 showed antagonism against phytopathogenic fungi: Phytophthora sp. - 19 mm, Fusarium sp. - 16 mm, Alternaria sp. - 23 mm, Cladosporium sp. - 18 mm. When using a strain Azotobacter A-6, the growth inhibition zone for Phytophthora sp. was 22 mm, Fusarium sp. - 24 mm, Alternaria sp. - 23 mm, Cladosporium sp. - 20 mm. Strain Azotobacter A-6 also enhanced the tomato germination capacity by 22.0% as compared with the control, increased the plant height by 15.5%, root length by 26.3%, plant mass by 15.0 %, and number of leaves by 33.3%.

A composition of the new biological preparation for crop production has been formulated based on the strains of rhizosphere bacteria Pseudomonas P-7 and Azotobacter A-6, compatibility of the strains under cultivation and storage examined, optimal ratio of bacterial microorganisms and soil extract in the combined culturing selected, ensuring the continued viability of cells within a scheduled storage period.
CLONING AND OVEREXPRESSION OF A NEW LIP GENE FROM STENOTROPHOMONAS MALTOPHILIA PSI-1 - PURIFICATION AND PARTIAL CHARACTERIZATION OF THE RECOMBINANT LIPASE

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Background

Lipases catalyze esterification, interesterification and transesterification reactions and have gained serious attention for the production of sustainable biofuels, motivated by environmental concerns combined with the depletion of fossil fuels.

Objectives

The aim of this study was the isolation of new strains bearing high lipase activity in order to exploit their lipases in biofuel production.

Methods

42 environmental strains, isolated from sludge of a waste treatment facility (Psittaleia, Greece), were screened for the presence of lipase activity, using a rhodamine B procedure. One isolate, identified as Stenotrophomonas maltophilia (named Psi-1) by means of IGS-based molecular taxonomy, was selected for further analysis. A 1.203 bp amplicon of Psi-1 corresponding to a putative secreted lipase gene (as revealed by bioinformatic analysis) was cloned and overexpressed in a pET29c - E. coli BL21(DE3) system. The recombinant protein was purified using Ni-NTA chromatography. Lipase activity was verified by in-gel assays (zymograms) and photometric biochemical assays.

Conclusions

A novel lipase gene from a new Stenotrophomonas maltophilia isolate was identified, cloned and overexpressed. The recombinant enzyme was proved to exhibit lipase activity and has a potential as a catalyst for biofuel production.

Acknowledgments
This project is co-financed by the European Union (European Regional Development Fund — ERDF), through the operational programs for "competitiveness and entrepreneurship" and regions in transition 'Cooperation 2011 — Partnerships of Production and Research Institutions in Focused Research and Technology Sectors”, of the National Strategic Reference Framework (NSRF) 2007–2013, and the Hellenic Ministry of Education, Lifelong Learning and Religious Affairs — General Secretariat for Research and Technology.
Background

The use of enzymes as biocatalysts for the preparation of optically pure compounds is becoming a common method especially in the pharmaceutical industry [Nestl B.M., 2011, *Curr Opin Chem Biol*]. Obtaining enantiopure active compounds with a defined configuration appeared to be more desirable, not only because of effectiveness of such pharmaceuticals, but also because of safety requirements [Ogawa J., 2002, *Curr Opin Biotech*]. Immobilized whole-cells are one of the biocatalyst form that can be used in biotransformation processes. This system allows reusing biocatalyst, facilitates downstream processes and enhance the stability of the enzyme. Moreover, use of immobilized catalyst in flow reactor allows carrying out continuous processes.

Objectives

The aim of the study was the kinetic resolution of a racemic mixture of 2-butyryloxy-2-(ethoxy-P-phenylphosphinyl)acetic acid by biotransformation using immobilized whole-cell biocatalyst. *Penicillium oxalicum* has been reported as an active biocatalyst in the hydrolysis of tested compound in shake flask system [Serafin M., 2014, *Chemik*]. Optically pure products can find application as chiral discriminator and building block of defined absolute configuration in synthesis of biologically active compounds such as pharmaceuticals or pesticides.

Methods

The activity of biocatalyst, cells immobilized in two ways, was tested toward racemic mixture of starting compound. Products were analyzed by nuclear magnetic resonance spectroscopy $^{31}$P NMR with the addition of quinine as a chiral solvating agent.

Conclusions

Simple and effective method of whole-cell immobilization was elaborated. Activity of biocatalyst was confirmed by hydrolysis of tested compound with 50% of conversion.
degree and high enantiomeric excess. Developed system allowed to enlarged the scale of the process.
Background
The filamentous fungus Aspergillus niger has a long tradition of safe use in industry in the production of enzymes and organic acids. Furthermore, it can grow on a wide range of substrates under various environmental conditions and has an optimized pathway towards producing organic acids, as indicated by its high production titers of citric acid in submerged fermentations (>300 g/L). These features make it an interesting host for the production of itaconic acid (IA), a high-value building block chemical with a broad application potential in the chemical industry. We have successfully identified the IA production pathway of Aspergillus terreus and engineered it into A. niger. The resulting A. niger strain is able to produce IA at low levels.

Objectives
In order to increase the yield we have tested various medium compositions and fermentation conditions. This resulted in an optimization of the fermentation medium and cultivation conditions, accompanied with an enhancement of yield and titer of IA production. Moreover, recently we have discovered a novel pathway found, to our knowledge, exclusively in A. niger that enables higher productivity, titer and yield of IA. This novel pathway provides a platform for further improvement of organic acid production in A. niger.

Methods
Fungal transformation and fermentation

Conclusions
This novel pathway enabled increased IA acid productivity, yield and titer.
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COMPARATIVE GENOMICS APPROACH FOR BUILDING A COMPLETE AGAROLYTIC SYSTEM
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Background
In a marine carbon cycle, the metabolic pathway of agar is not fully understood. Agar is a recalcitrant hetero-polysaccharide. Many agarases are known to fragment agar polymer into disaccharide unit, neoagarobiose, that is split into two monomeric sugars (D-galactose and 3,6-L-anhydrogalactose). Metabolic fate of 3,6-L-anhydrogalactose is still mysterious.

Objectives
We aim to elucidate full repertoire of agarolytic system in nature not only for understanding the marine carbon cycle but also for construct a synthetic agarolytic system for biomass utilization.

Methods
Using experimentally verified functional agarolytic genes as a probe, we searched all available microbial genomes in the public databases and collected all microbial genomes having potential agarolytic activity. In comparative genomics approach, we combined three distinct procedures such as ortholog prediction, gene cluster analysis and transcriptomic analysis (RNAseq) of three genomes, Saccharophagus degradans, Marinimicrobium agarolyticum and Vibrio sp. EJY3, we predicted potential genes related to agar metabolic pathways.

Conclusions
The predicted gene set from comparative genomics was divided into several functional categories: i) metabolic enzymes, ii) regulatory proteins (e.g. transcription factor), iii) signaling proteins and transmembrane proteins. We identified phylogenetically conserved genes and designated those genes as the core gene set. By functional examination, we revealed new genes essential for 3,6-anhydrogalactose and designed a synthetic E. coli system metabolizing agar as a sole carbon source.
Fungal strain development for improved protein production: protease mutant approaches and genome mining for novel enzyme discovery.

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Background

Optimized protein production in filamentous fungi requires the availability of fungal strains with low levels of secreted protease activity in order to improve secreted protein levels, including shelf life. Already for several decades research has been carried out to obtain these type of mutants, leading to the isolation of mutants with very favorable characteristics, one being a mutation in a transcriptional regulatory gene, prtT (e.g. Punt et al.;, 2008).

Objectives

Based on these results further improved strains have been developed using different selection approaches.

Methods

Controlled fermentation experiments with selected mutant strains revealed different improved characteristics, whereas full genome sequencing was carried out to identify the genetic basis of the mutant phenotypes. Identification of relevant protease-regulatory genes has also been carried out using collections of regulatory gene knock-out strains in N. crassa and A. niger.

Conclusions

Several examples of the use of selected mutant strains in our research to discover and produce novel hydrolytic enzymes will be presented.
DIRECT ETHANOL PRODUCTION FROM BAGASSE CLOSTRIDIUM CELLULOLYTICUM
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Background
In recent years the direct process of ethanol production from lignocellulosic materials, has been considered greatly due to its efficiency and money-saving manner of manufacturing. Some Clostridium species such as C.thermocellum sp. and C.cellulolyticum sp. has received many attentions because of their extracellular enzyme production ability.

Objectives
The ultimate goal of present study is evaluating the ethanol production ability of C.cellulolyticum from untreated bagasse and delignified bagasse as cellulose source. The ability of C.cellulolyticum H10 for direct consumption of cellulose, eliminate the hydrolysis processes and lowered the end cost of ethanol production and also decreased the environmental threats of the existing processes.

Methods
In this study the bacterial strain was cultured in modified CM3 culture medium in anaerobic condition and after 20 days the produced ethanol was determined by gas chromatography, then cellulose was replaced with untreated bagasse and the ethanol production was examined. Untreated bagasse was delignified with three different pretreatment methods which include sodium hydroxide, sodium chlorite and acetic/nitric acid and the delignified outputs of all these delignification methods were used as carbon sources for bacterial growth and ethanol production were determined.

Conclusions
Acetic/nitric acid delignification methods showed the highest delignification value with 86.7% lignin content decrease and the highest ethanol per cellulose yield with values of 0.19 (g ethanol/g cellulose).
LOW CYTOSOLIC PH AS A SIGNAL FOR CELL SURVIVAL IN H2SO4 STRESSED CELLS
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Background
In the process of bioethanol production yeast biomass is recycled using treatment with dilute sulfuric acid to control the bacterial population. This treatment can lead to loss of cell viability, with consequences on the fermentation yield.

Objectives
To understand and ideally enhance yeast cell survival in these conditions, we analyzed the functional cellular responses of S. cerevisiae to inorganic acid stress.

Methods
Was analyzed growth and intracellular pH (pHi) in low pH conditions, using the pH-sensitive GFP derivative ratiometric pHluorin expressed in the cytoplasm of the strains.

Conclusions
S. cerevisiae mutants in cell wall integrity (CWI) and calcium signaling have reduced growth and viability at a low external pH (pHex) of 2.5. WT yeast showed evident defect in growth only at a pHex of 1.5, showing that a functional cell wall biogenesis is crucial for survival. To our surprise, the CWI mutant showed enhanced viability at pHex of 1.5 compared to pHex 2.5. While at a pHex of 2.5 the pHi was unaffected compared to pHex 5.0, at pHex 1.5 the pHi of a CWI mutant was reduced. This reduction by itself did not lead to loss of viability. Rather, artificial reduction of pHi at pHex 2.5 rescued the CWI and Ca^{2+} signaling mutants. Also, the lethal effect of low pHex on CWI and Ca^{2+} mutants takes place only in growing cells. However, a lowered pHi leads to a reduction in growth rate, and thus protects the cells from death. Likely, cytosolic pH is a signal that directs the growth-stress tolerance trade-off in yeast.
PRELIMINARY STUDY ON ANTIMICROBIAL ACTIVITY OF BACTERIAL SIDEROPHORE

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Background

Bacteria must acquire iron by competing with environmental chelation. One mechanism for bacterial iron acquisition utilizes siderophores. These molecules are small-molecule chelating agents that have been isolated from many microorganisms. As a current study, siderophore is used to deliver an antibiotic payload, through production of siderophore linked antibiotics e.g. salmycins.

Objectives

In this study we aimed siderophore production from soil bacteria and detection of their antimicrobial activity.

Methods

Bacteria was isolated from soil samples. For detection of siderophore existence in these bacteria, Cas agar medium was used. Identification of siderophore producer bacteria were used 16s rRNA analysis. The crude siderophore was produced and extracted from different soil bacteria. Antimicrobial effect of siderophore was determined on several pathogens (Candida albicans ATCC 10231, Pseudomonas aeruginosa 27853, E.coli 25922, Staphylococcus auerus 25923, Bacillus subtilis 6633, Proteus mirabilis, Enterobacter aerogenes).

Conclusions

As a result of study, we were observed that siderophore from soil bacteria have antimicrobial effect on pathogen microorganisms. Nowadays, these studies are taught as a new methods against the growing antibiotic resistance. Mimicry of Iron-uptake dependent mechanism will can be new approach in medicine.
Background
Solvent tolerant strains of the ubiquitous soil bacterium *Pseudomonas putida* are increasingly important for a wide range of industrial biotechnology applications. Because of its remarkable solvent and stress tolerance and its energy efficiency, *P. putida* S12 has been described as an important strain for production of value added aromatic compounds and for efficient whole cell biotransformation of HMF to FDCA (1, 2, 3, 4).

Objectives
We have undertaken detailed genome sequence analysis to identify and understand the background of solvent tolerance and energy efficiency in *P. putida* S12.

Methods
Whole genome sequencing was performed through Illumina HiSeq and PacBio RSII sequencing. Assembly was performed using the CLC Genomic Workbench (BLASR, SSPACE-LongRead scaffold, Gapfiller version 1.10) and manual closure of remaining gaps. The completed genome sequence was subject to automated annotation by the NCBI Prokaryotic Genome Annotation Pipeline with manual annotation of prominent *Pseudomonas* features.

Conclusions
The full *P. putida* S12 genome consists of a 5.8 Mb circular genome and a 557 kb circular megaplasmid. Both the genome and the megaplasmid contain various mobile elements among which multiple copies of the ISS12-type repeat sequence (5). Location of ISS12 repeats and number variation in various strains indicate a role in acquisition and maintenance of solvent tolerance. In addition, gene composition of the megaplasmid indicates an essential role in instigation of solvent tolerance and resistance to other stress conditions. Further comparative analysis of the S12 genome will be presented together with functional analysis of predicted gene and
metabolic functions.

References

EFFECT OF SILENCING ASPERGILLUS TERREUS YAP1 GENE, ON REACTIVE OXYGEN SPECIES (ROS) ACCUMULATION PATTERN AND LOVASTATIN BIOSYNTHESIS.

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Background

In an earlier work we found a ROS build up during lovastatin production phase, which coincided with sod1 gene (oxidative stress defense enzyme) down regulation. Later, we showed that ROS regulate lovastatin biosynthetic genes. The mechanism by which ROS induce these genes is unknown, but it is considered to be through oxidative-stress-response transcription factor(s). Expression profiles suggested that A. terreus Yap1 is a candidate, although probably as a negative regulator. These findings could be used to design new genetic improvement methods.

Objectives

Study the role of Yap1 in the control of ROS and in lovastatin biosynthesis, in liquid submerged fermentation (SmF) and in solid-state fermentation (SSF). Also investigate if the perturbation in the ROS profile (caused by silencing yap1) can increase lovastatin production.

Methods

Silencing vector was constructed by ligation of a fragment of At yap1 gene to pGdpPki-RNAi vector. A. terreus TUB F-514 transformants were characterized. Lovastatin was quantified by HPLC, and ROS concentration by diclorofluoresein and gene expression by Northern Blot.

Conclusions

1) yap1 silencing in A. terreus also caused decreased expression of gene sod1.

2) ROS build up began before schedule, reaching higher levels than in the parental.
Hence,

3) Gene *brlA* (regulator of conidiation) expressed earlier, and reached higher spore densities.

4) Also, gene *lovE* (lovastatin genes specific transcription factor) expression and lovastatin production started earlier and reached higher production levels than the parental in both culture systems. yap1-silenced mutants displayed lovastatin production increases of: 60% in SmF and 70% in SSF.
Background

The limited supply and the negative environmental effects of the use of petroleum-derived fuels and chemicals have stimulated efforts for the development of more environmentally-friendly processes.

Objectives

The bacterial fermentation of carbohydrates is a promising way for the production of green chemicals and biofuels. My project aims at improving the efficiency of the anaerobic conversion of sugars by *Clostridium* species into C3 alcohols, namely isopropanol.

Methods

Genome sequencing and fermentation performance studies were carried out on multiple *Clostridia* strains including the natural Isopropanol-Butanol-Ethanol producing strain *Clostridium beijerinckii* DSM 6423 (NRRL B593). This strain was then used to generate, through N-methyl-N'-nitro-N-nitrosoguanidine (NTG) mutagenesis, 36 strains with altered fermentation profiles. The selective growth was performed in the presence of 3 suicide substrates or a high concentration of isopropanol (> 35g/L). Further improvement of this first set of mutated strains was achieved through genome shuffling which uses recursive protoplast fusion and offers the advantage of entire genome recombination (Gao, Zhao, Zhang, He, & Jin, 2012). Screening of the final strains was based on enhanced isopropanol tolerance, the best of them being able to tolerate concentrations of up to 50g/L.

Conclusions
Some improvement in isopropanol production compared to the wild type strain was attained with strains showing higher solvent yields and/or better selectivity. The best *Clostridium* strains obtained produce less butanol and more isopropanol allowing a more efficient bioprocess.
EVALUATION OF OENOCCUS OENI EXOPOLYSACCHARIDES AS ENDOGENEOUS PROTECTIVE AGENTS FOR MALOLACTIC STARTER PRODUCTION

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Background

The malolactic fermentation (FML) is an essential winemaking step driven by lactic acid bacteria. It takes place after the alcoholic fermentation and mainly consists in the conversion of L-malic to L-lactic acid, with subsequent improvement of the wine microbial stability, aroma and flavor. Oenococcus oeni is the most suitable species for driving FML in wine. O. oeni may come from the indigenous microflora found on grapes or in the cellar, or it may be massively added to wine as a malolactic starter. O. oeni malolactic starters are currently selected for their resistance to wine and their ability to produce polysaccharides is so far not considered. However, this property could be crucial for the protection of the bacteria during starter production and use.

Objectives

Our main objective was to examine whether the stimulation of the production of exopolysaccharides by the bacteria in situ before freeze drying could protect the bacteria and improve the survival rates towards production steps but also during inoculation in wine.

Methods

The genome sequencing of 50 O. oeni strains permitted us to realize the inventory of the exopolysaccharides genes (RAST, Kaas). The exopolysaccharides analysis included the quantification (colorimetric and analytical methods) and phenotypic observation (optical and electron microscopy) of the produced exopolysaccharides. The production of O. oeni strains in a lyophilized form and their later inoculation in wine (freeze-drying, cells viability).

Conclusions
We observed that the tested strains showed specific survival behaviors, which were perfectly correlated with their EPS biosynthetic ability. The protecting role of the bacterial exopolysaccharides is now to be proposed at industrial level.
Background

Background: The second generation ethanol using agro-industrial waste is a promising alternative biofuel but the challenge is a technological approach to deconstruction the recalcitrant lignocellulose.

Objectives

Objective: The present work aimed to study the effect of pre-treatment of sugar cane bagasse with microwave radiation in the presence of glycerol on chemical composition and the efficiency of subsequent enzymatic hydrolysis.

Methods

Bagasse immersed in glycerol 10% were subjected to 2 min of microwave irradiation and the solid fractions resulting were used in the analysis of fibers, TGA, DTG, DSC, FTIR, X-ray and transmission electron microscopy and the liquid fractions were used to determine sugar and phenol contents. Samples of bagasse treated and untreated (control) were submitted to enzymatic hydrolysis for 24 to 72 h at 55 °C with enzymatic solutions obtained by the cultivation of M. thermophila M.7.7.

Conclusions

Infrared spectra and thermal analysis showed that pre-treatment acted on the lignin and hemicellulose of bagasse. The hydrolysis with the enzyme produced by M. thermophila supplemented with β-glucosidase, afforded 74.0 mg/g and after 72 h the highest yield of reducing sugars was 240.9 mg/g, with β-glucosidase or not. The pre-treatment with microwave and glycerol improve in 40% the enzymatic hydrolysies of the cellulose.
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FUNGAL AND BACTERIAL CELLULOXYTIC ENZYMES TRCEL7B AND CFXYN11A ACTING IN SYNERGY DURING THE HYDROLYSIS OF SUGAR CANE BAGASSE
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Background

Enzymatic bioconversion of lignocellulose to sugar monomers is performed by cellulases and hemicellulases acting synergically. Synergy among cellulases has been documented [1]; synergy between cellulases and accessory enzymes has been studied in cellulosome systems from Clostridium cellulovorans [2]. Trichoderma reesei and Cellulomonas flavigena are two no related cellulolytic organisms which produce enzymes with reported cellulase and xylanase activities, respectively[3,4]


Objectives

To evaluate synergy between two no complex enzymes, fungal cellulase (TrCel7B) and bacterial xylanase (CFXyn11A), in the sugar release of enzymatic hydrolysis of sugar cane bagasse.

Methods

Full length CfXyn11A xylanase from C. flavigena was expressed in E.coli [4]. Core Tr Cel7B was expressed in T.reesei [3]. Enzymatic bagasse hydrolysis was performed in equimolar ratios of cellulose(C) and xylanase(X): 100C,75C:25X,50C:50X,25C:75X, 100X. Release of sugar monomers was followed by HPLC after acid hydrolysis. Degree of synergy(D.S.) was quantified as the ratio of the specific activities of the enzymes in the hydrolysis mixtures divided by the sum of the individual specific activities at the concentration used.

Conclusions
Synergy between \textit{Cf}Xyn11A and \textit{Tr}Cel7b during bagasse hydrolysis was observed in glucose and xylose but not in arabinose release. The highest xylose and arabinose release observed at 12 h suggests a debranching role of the xylanase prior to cellulase hydrolysis occurring mainly after 24 h. However, a high D.S. observed in the 50C:50X at 12h, enforce the best performance of enzymes when they act together.
DEVELOPMENT OF YARROWIA LIPOLYTICA AS AN OLEAGINOUS CELL FACTORY PLATFORM FOR PRODUCING NUTRACEUTICAL AND COSMETIC MATERIALS

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Background

Yarrowia lipolytica which has been extensively used as a model oleaginous yeast for producing heterologous proteins has capability to accumulate lipids (lipidbody, peroxisome etc) up to 50% of the DCW (dry cell weight). In addition, Y. lipolytica can be easily manipulated in gene deletion and insertion because it possesses a haploid genome and its genome sequence was completed and it is also known as GRAS (generally recognized as safe). Therefore, Y. lipolytica can be a high potential host strain in producing oleaginous (hydrophobic) and high value-added compounds used in nutraceutical and cosmetic materials which are hydrophobic and produced from plants by conventional extraction processes.

Objectives

In this study, we aim to construct basic and potential platform strain suitable for production of nutraceutical and cosmetic materials by using Y. lipolytica.

Methods

We used Y. lipolytica Po1g and constructed an ura3 auxotroph strain, Y. lipolytica (PolgDura3), in order to use URA3 blaster system which is a useful tool to carry out repeat genetic deletion and insertion on the genome. In addition, we constructed a Y. lipolytica (PolgDku70Dura3) in which Ku70 involved in non-homologous recombination was deleted to increase recombination efficiency in this strain.

Conclusions

We confirm that Y. lipolytica (PolgDku70Dura3) showed higher homologous recombination efficiency (>5 folds) than Y. lipolytica (PolgDura3). Next, we are introducing foreign genes involved in biosynthesis of some specific nutraceutical and cosmetic materials into Y. lipolytica (PolgDku70Dura3). This strain and genetic tool will be very useful to construct various customized strains for producing nutraceutical and cosmetic materials.
MODULATION OF OXIDATIVE FOLDING TO IMPROVE RECOMBINANT PROTEIN PRODUCTION IN THE YEAST SACCHAROMYCES CEREVISIAE

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Background
The yeast Saccharomyces cerevisiae is widely used in biotechnology for the production of both bulk chemicals, like ethanol and organic acids, and higher-value compounds such as recombinant proteins, like pro-insulin. Recombinant protein production in S. cerevisiae induces different stress responses, including oxidative stress response. This is especially the case when the overall protein folding rate and the oxidative folding rate, i.e. the formation of di-sulfide bonds, are not properly balanced (1).

Objectives
In the present study we aim to develop metabolic engineering strategies to increase recombinant protein production by specifically adjusting oxidative folding.

Methods
Two industrially relevant proteins were selected based on different overall folding rates: pro-insulin and alpha-amylase. The oxidative folding of these proteins was modulated by A) altering the expression levels of several key players in oxidative folding, e.g. the thiol oxidases Ero1 and Erv2, and B) changing the number of possible di-sulfide bonds, i.e. the number of cysteine residues present.

Conclusions
The effects of these modulations on reactive oxygen species (ROS) levels, unfolded protein response, oxidative state, protein titers and overall physiology are monitored under strictly controlled growth conditions.

Reference
PRODUCTION OF BIO-OIL BY OLEAGINOUS YEAST

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Background

The Dutch government is stimulating the transition to a more biobased economy. Microbial oil can be produced from biomass in a green and sustainable way without competing to food crops by using oleaginous microorganisms such as yeast, fungi, bacteria and microalgae. At the HAN BioCentre a unique robust oil producing yeast strain has been selected called HBC025. The strain grows on a broad range of C5 and C6 sugars. The oil is accumulated by HBC025 in the form of triacylglycerols (TAGs) containing predominantly oleic (18:1), linoleic (18:2) and palmitic (16:0) acid. The produced microbial oil can be used as biodiesel but also in other applications such as paint, roads, floors and cosmetics.

Objectives

Objective is the development of an economically feasible fermentation and extraction process for second generation oil production by HBC025.

Methods

A fed-batch fermentation process was developed on glucose as model substrate. Hydrolyzed paper pulp was obtained from Parenco (Renkum, The Netherlands). Extraction methods tested were homogenization and enzyme assisted auto-lysis.

Conclusions

Maximum productivity on glucose as substrate was 0.5 g bio-oil/l/h, which is comparable to the best lipid producing strains in literature (Ageitos et al. 2011). HBC025 was also able to grow on hydrolyzed paper pulp as substrate with the same growth rate. Calculations showed that the process can be economically feasible only in case components from the yeast biomass are also marketed besides the yeast oil.

References
REVERSIBLE PSEUDOHYPHAE FORMATION IN PICHIA PASTORIS DURING HETEROLOGOUS PROTEIN PRODUCTION.

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Background

The methylotrophic Pichia pastoris yeast expression system is widely used for production of heterologous proteins. However, till now morphological changes have not been reported in P. pastoris. This study reports for the first time, a comprehensive characterization of Granulocyte colony stimulating factor (GCSF) expression in filamentous P. pastoris.

Objectives

To investigate the morphological changes in P. pastoris during protein production phase.

Methods

Mut¹ transformants, containing the GCSF-cDNA fused with modified kex2 cleavage site (in the α-factor secretion signal) under the control of the alcohol oxidase 1 promoter, were used in the study. The transformants were cultivated in BMGY (buffered glycerol complex medium) followed by BMMY (buffered methanol complex medium). Extracellular protein was quantitated by gel densitometry from 20-fold concentrated culture supernatant. A combination of methanol level and feeding strategy was developed to increase productivity of GCSF at fermenter level.

Observation:
Reversible Pseudohyphae was observed, resulting in increased surface to volume ratio with an increase in extracellular protein relative to the normal phenotype of spherical cells at same cell dry weight.
Fig1.: Morphology of recombinants (A) Mut⁺ (methanol utilization plus) and (B) Mut⁻ (methanol utilization slow) strains of *P. pastoris*, SMD1168 cultivated in BMMY media + 0.4% histidine.

Conclusions

*P. pastoris* initiates a striking developmental transition to filamentous form, more so in the Mut⁻ phenotype, that results in more surface to volume ratio, and affects productivity of the culture. To the best of our knowledge, data presented here opens the possibility of a filamentous form of *P. pastoris* for production of heterologous proteins.
HIGH-LEVEL PRODUCTION OF MONO-COMPONENT AND ENZYME MIXTURES IN MYCELIOPHTHORA THERMOPHILA
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Background

The fungus Myceliophthora thermophila C1 was developed into an efficient and versatile platform for high-level production of industrially relevant enzymes.

Objectives

Through strain development strategies, two strain lineages were obtained that are currently being developed and exploited as production hosts. One lineage (HC) is able to produce and secrete high amounts of enzyme mixtures that contain large amounts of (hemi-) cellulases. The other lineage (LC) is impaired in its cellulase producing capability, resulting in low background-protein production. The LC strain has been further developed for high-level enzyme production. Total protein levels up to 80 g/L has been reached of which ~ 80% correlates to the introduced enzyme.

Methods

By transforming the LC strain with selected C1 genes, a wide collection of strains was obtained, each of which produced mainly one enzyme. This has led to an enzyme library of over 100 functional enzymes of which many have been purified and characterized. The LC strain can also be designed and constructed in such a way that it produces only those enzymes that are functional under dedicated application conditions.

Conclusions

In conclusion, M. thermophila was developed into a high-level protein-production platform. The HC strain is successfully applied to produce enzymes for the production of biofuels and biobased-chemicals. The LC strain is being used to produce single enzymes and defined combinations of enzymes. The obtained C1-enzyme library is a rich source for academic and industrial research. The properties of M. thermophila C1
make this fungus a highly suitable alternative for traditional fungal protein production hosts.
NEW φBT1 SITE-SPECIFIC INTEGRATIVE VECTORS WITH NEUTRAL PHENOTYPE IN STREPTOMYCES

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Background
Integrative plasmids are one of the best options to introduce genes in low-copy-number and a stable form into Streptomyces, but they generate mutations in their integration sites. Phage (PHI)C31, the most common integrative site used in Streptomyces, integrates at different positions (attB and pseudo-attB sites) in the Streptomyces genome generating different mutations and phenotypes. The less common integration site (PHI)BT1, integrates at a unique attB site localized in gene SCO4848 (S. coelicolor genome) or their homologues in other streptomycetes.

Objectives
Constructing an integrative plasmid with neutral phenotype.

Methods
Four plasmids were created modifying the φBT1 integrative vector pMS82: pNG1, in which the SCO4849 was introduced under the control of the promoter region of SCO4849 into pMS82 to restore the phenotype generated by the integration of the plasmid; pNG2, in which the promoter P_{ermE} was introduced into pNG1 to facilitate heterologous gene expression; pNG3 in which the bla gene for ampicillin resistance was included into pNG1 in order to facilitate selection in E. coli; and pNG4, in which the bla gene was introduced into pNG2.

Conclusions
The plasmids pNG1, pNG2, pNG3, and pNG4, are the only integrative vectors designed to produce a neutral phenotype when they are integrated in streptomycetes.
THE INFLUENCE OF LACTIC ACID BACTERIA ADDITIVE ON THE LEVEL OF RESIDUAL WATER-SOLUBLE CARBOHYDRATES IN CORN SILAGE

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Background
Ensilaging is a high moisture crop conservation method based on the fermentation of water soluble carbohydrates (WSC) guided by epiphytic microflora, specifically by lactic acid bacteria (LAB). LAB additives (inoculants) in ensilaging are used for promotion of optimal fermentation and rapid pH drop thus improving dominant nutrients preservation. However, the impact of inoculation on the level of residual WSC, which in high levels may result in silages aerobic deterioration, is not clear.

Objectives
During the present study, the application of LAB inoculant on residual WSC was investigated.

Methods
The three different yellow corn hybrid (Bc 418b, Bc 678 and Bc exp 6) produced in the same production conditions (31,01 – 38,52% DM) were ensiled in five replications in laboratory scale silos with and without Sil-All®4×4 LAB inoculant in a concentration 1x10⁷ CFU/g of fresh material. The silages were sampled on the 21st and the 60th day. The contents of lactic acid, WSC and pH were monitored.

Conclusions
The analyses showed that the inoculated silages had a significant increase (P<0,05) of lactic acid concentrations (g/kg DM basic) both on the 21st (31,45) and the 60th day (33,28) in contrast to silages without inoculant (21st 22,49; 60th 26,38). The same pattern was observed with pH. At the end of ensiling the difference in WSC content between silage with (8,63 g/kg DM) and without inoculant (7,39 g/kg DM) was not statistically significant (P>0,05). Inoculation of silages with Sil-All®4×4 stimulates lactic acid production and rapid pH drop whereas has no influence on the level of residual water-soluble carbohydrates.
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FUNCTIONAL ANALYSIS OF FILIPIN TAILORING GENES FROM STREPTOMYCES FILIPINENSIS
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Background

Streptomyces filipinensis is the industrial producer of filipin, a 28-membered non-glycosylated pentaene macrolide widely used for the detection and the quantitation of cholesterol in biological membranes.

Objectives

Identification of tailoring genes belonging to the biosynthetic cluster for the biosynthesis of filipin, and generation of engineered derivatives for pathway elucidation.

Methods

A region of 13778 base pairs of DNA from the S. filipinensis genome was isolated, sequenced, and characterized by gene replacement and complementation techniques.

Conclusions

This set of genes shows synteny with the homologous pte genes from the filipin cluster of S. avermitilis, and includes two cytochrome P450 monooxygenase encoding genes, filC and filD, which are proposed to catalyze specific hydroxylations of the macrolide ring at C26 and C1’ respectively. Gene deletion and complementation experiments provided evidence for their role during filipin III biosynthesis. Filipin III derivatives were accumulated by the recombinant mutants at high yield. These have been characterized by mass spectrometry following high-performance liquid chromatography purification thus revealing the post-polyketide steps during polyene biosynthesis. Two alternative routes lead to the formation of filipin III from the initial product of polyketide synthase chain assembly and cyclization filipin I, one trough filipin II, and the other one trough 1’-hydroxyfilipin I, both compounds being biologically active.
SCREENING OF LACTIC ACID BACTERIA FOR PRODUCTION OF BIOSURFACTANTS
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Background
Lactic acid bacteria (LAB) have been extensively used for thousands of years in food fermentation and nowadays in the industrial production of lactic acid and other metabolites.

Objectives
In this study, the biosynthetic potential of lactic acid bacteria was exploited for biosurfactant (BS) production.

Methods
66 Lactobacilli strains were screened for BS production when growing in different MRS-based media at static conditions. To determine BS production (extracellular and cell-bound), surface tension (ST) of supernatant and phosphate buffered saline (PBS) extract of the strains were measured by tensiometer. To determine critical micellar concentrations (CMC), PBS extracts of the strains remarkably reduced ST were diluted.

Pyrene solubilization assays were further done as a complementary assay to prove the presence of BSs. 1 mg of pyrene was distributed into glass test tubes and was subjected to supernatant and PBS extract of the strains in shaking conditions for 24h. The values were then compared with those of pyrene solubilization at the presence of rhamnolipid in different CMCs.

Conclusions
The results showed there is a significant potential for BS production among the strains screened and 9 out of 66 Lactobacilli strains reduced ST of PBS extract between 19 and 22 units. The values are close to the data reported by Moldes et al. 2007 for Lactobacillus pentosus CECET-4023 (20 units) and Gudina et al. 2011 for Lactobacillus paracasei spp. paracasei A20 (22 units).
Pyrene solubilization assay can be taken into account as a promising method to identify BS-producing strains.

Concentrations and characterization of BSs produced are under investigation.
CHARACTERIZATION OF SCO4439, A D-ALANYL-D-ALANINE CARBOXYPEPTIDASE INVOLVED IN SPORE CELL WALL MATURATION, SPORE RESISTANCE, GERMINATION AND MYCELIAL RESISTANCE TO VANCOMYCIN AND TEICOPLANIN

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Background

Streptomycetes are model microbes with complex developmental cycles. The last advances in system biology methodologies allowed the identification of large datasheets of proteins and genes differentially expressed during development whose biological function remains, in many cases, unexplored.

Objectives

Characterization of the biological function of SCO4439, a developmentally associated D-alanyl-D-alanine carboxypeptidase (DD-CPase) overexpressed during the aerial mycelium and sporulation stages.

Methods

Strain: Streptomyces coelicolor M145

Conclusions

1- SCO4439 is a multidomain protein harbouring a DD-CPase at its carboxyl end, and a putative cytosolic transcriptional regulator domain at the N-terminal end. Both domains are separated by a putative transmembrane region.
2- The DD-CPase activity was demonstrated ‘in vitro” to have a preference for DAla-DAla peptides, and to be inhibited by penicillin G.
3- SCO4439 was demonstrated ‘in vivo” to be involved in the resistance of the spores to heating and acid, in the spore swelling during germination, and in the resistance of the mycelium to glycopeptide antibiotics (vancomycin and teicoplanin).
4- The DD-CPase domain, together with the putative hydrophobic transmembrane region are highly conserved in Streptomyces, and the presence of both domains is essential to restore the phenotypes observed in the SCO4439::Tn5062 mutant. The conservation of the putative transcriptional regulator domain is much lower, and it is not necessary to restore the phenotypes detected in the SCO4439::Tn5062 mutant.
5- SCO4439 controls PG cross-linking releasing DAla from peptidoglycan lateral chains and reducing the substrate for the transpeptidation.
The level of peptidoglycan cross-linking is regulating spore resistance and germination.
AN EFFICIENT FED-BATCH FERMENTATION PROCESS: THE PRODUCTION OF STREPTOKINASE FROM E. COLI
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Background
Streptokinase is a therapeutic protein drug of bacterial origin. Its role in thrombolytic therapy has been well established. It is either produced from Streptococcus, its natural producer or from E.coli by recombinant means. The production from the former, however, is problematic due to low yield and simultaneous production of toxic endolytic chemicals. At IMTECH, the gene encoding for streptokinase from Streptococcus equisimilis was cloned and expressed in E. coli BL21 (DE3). Initially the expression in the latter was very low, hence the coding sequence was appropriately modified using molecular biology approaches and as a result of such optimizations, a high expression level of upto 80 mg l⁻¹ was obtained in shake flask cultivation.

Objectives
The main objective of this study was to develop a bioprocess that could be used for mass scale production and purification of this high value therapeutic protein.

Methods
To achieve the above objective, a fed-batch fermentation process was developed and optimized. Cultivation parameters like dissolved oxygen concentration (DOC), pH, media constituents and feed rate were carefully optimized and product yield was monitored.

Conclusions
The process optimization resulted in an intracellular expression level of 1120 mg l⁻¹ in high cell density fermentation. This represented a 14-fold increase in production levels from flask to bio-reactor. Also the specific expression level i.e. gram protein produced per gram wet cell mass remained unchanged. This yield is the highest reported so far for recombinant Streptokinase.
Background
This research investigates the mechanism of lignocellulose degradation by brown rot fungi and potential industrial applications of this system. The brown rot non-enzymatic degradative mechanism is largely unexplored for use in biomimetic industrial/biorefinery applications and understanding the degradative mechanisms of brown rot fungi will lead to better methods of addressing the recalcitrance problems associated with lignin and crystalline cellulose in the pretreatment of biomass in biorefinery applications. A chelator mediated Fenton (CMF) system has evolved to substitute for components of the cellulolytic enzyme machinery in the brown rot fungi, generating an alternative efficient mechanism for depolymerization of biomass. The CMF system is unique among biological systems in being the only reported substrate deconstruction system based on oxygen radical chemistry that permits non-enzymatic deconstruction at a considerable distance (several microns) from the organism.

Objectives
Specific objectives of this research include: 1) Examining the morphology of the fungal sheath in a brown rotted fungal-wood complex to determine how the sheath may aid in the movement of CMF components and enzymes; and 2) Analyzing the structure of lignocellulose in early and late-stages of brown rot fungal degradation to assess how the CMF system modifies the wood cell wall non-enzymatically.

Methods
This research utilized several different techniques including the ORNL Bio-SANS (small angle neutron scattering) beam line to obtain the mechanistic information necessary to better characterize processes involved in non-enzymatic biodegradation of lignocellulose by brown rot fungi.

Conclusions
This research supports a CMF model system being employed by brown rot fungal organisms in lignocellulose deconstruction.
CHARACTERIZATION OF 3-KETOSTEROID-9Α-HYDROXYLASES IN RHODOCOCCUS RUBER STRAIN CHOL-4

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Background
Bacterial catabolism of steroid compounds provides a broad range of intermediaries needed for the synthesis of pharmaceutical steroid drugs such as 4-androsten-3,17-dione (AD) and 1,4-androstadiene-3,17-dione (ADD). Rhodococcus ruber strain Chol-4, isolated from a sewage sludge sample, is able to grow in minimal medium supplemented with steroids, showing a large catabolic capacity. The 3-Ketosteroid 9α-Hydroxylase (KSH) is a key enzyme in the general scheme of bacterial steroid catabolism; it initiates the opening of the steroid ring by the 9α-hydroxylation of the C9 carbon of 4-ene-3-oxosteroids (e.g. AD) or 1,4-diene-3-oxosteroids (e.g. ADD), transforming them into 9α-hydroxy-4-androsten-3,17-dione (e.g. 9OHAD) or 9α-hydroxy-1,4-androstadiene-3,17-dione (e.g. 9OHADD), respectively.

Objectives
Study and characterization of 3-ketosteroid-9α-hydroxylases in Rhodococcus ruber strain chol-4

Methods
The ORF finder and the pDRAW32 programs were used to detect the ORFs. Determination of KSHs induced or constitutive transcription, along with the co-transcription of each cluster, was made by RT-PCR. Mutagenesis of both single, double and triple kshA and kshB were made by unmarked gene deletion. Growth studies were analyzed on different substrates.

Conclusions
This work presents a set of data that clarifies KHAs specific roles in strain chol-4. We have characterized three different kshA and one kshB homologous ORFs in strain Chol-4. All of them present constitutive transcription and co-transcription. The mutants have shown differences in growing depending on the substrate.
Background

Transposase is an enzyme that binds to the ends of a transposon and catalyzes the movement of the transposon to another part of the genome by a cut and paste mechanism or a replicative transposition mechanism. Rhizobium is a genus of Gram-negative soil bacteria, many of which fix nitrogen. Nitrogen fixation is an important part of the nitrogen cycle. Plants cannot use atmospheric nitrogen (N₂) they must use nitrogen compounds such as nitrates.

Objectives

The objective of this research was to determine whether at random transposon mutagenesis could be applied in genetic studies of Rhizobium japonicum. One major reason for this is that a range of defined mutations is not available.

Methods

In present study R. japonicum at frequencies sufficient to allow the isolation of large numbers of insertion mutants. The selection of Tn5 mutants was facilitated by the expression, in all the R. japonicum strains we have tested, of the Tn5 encoded kanamycin resistance. A number of auxotrophic and symbiotically defective, single, random transposon (Tn5) mutants were obtained in three slow-growing strains of different DNA homology and serogroups and the single fast growing strain of R. japonium.

Conclusions

In conclusion, the diversity of auxotrophs detected and the isolation of symbiotic mutants suggests that transposon Tn5 can be used as a generalized mutagen to isolate a variety of mutants with defects in symbiotic nitrogen fixation. The analysis of such mutants should prove to be useful in elucidating the biochemical, genetic, and regulatory events involved in the R. japonicum which effectively nodulates certain Indian soybean cultivars.
ACINETOBACTER STRAIN XB174 DEGRADES QUORUM SENSING SIGNAL OF RALSTONIA SOLANACEARUM, PREVENTS WILT AND PROMOTES GROWTH IN EGGPLANT
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Background

*Ralstonia solanacearum* is one of the most destructive plant pathogens and causes bacterial wilt in about 450 plant species worldwide. During infection, the pathogen primarily colonizes plant vascular system in high density leading to wilt and eventually plant death. Expression of virulence factors by the pathogen is regulated by a quorum sensing system activated 3-hydroxy palmitic acid methyl ester (3OH-PAME).

Objectives

The objective of this study was to determine 3OH-PAME degradation by bacterial strains isolated from eggplant xylem and to determine their wilt prevention and growth promotion ability in eggplant.

Methods

Bacterial strains (n = 6) were earlier isolated from xylem sap of eggplant. Using a bioassay involving indicator *R. solanacearum* strain AW1-3 and High Performance Liquid Chromatography Mass Spectrometry, 3OH-PAME degradation by bacterial isolates was estimated. Strain XB174 degraded 3OH-PAME and was evaluated for wilt prevention and growth promotion in eggplant under greenhouse conditions. XB174 was identified by 16s rRNA gene sequencing and its activity against Tween-20, Tween-80 and tributyrin was determined.

Conclusions

Among the strains screened using bioassay, only XB174 degraded 3OH-PAME to 3OH-palmitic acid. Specific activity of crude extracellular enzyme of XB174 towards 3OH-PAME was 17.78 U mg⁻¹. XB174 prevented wilt (55.0 ± 7% over control) and improved growth (19.2 ± 10% more growth promotion efficacy over control) in eggplant. XB174 is identified as *Acinetobacter* sp., degraded Tween-20, Tween-80 and tributyrin.
The study reveals the quorum quenching activity of *Acinetobacter* strain XB174 against *R. solanacearum*. After additional screening, extracellular enzyme from XB174 may have applications in agriculture and industries.
FUNCTIONAL CHARACTERIZATION OF A LUXR TYPE REGULATOR, ABAR IN ACINETOBACTER BAUMANNII: INVOLVEMENT OF ABAR IN PATHOGENESIS

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Background

Quorum sensing is a cell to cell communication system that coordinates gene expression in many bacterial species. It has become an increasingly interesting target for the development of alternative drugs for treating multi-drug resistant Acinetobacter baumannii infection. However, the role of quorum sensing system in A. baumannii has not been characterized.

Objectives

The aim of this study is to characterize the functions of AbaR, a putative LuxR type receptor, using an isogenic mutant, and to shed some light on to the development of effective alternative drugs targeting AbaR.

Methods

The disruption of abaR in A. baumannii lead to significant decrease in biofilm formation. In addition, the pellicle production and swarming motility of the abaR mutant was substantially decreased compared to that of wild-type. The abaR mutant complemented with the intact abaR gene reverted back to that of wild-type capable of producing biofilm and displaying swarming motility. The survival rate of mice infected with the abaR mutant strain was significantly high than that of those infected with the A. baumannii wild-type. Also, whole transcriptome analysis was performed in order to compare the gene expression in wild-type and abaR mutant strains. Furthermore, chemical screening was carried out for the possible quorum quenching compounds targeting AbaR.

Conclusions

The data from the current study reveals that AbaR plays an important role in A. baumannii pathogenesis and this regulator protein could be a putative target of quorum quenching compounds. Supported by KHIDI (grant number: HI14C0257)
How does Hamametannin Increase Staphylococcus Aureus Biofilm Susceptibility Toward Glycopeptides?

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Background
Staphylococcus aureus biofilm-associated infections are difficult to treat and novel targets are needed to combat these infections. The QS modulator hamametannin increases susceptibility of S. aureus biofilms towards vancomycin in vitro as well as in vivo. However, it is unclear how HAM affects susceptibility.

Objectives
Elucidate the mechanism of action of HAM at the molecular level.

Methods
Two parallel strategies were followed. First, we evaluated the effect of HAM on biofilms of S. aureus strains with mutations in the QS systems or in genes involved in biofilm formation and virulence. Secondly, using illumina sequencing we identified genes that were differentially expressed in untreated biofilms and biofilms treated with VAN alone or in combination with HAM. Results obtained with both strategies were further investigated using the appropriate tools.

Conclusions
No loss in HAM activity was observed for most of the mutants. In contrast, HAM did not affect biofilm susceptibility of S. aureus strains with mutations in the QS system. Using sequencing, we identified a large number of genes that were differentially regulated after treatment (e.g. genes involved in biosynthesis of lysine, D-alanine, glutamine consuming pathways and virulence). This indicates that HAM reduces the upregulation of peptidoglycan biosynthesis normally observed after treatment with VAN. This possibly leads to the increased susceptibility of S. aureus biofilm cells towards VAN. Our results further indicate that combination therapy could positively affect morbidity since the upregulation of virulence factors observed for VAN treatment are not observed when VAN is combined with HAM.
Background
MELiSSA (Micro-Ecological Life Support System Alternative) has been conceived as a 5 compartments microorganisms and higher plants recycling system for long haul space flights. *Rhodospirillum rubrum* S1H colonizes compartment II and grows under light anaerobic conditions (LAN) using acetate as carbon source (MELiSSA conditions). Previous work reported that continuous culture of the bacterium in a photobioreactor lead to thick biofilm formation, leading to bioreactor arrest.

Objectives
The aim of this research is to investigate the relation of the quorum sensing (QS) system and biofilm formation of *R. rubrum* S1H (wild type, WT) under MELiSSA relevant culture conditions.

Methods
In this context we have constructed a mutant strain named M68 that does not produce acyl homoserine lactones (AHLs) signaling molecules. The transcriptomic and proteomic profiles and phenotype of WT and M68 under MELiSSA conditions were compared. In addition, the biofilm development of WT and M68 in a flow cell system under light microaerobic conditions was studied.

Conclusions
*R. rubrum* has a cell-to-cell communication system based on AHLs which regulates 8% (326 genes) of the genome of M68. In *R. rubrum* QS regulates pigmentation, photosynthesis, energy generation, carbon metabolism, motility and biofilm formation. To our knowledge this is the first report where QS is linked to biofilm formation in *R. rubrum* S1H under light microaerobic conditions. Further flow cell experiments under MELiSSA conditions will help us to study biofilm formation and to test substances with known anti-biofouling properties.
Background

In addition to causing various infections, Acinetobacter species are described as nosocomical pathogens. These species are increasingly developing resistance against antibacterial agents due to their capability to adapt to any environment. Acinetobacter species develops these capabilities against their environment by quorum sensing (QS) mechanism which is a bacterial communication or ambient sensing mechanism. Quorum sensing systems needed for iron uptake comprising genes required for siderophore biosynthesis.

Objectives

In this work, effect of signal molecules on siderofor biosynthesis and its relation with QSS in MDR A.baumannii were studied.

Methods

Twenty strains of MDR A.baumannii from different clinical specimens were identified by standard methods. QSS molecules were identified by cross-validation test and siderophore biosynthesis was evaluated on CASagar and thin layer chromatography(TLC). AHL profiles were determined by using Agrobacterium tumefaciens biosensor strains (A136 or NTL1). QSS molecules combined with the biosensor strains were separated by TLC from each other.

Conclusions

According to results of the study, it was found out that all of the MDR A. baumannii strains produced the following miscellaneous QSS molecules in varying amounts: C8-HSL, C10-HSL and C12-HSL and these strains produced hydroxamate- and catechol-type siderophores. Iron uptaking in the environments is thought to be a direct relationship with quorum sensing system. The variations in the amounts and types of signal molecules were observed to affect virulence properties. QSS molecules can be used as target molecules in medical therapy. The study provides new information about the virulence factors for the better understanding of the infection properties.
of *A. baumannii* in the host.
BIO-CONTROL APPLICATION OF BACILLUS QUORUM SENSING ON AFLATOXIGENIC FUNGI THROUGH THEIR INTER-KINGDOM CROSS-TALK

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Background
Quorum sensing molecules (QSM) are involved in the regulation of complicated processes helping bacterial population benefit from their cell-density. Prokaryotes and eukaryotes' co-evolution raises the prospect of the existence of inter-kingdom signalling pathways. The role of hormone-like molecules in cell communication approves QS role in this cross-talk. Bacterial antagonistic activity against fungi is considered as an inter-kingdom talk. Bacillus and Pseudomonas have the ability to inhibit Aspergillus growth and aflatoxins production. Screening of antagonistic bacteria against Aspergillus flavus in vitro, Bacillus subtilis was identified with high antifungal activity.

Objectives
Bacillus licheniformis has industrial application due to its production of antimicrobial compounds and is related to B. subtilis genetically, whose control of competence sporulation is regulated by a QS mechanism. QS in B. subtilis is regulated by ComX pheromone. As QS genes have been identified in B. licheniformis NCIMB-8874, our study has focused on QSM in this strain.

Methods
To investigate cell-cell communication, the comQX locus was sub-cloned into a shuttle vector which was expressed in E. coli and pheromone was isolated by reverse phase chromatography.

Conclusions
Pheromone as a QSM is potential signal for communicating between kingdoms and could be applied for bio-control purposes. Identification of new antifungal peptides against A. flavus could lead to the development of biotechnological strategies which facilitate control of aflatoxin contamination.
THE FUNCTIONAL UNITS OF QUORUM SENSING (QS) IN AMMONIA-OXIDIZING BACTERIA

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Background

Quorum sensing (QS) is an important form of cell-to-cell communication that regulates gene expression in response to fluctuations in cell density. One of the best-studied examples of quorum sensing is N-acyl homoserine lactones (AHLs)-mediated cell-cell communication in gram-negative bacteria. While it has been able to describe the QS system in some individual microorganisms for some time now, the functional QS system of nitrifying bacteria has not been completely proved to date.

Objectives

Testing the existence and characterization of quorum-sensing signal produced by nitrifying bacteria (like *Nitrosomonas europaea*, *Nitrosospira multiformis*), and the functional QS signal synthase, the cognate receptor responding to the synthase-generated signal in these microorganisms.

Methods

The AHLs product was obtained by extracting bacteria culture supernatants with acidified ethyl acetate, and then detected by AHLs biosensor and liquid chromatography-mass spectrometry (LC-MS). To identify gene(s) encoding AHL synthases and transcriptional activators, the genome of related nitrifying bacteria was searched for genes with similarity to all QS-related genes. AHL synthases gene was expressed in the heterologous host *E. coli* BL21. Putative AHLs in the recombinant extract were confirmed by comparing the retention times and mass spectra from LC-MS with those of standard AHLs. As a test of the hypothesis that these AHLs were cognate signals of probable receptor, we also examined the solubility of the R protein.

Conclusions
We have shown a LuxI/R type QS signal synthase and regulator in an ammonia-oxidizing strain, and the results provide an opportunity to complete the QS regulatory networks.
Background
Burkholderia glumae, the causal agent of rice panicle blight, contains a single LuxI-R type quorum sensing (QS) system. TofI synthesizes N-octanoyl homoserine lactone (C8-HSL), which is recognized by TofR to regulate various private and public goods. The complex of TofR and C8-HSL activates expression of genes for toxoflavin and flagellar biosynthesis and an IclR type transcriptional regulator gene, qsmR. Among QS-dependent public goods, oxalate is produced to protect cells from ammonia-mediated alkaline toxicity in LB during stationary phase. RNAseq analysis indicated that expression of isocitrate lyase gene (aceA) might depend on QsmR.

Objectives
We aimed to confirm that glyoxylate cycle is controlled by QS and to determine roles of QS-dependent glyoxylate cycle in B. glumae.

Methods
To ascertain control of glyoxylate cycle by QS, we measured expression level of aceA and malate synthase (glcB) genes from the chromosomal Tn3-gusA fusions in each gene in the wild type and the QS mutants. Electrophoretic mobility shift assays were performed using the promoter regions of aceA and glcB and purified QsmR. AceA enzyme activity was determined by measuring the formation of glyoxylate-phenylhydrazone in the presence of phenylhydrazine and isocitrate at 324 nm. Levels of oxalate production of each strain were measured by using oxalate assay kit (Libios).

Conclusions
Glyoxylate cycle is controlled positively by QS and is important for oxalate biosynthesis in B. glumae.
Background

*Ralstonia solanacearum* causes a destructive disease called “bacterial wilt” in numerous plant species. Its virulence is controlled by the transcriptional regulator PhcA, the activity of which is, in turn, regulated in a cell-density dependent manner, termed quorum sensing.

Objectives

We herein described the identification and characterization of ralfuranones J–L, new PhcA-regulated secondary metabolites, and the known derivatives, ralfuranones A and B, from *R. solanacearum* strain OE1-1.

Methods

Their structures were determined by spectroscopic and chemical methods. These ralfuranones were also detected in vascular exudates from host plants infected with OE1-1. Deletion of *ralA*, which encodes an enzyme for ralfuranone biosynthesis, reduced the virulence of OE1-1 in tomato plants. Virulence was restored by complementation of the *ralA* gene.

Conclusions

The results suggest that ralfuranones play important roles in the virulence of OE1-1.
LOSS OF QUORUM SENSING SYSTEM CAUSES OUTER MEMBRANE VESICLE FORMATION IN BURKHOLDERIA GLUMAE

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Background
Burkholderia glumae possesses one LuxI-R type quorum sensing (QS) system. The QS system of B. glumae activates expression of oxalate biosynthesis genes to avoid ammonia-mediated alkaline toxicity as results of amino acid catabolism. However, QS mutants do not produce oxalate, which causes population crash in stationary phase.

Objectives
We aimed to address a question whether QS mutants experience envelop stress to cause vesicle formation.

Methods
Outer membrane vesicles (OMVs) were purified from QS mutants using density-gradient sedimentation techniques. Total proteins were identified from the OMVs by LC-MS/MS analyses. The membrane ultrastructure of QS mutants and their vesiculation were visualized by transmission electron microscopy (TEM) after ultrathin sections.

Conclusions
Total 368 vesicular proteins were identified from the OMV proteomes. Most vesicular proteins were found to be involved in molecular functions associated with transporter, protein secretion, and protein localization. TEM revealed that periplasmic space of QS mutants is swollen due to hyperhydration in exponential phase. Previous metabolome analyses showed that serious imbalance of glutamate in QS mutants, which suggests that QS mutants might experience a turgor pressure problem. We propose that OMVs of B. glumae might be induced to relieve turgor stress caused by imbalanced metabolism.
THE ACTIVATION OF STAPHYLOCOCCUS EPIDERMIDIS SORPTION BY THE AUTGENOUS LOW MOLECULAR WEIGHT FACTORS

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Background: The bacterial occupation of a surface can be associated with the production and realizing of compounds which can optimize and accelerate this process.

Methods: The exponential phase cells of Staphylococcus epidermidis strain GISK 33 were washed twice (0.85% NaCl) and diluted to 10⁷ CFU/ml (PBS, pH 7.2). The cell-free medium filtrates were obtained after the S.epidermidis sorption on a polystyrene and glass surface for 60 min. The effect of filtrates media on the bacterial cell adhesion was evaluated. In addition these filtrates were subjected to MALDI-TOF analysis and treatment by trypsin and proteinase K (100 mkg/ml, 37°C, 2 h) with followed by enzymes separation by membrane filtration (10 kDa).

Conclusions: The cell-free medium after S.epidermidis adhesion to the polystyrene had a stimulating effect on the adhesion of the same bacterial cells on the polystyrene surface compare to the filtrates after the bacterial contacting with the glass surface. The level of found by MALDI-TOF analysis compounds with low mol. weight 552, 574 and 596 Da in the filtrate from polystyrene which showed the greatest stimulatory effect on bacterial adhesion was higher than the glass filtrate. The peptide nature of these compounds was proved by the reduction of the stimulatory effect on the S.epidermidis 33 adhesion to the control level (PBS) after the treatment by proteases. Thus, the evidence of the possibility of autoregulation of S.epidermidis cell adhesion by using the low molecular weight peptide compounds was provided.

This work was supported by RFBR (14-04-00687).
Background
The enterotoxin produced by Clostridium perfringens is associated with the sporulation process. The regulation of this process is therefore of interest. We previously described the presence of a low MW, heat- stable compound in the culture fluids of this organism which stimulated the sporulation of homologous and heterologous strains, including enterotoxin-positive strains.

Objectives
Isolate and characterize the presumptive peptide involved in promoting sporulation of Clostridium perfringens

Methods
The peptide was isolated from a defined medium following Sephadex LH-2, SP Sephadex, DEAE Sephadex chromatography followed by HPLC using C8 and C18 columns. Mass spectroscopy, amino acid analysis, and N-terminal sequencing revealed a 1018 Da peptide composed of glutamic acid, glycine, alanine, and an unidentified residue. N-terminal sequencing suggests a branched chain or cyclic structure. The isolatee compound was effective in stimulation sporulation of C. perfringens in a dose-dependent manner at levels <2 nm/l and differs from sporulation-stimulating peptide previously identified from Bacillus subtilis and which was ineffective in promoting sporulation of C. perfringens.

Conclusions
We believe the peptide functions as a quorum-sensing molecule. The identification of a sporulation signal could help identifying the molecular events leading to sporulation and enterotoxin production by this organism.
QUORUM SENSING INHIBITORY AND ANTI-BIOFOIOM ACTIVITY OF DIETARY PLANTS CULTIVATED IN KOREA

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Background
The discovery of new agent is demanded by the appearance of multidrug resistant bacteria, like super bacteria, for many years. Also, quorum sensing, cell-to-cell communication, is regarded as remarkable target for development of new antibiotics in recent

Objectives
In this study, we investigated antibacterial and anti-quorum sensing (anti-QS) capacity of dietary plants cultivated in Korea and their effect on formation of bacterial biofilms.

Methods
Inhibition of quorum sensing (QS) was investigated using the biosensor bacteria Chromobacterium violaceum and Pseudomonas aeruginosa PA01. The crude extracts of five plants were evaluated on antibacterial activity against five major food pathogen by bi-layer agar well method.

Conclusions
Especially, the extract of Zingiberaceae plant inhibited production of violet pigment in C. violaceum at 10mg/ml and the extracts of Punica granntum L. and Prune mume showed anti-QS activity against C. violaceum. Biofilm formation was measured by crystal violet and resazurin staining and the extract from Zingiberaceae plant had lower minimal inhibitory concentration than that of Citru junos and Punica granntum L. All of the extracts showed antibacterial activity against S. aureus, L. monocytogens, and B. cereus. The extract of Punica granntum L. also had high activity against E. coli and S. Enteritidis. The extract from Zingiberaceae plant had lower minimal inhibitory concentration (MIC) than that of Punica granntum L. against gram positive bacteria. The extract of Zingiberaceae plant had the lowest MIC of 50ug/ml against S. aureus and L. monocytogenes. These results exhibit the potential of five dietary plants to use as food preservatives and new antibacterial agents.
Background

The regulation of biofilm–formation in many pseudomonads is controlled by quorum sensing (QS) and is key to their ability to colonize different environments. However, in the soil and plant-associated *P. fluorescens* SBW25, mutations in diguanylate cyclase-associated genes increase c-di-GMP levels to induce attachment and cellulose expression, resulting in the formation of biofilms at the air-liquid interface of static microcosms, and no involvement of QS-dependent behaviour has been reported.

Objectives

The objective of the study was to examine the ability of SBW25 to quorum sensing.

Methods
Bioinformatics, cultivation assays, TLC and CLSM were used.

Conclusions

A bioinformatics analysis of the SBW25 genome has identified putative N-acyl homoserine lactone (AHL) and α-hydroxy ketone (AHK)-dependent QS pathways, including an AHL/AHK synthase-like protein belonging to the HdtS family (PFLU0050) and a CqsA-like protein (PFLU5614). Furthermore, a putative link can be established between QS and the regulation of c-di-GMP levels in SBW25 based on the TpbA/TpbB system of *P. aeruginosa*. In this preliminary work, bioassays with AHL/AHK reporter strains overlaid onto TLC of liquid culture extracts demonstrate that SBW25 produces detectable levels of AHL and AHKs. Although these quorum compounds have yet to be identified, tests using exogenous dodecanoyl homoserine lactone suggest that in SBW25, biofilm structure, eDNA, and possibly siderophore production, may all be regulated by QS pathways. This work is the first to provide experimental proof that SBW25 is capable of responding to AHL/AHK quorum signals like many other pseudomonads, and to suggest that biofilm-formation may also be AHL-regulated under different environmental conditions.
CELL-CELL COMMUNICATION PROVIDES FITNESS BENEFITS TO POPULATIONS OF LISTERIA MONOCYTOGENES IN THE SOIL ENVIRONMENT

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Background

Listeria monocytogenes is a ubiquitous opportunistic human pathogen detected in many habitats spanning from the farm environment to food industry and the gastrointestinal tract.

Cell-cell communication participates to the adaptation of bacteria to their environment. In the species Listeria monocytogenes, the Agr system is required for full virulence and biofilm formation but its actual role is still poorly understood.

Objectives

To investigate whether the ability to communicate provides a benefit to L. monocytogenes in soil, a complex environment combining biotic and abiotic characteristics.

Methods

Deletion of the gene coding the response regulator or the signal propeptide. Population dynamics of the mutant and/or parental strains in soil microcosms. Both unsterilised and sterilised soils were investigated.

Conclusions

Deletion of the gene coding the regulator or the signal did not affect population dynamics in sterilized soil but survival was altered in biotic soil suggesting that the Agr system was involved to face the complex soil biotic environment. This was confirmed by co-incubation experiments. The fitness of the response negative mutant was lower either in the presence or absence of the parental strain but the fitness of the signal negative mutant depended on the strain with which it was coincubated. Survival of the signal negative mutant was higher when cocultured with the parental strain than when cocultured with the response negative mutant. These results showed that the ability to respond to communication provided a benefit to listerial cells. These results might also indicate that in soil, the Agr system controls private goods rather than public goods.
SCREENING OF QUORUM SENSING INHIBITORS WITH AN E. COLI BIOSENSOR AND ENHANCEMENT OF THEIR QUORUM QUENCHING ACTIVITY BY ENCAPSULATING THEM IN CHITOSAN-BASED NANOCAPSULES

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Background
The quorum sensing (QS) hypothesis explains a type of bacterial cell-cell communication which is mediated by means of autoinducer low molecular weight exocellular compounds. This phenomenon has profound implications in the control of many important bacterial traits (e.g. biofilm formation, secretion of virulence factors, etc.). A typical QS system in gram-negative bacterial involves the production and response to acylated homoserine lactone (AHL) signals that are recognized by the LuxR receptor. Several strategies have been conceived to block or disrupt QS (known as quorum quenching, QQ) and thus offer an alternative strategy to reduce the collective power of bacterial pathogens.

Objectives
In this regards, we have screened a library of total 24 purified compounds and have tested their QQ activity when applied at a concentration of 1mM.

Methods
To this end an E. coli Top 10 biosensor reporter of AHL-mediated QS which constitutively expresses green fluorescent protein (GFP) upon exogenous addition of 3-oxo-hexanoyl homoserine lactone (3OC\(_6\)HSL) has been used. A computer-based docking approach was used to elucidate in further detail the interaction between the hit compounds with the TraR protein 3D crystal structure.

Conclusions
The hit compounds (i.e. able to inhibit significantly the production of GFP but without inducing a toxic response) were vanillin, trans-cinnamaldehyde, caffeine, genipin, PQS (Pseudomonas quinolone signal) and MOQ (1H-2-methyl-4-quinolone). We have also tested the effect of loading the hit compounds into chitosan-based nanocapsules on their QQ activity and found that this is a strategy to further control and modulate their QQ bioactivity.
A LARGE SET OF GENES LINKED TO QUORUM SENSING-DEPENDENT REGULATION IDENTIFIED BY A GENOME WIDE TRANSCRIPTOME ANALYSIS OF SINORHIZOBIUM FREDII NGR234

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Background

The alphaproteobacterium Sinorhizobium fredii NGR234 is outstanding and unique among the rhizobia with its ability to form nitrogen-fixing nodules with a wide range of legumes (1). Beside many other striking features, its 6.9 Mbp genome encodes for two N-acyl-homoserine-lactone synthase genes (i.e. traI and ngrI) which are involved in the biosynthesis of two distinct autoinducer I-type molecules (2,3).

Objectives

Here we report on the construction of a NGR234-ΔtraI and a NGR234-ΔngrI mutant and their genome wide transcriptome analysis. The high-resolution RNA-seq analysis of early stationary phase cultures in the NGR234-ΔtraI background suggested that up to 316 genes were differentially expressed in the NGR234-ΔtraI mutant vs. the parent strain and 466 in the background of NGR234-ΔngrI vs. the parent strain. Accordingly, a common set of 186 genes was regulated by the TraI/R and NgrI/R regulon including flagella biosynthesis genes and genes linked to EPS succinoglycan biosynthesis. Among the genes that were differentially regulated in NGR234-ΔtraI were those linked to replication of the pNGR234a symbiotic plasmid and cytochrome c oxidases. In the NGR234-ΔngrI mutant biotin and pyrroloquinoline quinone biosynthesis genes were differentially expressed as well as the entire cluster of the NGR234 type III secretion system (T3SS-II). Further we also discovered that genes responsible for rhizopine catabolism in NGR234 were strongly repressed in the presence of high levels of N-acyl-homoserine-lactones.

Methods

high-resolution RNA-seq analysis

Conclusions
Together with nodulation assays, our RNA-seq-based findings suggested that QS-dependent gene regulation appears to be of higher relevance during non-symbiotic growth rather than for life within root nodules.
REGULATORY RNA ARISES FROM 3´-UTR OF AHL SENSOR ENCODED RNA AFTER ITS PREPROCESSING IN PECTOBACTERIUM ATROSEPTICUM

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Background
In a broad group of proteobacteria two quorum sensing-related genes, which encode synthase of autoinducers and their sensor, respectively, have convergent topology and overlap by their 3´-ends. It was suggested that the expression of one gene may antagonize the transcription of another one because of the convergent arrangement of these genes.

Objectives
We proposed that the topology of quorum sensing-related genes of plant pathogenic bacterium Pectobacterium atrosepticum, expI and expR, determines the mechanism of regulation of their expression.

Methods
We noticed that expR gene lacks any obvious transcriptional stop signals. Using chain-specific RT-qPCR we found that at the stationary growth phase expR mRNAs having long 3´-untranslated region (UTR) were formed and the expression of oppositely oriented expI gene was simultaneously decreased. It is likely, that RNA products of the expR regulatory gene may inhibit the expression of the autoinducer synthase gene through the interactions with expI mRNA. Using 3´-RACE method we found that at the stationary growth phase the extended expR transcripts underwent preprocessing resulting in their break down into two fragments. The first fragment encompassed the entire length of open reading frame (ORF) of expR gene and the second one included only a long 3´-UTR having a predicted regulatory function.

Conclusions
We found that cis-encoded RNA may participate in the regulation of expression of two convergent functionally related genes. The length of one of two transcripts varies respective to the cell growth phase affecting the expression of oppositely oriented gene. Moreover, cis-encoded regulatory RNAs arises from protein encoded RNAs that undergo preprocessing in their 3´-UTR.
Background
Burkholderia pseudomallei, the causative agent of melioidosis, is a gram-negative soil bacterium in tropical areas. B. pseudomallei employs several N-acyl-homoserine lactone (AHL)-mediated quorum sensing (QS) systems, which activate specific sets of genes as a function of cell density [1]. The genome of B. pseudomallei encodes genes for three QS systems with one luxI and one luxR homologue, respectively, and additionally three orphan luxR homologues [2]. The luxI homologues encode AHL synthases, which produce specific AHLS binding to the respective transcriptional regulator and thus regulating expression of specific genes involved in virulence like biofilm formation, siderophore biosynthesis or swarming motility [3].

Objectives
The aim was to clarify the influence of the different QS systems on virulence factor expression.

Methods
We constructed mutants in the synthase-coding genes. Subproteome fractions of the mutant strains were subjected to mass spectrometry analysis to identify targets of the different QS systems. We used the DIA approach IMS^E in combination with the Hi3 approach for quantification of cytosolic proteins and the GeLC MS/MS approach for the analysis of extracellular proteins. Furthermore, the AHL spectrum, synthesized by the different QS systems was determined by analytical thin layer chromatography.

Conclusions
The luxI homologues influence protein expression by up- and downregulation of AHL-dependent proteins in the QS circuitry of B. pseudomallei.
ENZYMATIC PRODUCTION OF MODIFIED STARCHES WITH SLOW DIGESTION PROPERTIES

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Background

In human nutrition, the proper rate of glucose release and absorption from digesting starch may play important roles in body health. Therefore, improving food quality with higher amounts of slow digestible starch is becoming an area of interest for researchers in academia and industry.

Objectives

The objectives of this study is to produce the modified starch with slow digestion properties by starch-modifying enzymes.

Methods

Two branching enzymes (BEs), PH1386 of Pyrococcus horikoshii OT3 and Athe_0558 of Caldicellulosiruptor bescii, were expressed in E. coli and purified by Ni-NTA affinity chromatography. The enzymatic properties of two BEs have been studied by high performance anionic exchange chromatography (HPAEC) analysis. The melting temperature range (Tr) and melting enthalpy (ΔH) representing the degree of retrogradation of modified starches were also studied by differential scanning calorimetry. Finally, the digestion properties of modified starches were examined by pancreatic enzyme and α-glucosidase.

Conclusions

HPAEC analysis of BE-modified starches revealed that PH1386 produces the branch chains with mainly DP 10-12, while Athe_0558 prefers DP 6-7. The proportion of A chains (DP ≤12) of amyllopectin was increased and the proportion of B1 (DP 13-36) and B2 chains (DP ≥37) was decreased compared to control. The Tr and ΔH also decreased compared to control, suggesting that retrogradation rate of the BE-modified starches was delayed. The increased resistant starch contents and
decreased rapidly digestible starch contents of BE-modified starches are thought to be due to the increase in α-1,6 linkages and the highly branched structure of starch.
MAXIMIZING BIOTECHNOLOGICAL PRODUCTION OF 3-HYDROXYPROPIONALDEHYDE AS A NATURAL FOOD PRESERVATIVE WHILE MINIMIZING ITS CONVERSION TO TOXIC ACROLEIN

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Background

3-hydroxypropionaldehyde (3-HPA) has strong antimicrobial activity and application potential as a natural food preservative for disinfection of minimally processed vegetables. During glycerol metabolism, Lactobacillus reuteri DSM 20016T can accumulate 3-HPA, which undergoes reversible dimerization and hydration, and can be further converted enzymatically into inactive 1,3-propanediol (1,3-PD), restoring intracellular redox potential. Dehydration of 3-HPA resulting in acrolein, a toxic metabolite, is also possible, however, conditions promoting this transformation have not been fully understood due to the lack of analytical methods to simultaneously quantify 3-HPA as well as acrolein.

Objectives

The aim of this study was to maximize the capacity for biotechnological processing of glycerol to 3-HPA using L. reuteri, while accounting for the accumulation of acrolein after production and storage to evaluate toxicological risk.

Methods

Strategies to improve 3-HPA production included the addition of redox-active compounds during glycerol fermentation to prevent conversion to 1,3-PD, the usage of immobilized cells resulting in high densities of viable cells as well as the optimization of process parameters in batch and fed-batch processes. Special attention was paid to cultivation conditions of L. reuteri prior to glycerol conversion. Acrolein and 3-HPA were simultaneously quantified using a newly established analytical method employing ion-chromatography with pulsed amperometric detection.

Conclusions

Redox-active compounds, process conditions and pre-treatment of L. reuteri cultures influenced 3-HPA yield as well as ratio of 3-HPA to 1,3-PD. Temperature and pH
optimization prevented formation of acrolein during biotechnological production and storage, promoting the safety of 3-HPA when applied as natural food preservative.
COMPARATIVE STUDY OF FERMENTATION PROCESSES BY FREE AND IMMOLIZED GRAPEVINE INDIGENOUS STRAINS FOR THE PRODUCTION OF HIGH QUALITY WINES


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Background
Vinification is a complex microbiological process. Immobilized microorganisms constitute an alternative approach aiming to lower the overall cost of production and improve wine quality.

Objectives
This study focuses on the immobilization of indigenous Saccharomyces and non-Saccharomyces strains for use in wine production.

Methods
Saccharomyces cerevisiae strains are known for their ability to produce high ethanol levels, while non-Saccharomyces species contribute to the enhancement of the wine flavor resulting in high quality wines. Two indigenous environmental yeast strains isolated from a Greek agricultural region (Zitsa, Epirus), Saccharomyces cerevisiae Z622 and Metschnikowia pulcherrima var Zitsae ZY6 were used as starter cultures in a sequential fashion for wine fermentation. Delignified cellulose is used as an immobilization support meeting all requirements and demands for use in food production and wine making.

Conclusions
Both yeast strains were successfully immobilized maintaining their metabolic activity during fermentation. Immobilized cells showed a better overall metabolic activity compared to control free cells.

Acknowledgments
This project is co-financed by the European Union (European Regional Development Fund — ERDF), through the operational programs for “competitiveness and entrepreneurship” and regions in transition “Cooperation 2011 — Partnerships of Production and Research Institutions in Focused Research and Technology Sectors”, of the National Strategic Reference Framework (NSRF) 2007–2013, and the Hellenic Ministry of Education, Lifelong Learning and Religious Affairs — General Secretariat for Research and Technology.
LACTOBACILLUS SAKEI ISOLATED FROM FERMENTED MEAT: EXTERNAL ANTIBACTERIAL ACTIVITY AGAINST LACTIC ACID BACTERIA AND PATHOGENIC STRAINS

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Background

Lactobacilli play an important role in the production of fermented sausages, due to their antimicrobial activity and their effect on the organoleptic properties of meat products. Lb. sakei is gaining relevance because it produces bacteriocins with antilisterial activity. However, the use of bacteriocins may be limited due to their action against phylogenetically related species and ineffectiveness against some pathogenic and undesirable spoilage bacteria. Another group of proteins with antibacterial activity are peptidoglycan hydrolases, which so far have not been reported for this bacterial species and probably play a role in food safety and in the organoleptic properties of meat products; because they prevent the growth of pathogens and spoilage bacteria, including heterofermentative lactic acid bacteria (LAB).

Objectives

The aims of this work were to identify lactic acid bacteria isolated from fermented meat, as well as the proteins responsible for their extracellular antibacterial activity against different pathogenic and LAB strains.

Methods

Isolated lactobacilli were identified by 16S rDNA sequencing. Antimicrobial and lytic activities were evaluated in the culture supernatant by agar diffusion tests and zymography, using in both cases different target microorganisms. A 79-kDa protein was identified which showed antimicrobial activity. Protein identification was performed by LC/MALDI TOF/TOF and the data showed a high correspondence with a putative N-acetylglucosaminidase.

Conclusions

A Lb. sakei strain isolated from fermented meat which has displayed external antibacterial activity has been cultured and studied for the first time. The aforesaid activity is probably related to a protein with peptidoglycan hydrolase activity. This may have a potential use in food preservation, considering that this activity was obtained from a bacteria isolated from a meat product.
Background

Gut microbiota has numerous roles in human life and its significance has thus far been underestimated. Therefore, more detailed studies on diversity and metabolism of gut microbiota are needed.

Objectives

Gut microbiota can be shaped by prebiotics. Levan-type (β 2,6-linked) fructans can be considered as potential prebiotics\(^1\). We have synthesized levan and levan-type fructooligosaccharides by heterologously expressed levansucrase Lsc3 of *Pseudomonas syringae* pv. tomato. These substrates should affect gut microbiota as were perfectly fermented by a gut generalist *B. thetaiotaomicron*\(^2\). Here we focus on structure-function relationships of the Lsc3 protein and its biotechnological potential.

Methods

The His-tagged Lsc3 protein was site-directedly mutated, biochemical methods were used to characterize the mutant proteins and 3D structure modelling was applied to interprete the results.

Conclusions

Levansucrase Lsc3 has high polymerizing activity (up to 80%) and extremely high stability confirming its biotechnological potential for the synthesis of fructans. Several novel catalysis-related positions for levansucrases were revealed and their predicted location on a 3D model of Lsc3 will be presented.

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References:


MATCHING GENOME AND TRANSCRIPTOME OF LACTOCOCCUS LACTIS STRAINS WITH ROBUSTNESS TOWARDS INDUSTRIAL STRESSES

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Background

Lactococcus lactis is industrially employed to manufacture fermented dairy products. Spray drying is the cheapest preservation method for L. lactis starter cultures, but during this process cultures encounter heat and oxidative stress, resulting in low survival. Viability of starter cultures is essential for their role in dairy acidification, supporting the need to understand and improve their robustness.

Objectives

We have measured the ability of 39 L. lactis strains to survive industrially relevant stresses. This set of strains contained different L. lactis strains from dairy as well as plant origin. The observed 4-log variation in heat and oxidative stress survival was compared with genomic content, resulting in the identification of genes associated with robustness. Presence and activity of genes can play an important role in the observed diversity in robustness.

Methods

Therefore, four L. lactis strains with varying robustness phenotypes were fermented under twelve different conditions, varying in temperature, salt concentration, pH, and oxygen level. Cells were harvested at exponential phase of growth for transcriptome analysis and survival measurements.

Conclusions

The varying growth conditions resulted in up to 4-log differences in robustness towards heat and oxidative stress. Moreover, clear differences in gene expression profiles were observed. Correlation of robustness phenotypes and gene expression levels revealed transcriptome signatures for oxidative and heat stress survival. For strain MG1363 this included the metC-cysK operon, involved in methionine and cysteine metabolism, which triggered us to grow this strain in the absence of
cysteine, resulting in elevated expression levels of the $\text{metC-cysK}$ operon and concomitant enhanced robustness towards oxidative stress.
Background
Brazzein is an intensely sweet-tasting protein with four disulfide bridges. In previous study, we constructed the secretory expression system of recombinant brazzein in yeast *Kluyveromyces lactis*.

Objectives
In this expression system, the control of protein disulfide-bond formation in the endoplasmic reticulum (ER) is often a bottleneck for secretory protein production. The major pathway for protein disulfide-bond formation in endoplasmic reticulum (ER) includes the conserved ER-membrane protein Ero1p and protein disulfide isomerase (PDI).

Methods
In the present study, the PDI gene was introduced in the yeast *Kluyveromyces lactis* to improve the secretory efficiency of brazzein. The expression of Ero1p was also induced by treatment of the cells with dithiothreitol. The culture condition of the yeast transformants for high yield secretion of the recombinant des-pE1M-brazzein was in YPGal medium for 96 hours at 30 °C.

Conclusions
The amount of misfolded or unfold recombinant des-pE1M-brazzein remaining inside the cell decreased and the amount of the secretory recombinant des-pE1M-brazzein having an intrinsic sweetness increased approximately 1.7-fold than that of the previous expression system. These results demonstrate that the modified *K. lactis* expression system could be applicable to mass production of the recombinant brazzein with attributes useful in the food industry.
IDENTIFICATION OF SULPHITE RESISTANCE WINE YEASTS 
SACCHAROMYCES CEREVISIAE FROM A POOL OF SELECTED STRAINS
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Background

Sulphite is frequently added in winemaking, hence sulphite resistance mechanisms have been studied extensively in Saccharomyces cerevisiae. Research turned out that yeasts possess plasma membrane protein encoded by the SSU1 gene (lies on chromosome XVI), which excretes sulphite from yeast cells. Some highly sulphite-resistance wine strains have an SSU1 allele – SSU1-R - on chromosome VIII. We tried to elucidate, how long adaptation to high sulphite concentrations affects to the evolution of yeast strains.

Objectives

The aim of experiment was to determine presence of SSU1 and SSU1-R genes and their expression in selected Saccharomyces cerevisiae strains.

Methods

After 10 years of storage with SO₂ (concentration ~ 400 mg/ml) the pool of 12 strains S. cerevisiae, include 11 sulphite strains and negative control strain, were tested for the presence of the SSU1 or SSU1-R gene. We analysed them by using PCR assays, gel electrophoresis and DNA sequencing. Obtained sequences were aligned, in CLUSTAL W tool, with known sequences from databases (BLAST and SGD). Gene expression was tested during alcoholic fermentation for few strains by real-time PCR.

Conclusions

The presence of the SSU1 gene was confirmed in 7 strains (include control strain) and SSU1-R gene in 3 strains. None of those genes was detected in 2 strains. The results indicate that possession of the SSU1 or its allele is independent of the long term storage in must with SO₂. It is well-known that evolution in yeast cells occur dynamic. In our research it was confirmed by the presence of SSU1 in negative control strain.
Background:

Citric acid is the most important and widely used organic acid produced by fermentation. Mineral composition of the fermentation medium has a critical role in citric acid production.

Objectives:

The aim of this study was to investigate the influences of different minerals on growth and citric acid production of *Yarrowia lipolytica* NBRC 1658 in a glucose medium.

Methods:

The mineral salts; FeSO$_4$, CuSO$_4$, MnSO$_4$ and ZnSO$_4$ were added to the fermentation medium separately with concentration ranges of 0.01-0.1, 0.001-0.02, 0.005-0.05 and 0.002-0.008 g/L, respectively. Experiments were performed in a batch system. Biomass concentration was determined spectrophotometrically. Concentration of citric acid was measured by pyridine-acetic anhydride method.

Conclusions:

In this study it was obtained that by the addition of FeSO$_4$ into the media, the specific growth rate of the yeast increased and although CuSO$_4$ addition had a positive effect on the decrease of the duration of the lag phase, it had an adverse effect on the citric acid production as well. The maximum specific growth rate of the yeast determined as 0.053 h$^{-1}$ by using 0.02 g/L of MnSO$_4$ in the glucose based medium. Although ZnSO$_4$ supported the growth of the yeast, citric acid concentration reduced and reached to a maximum value of 13.4 g/L. It was also concluded that citric acid production decreased by the supplement of the salts examined within the concentration ranges above, while certain concentrations of FeSO$_4$, MnSO$_4$ and ZnSO$_4$ had profound effects on the growth of the yeast.
ENHANCEMENT OF THE BIOAVAILABILITY OF EXTRACTABLE BERRY PHENOLICS BY SOLID STATE FERMENTATION

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Background
Increasing interest can now be observed for the utilization of plant-wastes rich in extractable phenols derived from fruit processing. Using these residues, free phenolics with high antioxidant potential can be liberated by solid-state fermentation, via the deglycosylation action of beta-glucosidases of the fermenting fungi. Thus, the bioavailability of the extractable phenols can be enhanced since aglycons get across the cell membrane easily.

Objectives
In our previous studies, isolates of *Rhizomucor miehei* and *Mucor corticulosis* showed intensive extracellular beta-glucosidase activity on plant-derived residues. Here, we analyzed the ability of these zygomycetes to generate free phenolics under jostaberry-based solid-state fermentation.

Methods
Solid pomace medium was supplemented with soy flour as nitrogen source. During incubation samples were taken every second or third day and extracted with distilled water, ethanol:water 50:50 or HCl:methanol 10:90 solutions. Besides beta-glucosidase activity, total phenolics and total anthocyanins were determined. Antioxidant activity tests including radical scavenging capacity and ferric reducing capacity assays were also performed.

Conclusions
The *R. miehei* and *M. corticulosis* bioconversion of jostaberry pomace leads to enhanced total phenolic content. Positive correlation between total phenolic content and beta-glucosidase activity was found in both systems. Ferric reducing antioxidant power was slightly increased during the first phase of fermentation; however, anthocyanin content showed significant drop after 3-5 days. Results indicate that *R. miehei* and *M. corticulosis* fermentations are applicable methods for liberation and enrichment of health-relevant extractable phenolics from jostaberry pomace. This research was supported by the Hungarian Research Fund (OTKA PD 112234).
INFLUENCE OF STARTER INOCULUM ON DYNAMIC GROWTH OF 
STREPTOVERTICILLIUM MOBARAENSE IN O3 MEDIUM. 

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Background

Streptoverticillium mobaraense KKP 2013 strain obtained from the Culture Collection of Industrial Microorganisms is Gram-positive Actinomycetes commonly occurring in natural environment. Under certain conditions, it has the ability to produce the enzyme MTG, protein-glutamine gamma-glutamyltransferase, EC 2.32.13, commonly called transglutaminase. Transglutaminase catalyzes the formation of a covalent bond between the proteins, it allows also crosslinking reactions, namely the production of new intra- or intermolecular bonds between the proteins and peptides. MTG due to its characteristics is widely used in the production of meats, dairy and bakery products.

Objectives

In the process for producing transglutaminase one of the important steps is to obtain an appropriate number of microorganisms, which allows to obtain the highest possible quantity of active enzyme.

Methods

In our study, we tried to determine the growth rate of strain KKP 2013 S. mobaraense in the proliferating medium O3. In the precise number of microorganisms on the breeding stage, we used the method of pour plate technique on PCA medium, according to PN-EN ISO 4833-1: 2013-2014.

Conclusions

We established the optimal volume of medium in relation to the number of bacteria in the starting inoculum, which will produce the highest activity of the resulting enzyme.
USING DYNAMIC SINGLE CELL BIOLOGY TO DISSECT INJURY, RESUSCITATION AND DEATH PHENOMENA IN STRESSED POPULATIONS OF ESCHERICHIA COLI

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Background

While inimical food preservation approaches often result in heavily damaged and stressed populations of foodborne pathogens and spoilage microorganisms, very little is known about the actual molecular and genetic events and dynamics that govern the (sub)lethal injury and subsequent resuscitation or death of these cells. Nevertheless, insights into these phenomena might be decisive for a proper understanding of the resulting behavior and evolvability of the surviving subpopulation.

Objectives

In this study, we set out to monitor and dissect (sub)lethal injury and resuscitation phenomena in populations and cells of Escherichia coli that are stressed by heat or high hydrostatic pressure treatment.

Methods

Directed evolution, genetics and time-lapse fluorescence microscopy.

Conclusions

Based on adaptive mutations, genetics and live cell biology, different subcellular processes and structures have been delineated and monitored that play a role in the heterogeneous injury and resuscitation dynamics of high temperature or pressure stressed E. coli cells and populations, thereby revealing a surprising impact on the spatial dynamics of nucleoids and protein aggregation inside stressed cells.
Background

*Staphylococcus aureus* is a pathogen of major concern for clinical infection and foodborne illness and has remarkable invasiveness and toxin-mediated virulence. Studies have shown that plant essential oils can influence production of toxins, avoiding overuse of antibiotics and the development of bacterial resistance.

Objectives

This study aimed to verify the influence of major terpenoid phenols compounds from plants essential oils in subinhibitory doses on enterotoxins A, B, C and D produced by *Staphylococcus aureus*.

Methods

Five phenolic compounds (eugenol, geraniol, cinnamaldehyde, citronellol and terpineol) were tested. Amounts corresponding to 60% and 80% of Minimum Inhibitory Concentration (MIC90%) were added to culture medium Tryptic Soy Broth (TSB). Each treatment was inoculated with a suspension of *S. aureus* producer of each enterotoxin A, B, C and D and incubated (37°C/24h). TSB without compounds was used as positive control. After incubation each treatment was centrifuged (9000g/4°C/30min) and supernatant was used to detect enterotoxin production by Reverse Passive Latex Agglutination. Interpretation of results was performed according to manufacturer’s instructions kit. Agglutination was characterized as: strong (+++); moderate (++); weak (+) and absence (-) of enterotoxin production. Assays were performed in duplicate. Kruskal-Wallis nonparametric test was performed using scores related to enterotoxin production.

Conclusions

Some phenolic compounds from plants essential oils tested demonstrated great capacity to inhibit enterotoxins produced by *S. aureus* and, consequently decreasing its virulence. Therefore, it may be a promising way to prevent diseases caused by enterotoxigenic *S. aureus* and to contribute for reduction of antimicrobial drugs consumption.
CHARACTERIZATION OF THE DAIRY STREPTOCOCCUS THERMOPHILUS ACA-DC 29 STRAIN THROUGH COMPARATIVE GENOMICS

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Background
Although the Streptococcus genus includes mainly pathogenic species, Streptococcus thermophilus is a widely used dairy starter culture of great economic importance for the food industry. S. thermophilus has been adapted to milk probably through a degenerative evolution process that has led to the loss of typical streptococcal pathogenic traits.

Objectives
The genome sequence of the yogurt isolate S. thermophilus ACA-DC 29 was analyzed for assessing the technological potential of this strain. Comparative genomics analysis was also performed between the genome of ACA-DC 29 and the existing complete genome sequences of S. thermophilus.

Methods
The genome sequence of ACA-DC 29 was annotated using online annotation tools. Full chromosome alignments were calculated with Progressive Mauve. The pangenome, the core genome and the unique genes were predicted with the GView Server. The genomic islands, the CRISPRs and the antimicrobial peptides were predicted with IslandViewer, CRISPRcompar and BAGEL3, respectively.

Conclusions
The analysis of the S. thermophilus ACA-DC 29 genome sequence revealed the absence of pathogenic features. Genes related to the adaptation to milk were identified. Full chromosome alignments showed a high degree of synteny among the different strains. The pangenome of the eight strains comprised of approximately 2,300 genes. Concerning the ACA-DC 29 strain, approximately 250 unique genes involved in various biological processes were also identified. Further analysis indicated that several of them may have been acquired through horizontal gene transfer. Five potential antimicrobial peptides and two CRISPR systems, which may confer resistance against phages, were also predicted.
GREEK TRADITIONAL DAIRY AND MEAT PRODUCTS: A BIOLOGICAL RESERVOIR FOR NEW PROBIOTIC STRAINS
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Background

Probiotics provide health benefits to the host upon ingestion. Lactic acid bacteria are frequently used as probiotics. Important qualifying characteristics of probiotics strains are the acid and bile resistance, the absence of pathogenic traits, the elevated adhesion ability, the production of antimicrobial substances and the inhibition of pathogens. Traditional fermented foods constitute a promising source for discovering new probiotic strains.

Objectives

Lactic acid bacteria isolated from Greek traditional dairy (Yogurt and Feta, Kasseri, Xynotyri, Graviera, Kopanisti, Formaella, Galotyri, Kefalotyri cheeses) and meat (Lountza and Salami) products, were screened for their probiotic potential.

Methods

The isolated strains were evaluated for survival to low pH and in the presence of bile salts. The safety of selected strains was assessed by studying their haemolytic activity and their susceptibility to commonly used antibiotics. Antimicrobial activity against pathogens, was also examined. Finally, the ability of the strains to adhere to collagen-coated 96-well microplates and to the human colorectal adenocarcinoma cell lines HT-29 and Caco-2 in vitro, was investigated.

Conclusions

Of the 128 isolated strains, thirty were selected for further study due to their robust performance under the adverse conditions of the gastrointestinal tract. None of the strains exhibited haemolytic activity while variable antibiotic resistance was detected. Two Streptococcus thermophilus strains presented inhibitory activity against Streptococcus anginosus LMG 14502T and Streptococcus mutans LMG 14558T. Three Lactobacillus plantarum strains were found to adhere better to both cell lines,
as well as to the collagen-coated microplates. The utilization of *in vitro* tests enabled the selection of strains with promising probiotic features.
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Food microbiology

NATIONWIDE OUTBREAK OF LISTERIOSIS DUE TO READY-TO-EAT SALAD IN SWITZERLAND

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Background
Listeria monocytogenes is an important foodborne pathogen with a significant impact on public health and economy worldwide. Human infections are rare, but L. monocytogenes has the potential to cause serious and life-threatening disease. Clinical conditions primarily comprise septicemia, meningitis, meningoencephalitis and abortion.

Objectives
To describe a nationwide outbreak of listeriosis having occurred during 2013 to 2014 in Switzerland.

Methods
Microbiological methods (serotyping, MLST, PFGE) and epidemiological investigations by telephone interviews were used.

Conclusions
From 26 October 2013 to 23 April 2014, 32 cases of listeriosis infected with an L. monocytogenes strain serovar 4b, sequence type 4 and belonging to a single distinct PFGE pulsotype were registered in patients from several cantons of Switzerland. L. monocytogenes was detected in blood (75%), CSF (16%), ascites (6%) and in joint fluid (3%) samples. By the end of March 2014, a retail company reported an L. monocytogenes contamination of ready-to-eat salads to the authorities after detecting the pathogen through its in-house routine quality control. Product and environmental samples collected during subsequent investigations yielded isolates, matching the outbreak strain, thus confirming that ready-to-eat salad from this company was most likely the outbreak source. The cause for the product contamination was related to a design-inherent hygienic problem of one specific product-feeding belt. Complementary patient interviews also identified ready-to-eat green salads bought at one retailer as the likely outbreak source.
RAPID AND ACCURATE LISTERIA SPP. IDENTIFICATION BY MALDI BIOTYPER DATABASE IMPROVEMENT

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Background
Listeria species are closely related to each other, however, only L. monocytogenes and L. ivanovii exhibit pathogenic features, (human cases of L. ivanovii infection being quite rare). Rapid, accurate identification of Listeria strains is essential for appropriate management and timely intervention for infection control.

The accuracy and speed of data acquisition by MALDI-TOF MS makes this an important tool for biological public health hazards, food processing, quality control, and disease diagnoses.

Objectives
Since Listeria species are closely related and have similar profile spectra, the objective was to reliably identify the different Listeria species by MALDI-TOF MS through establishment of a reliable library of mass spectral Listeria fingerprints from reference strains. Therefore the database was enhanced by 52 Listeria strain references, from 9 species.

Methods

Listeria strains were grown on Columbia Agar with 5 % Sheep Blood agar (BD, Heidelberg, Germany) at 37 °C. Biomass was collected after one day, and then processed by ethanol / formic acid standard extraction method. Mass spectra were acquired by using microflex LT (Bruker Daltonik, Bremen, Germany) and identified with MALDI Biotyper Compass software.

Conclusions

With the new, optimized references database, it was possible to identify all tested Listeria strains from 9 species. This was proven by analysis of more than 1200 mass spectra.

Although log (score) value differences were small between Listeria species, the correct species was always the best match, no false identification occurred. In consequence, the important identification of the pathogenic Listeria species is possible with easy, rapid and reliable low-cost MALDI-TOF MS method.
INCIDENCE OF ASPERGILLUS PRODUCER OF OCHRATOXIN A IN THE TROPICAL VITICULTURE IN THE NORTHEAST OF BRAZIL
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Background
The biodiversity of filamentous fungi is one of the most important contributions to the occurrence of mycotoxins in agricultural products. The species belonging to the genus Aspergillus has been blamed for the presence of mycotoxins in wines. The incidence of these genera in grapes will depend on climatic conditions, grape variety, cultivation form and agricultural practices.

Objectives
This study aimed to identify Aspergillus species isolated from wine grapes at harvest, from organic and conventional vineyards, in the São Francisco Valley.

Methods
Varieties evaluated were Touriga Nacional and Ruby Cabernet, and Tempranillo. For the isolation of fungi direct plating was carried out in DRBC culture medium, 25°C/7 days. The isolates were identified using standard culture media. The determination of toxigenic potential was performed by Thin Layer Chromatography.

Conclusions
There was no presence of Aspergillus in the Tempranillo. All A.carbonarius were producers of ochratoxin A. The presence of these species can spoil the grapes. All species identified are naturally present in the vineyards.
INCIDENCE OF OCHRATOXIGENIC SPECIES OF ASPERGILLUS AND OCHARATOXIN A ON COFFEE BEANS FROM CONVENTIONAL AND ORGANIC CULTIVATION

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Background
The natural microbiological contamination of fruit and coffee beans, are due to the presence of bacteria, yeasts and filamentous fungi, including fungi species of the genus Aspergillus ochratoxin-producing (OTA) has great relevance for coffee security.

Objectives
The objective of this study was to identify ochratoxigenics fungi and quantify the presence of OTA in conventional and organic cultivation of coffee beans in the southern region of Minas Gerais (Brazil) and compare the concentrations in both systems.

Methods
We analyzed 30 samples of coffee beans (Coffea arabica L.), 20 samples of conventional coffee and 10 organic coffee samples. The incidence of OTA in the samples of coffee beans was performed by high performance liquid chromatography (HPLC).

Conclusions
Samples were identified 480 fungi of the genus Aspergillus Section Nigri and Circumdati. The main producing species Aspergillus ochraceus was OTA (83% of the samples). For the OTA in the grain, only one sample collected organic coffee ground, presented OTA, 1.12 ug / kg. Considering these results, it can be stated that the presence of Aspergillus ochraceus OTA producer does not imply contamination of grain by mycotoxins and two coffee cultivation systems have the same risk of contamination.
RATIONAL SELECTION OF YEASTS BASED ON THEIR PECTINOLYTIC ACTIVITIES AND ITS INCIDENCE ON TECHNOLOGICAL AND SENSORIAL ASPECTS OF WINE QUALITY

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Background

Pectinase enzymes have shown a considerable influence in both, sensitive and technological properties of wines. They can help to improve clarification and filterability process, releasing more color and flavor compounds entrapped in the grape skin and facilitating the liberation of phenolic compounds. The addition of commercial enzyme preparations, with filamentous fungi as its main source, can be quite expensive for industry.

Objectives

This work aims to find yeasts that, because of their native pectinases, can be applied on combined fermentations with Saccharomyces cerevisiae obtaining significant benefits over traditional wine fermentations.

Methods

785 yeast strains isolated from wineries were identified and tested for several enzymatic activities of recognized interest for enology industry. The impact of Metschnikowia pulcherrima as a source of pectinolytic enzymes during wine fermentation was analyzed by measuring its influence on filterability, turbidity and the increase on color, anthocyanin and polyphenol content of wines fermented in combination with Saccharomyces cerevisiae. Further metabolites with enological interest were analyzed during the entire fermentation period.

Conclusions

Positive results were obtained in all expected parameters when M. pulcherrima was used by comparing wines fermented with S. cerevisiae alone and combined with Kluyveromyces thermotolerans, even working better than commercial enzymes preparations in most parameters. Additionally, M. pulcherrima selected strain was used in a semi-industrial scale combined with three different S. cerevisiae strains, obtaining meaningful results on sensorial parameters like total polyphenol index, color intensity and turbidity of wines and on technological properties as wine filterability.
DETECTION AND TYPING OF LACTOBACILLUS PARABUCHNERI, A POTENTIAL SPOILAGE ORGANISM, IN MILK AND CHEESE

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Background

Histamine can be formed by microorganisms in fermented products such as cheese. High concentrations of this compound can trigger symptoms in consumers who have a histamine intolerance. Microbial decarboxylation of histidine is probably the major cause of histamine formation in cheese. We isolated histamine-producing bacteria from various raw-milk cheeses containing histamine levels higher than 50 mg kg⁻¹. Partial sequence analysis of the tuf gene showed that all these bacteria belong to Lactobacillus parabuchneri which is closely related to Lactobacillus buchneri, a bacterium associated with spoilage in plant-based fermented food and beverages but which is also used in silage fermentation.

Objectives

The aims were to develop a real-time PCR for the specific and quantitative detection of L. parabuchneri in milk and cheese as well as a genotyping methods for strain discrimination and biodiversity estimation.

Methods

We sequenced the genome of L. parabuchneri FAM21731 and twelve other strains.

The draft genome sequence was compared to publicly available genome sequences of L. buchneri and other related species to identify a gene sequence which seems to be unique for L. parabuchneri. Sequences were submitted to Tandem repeats finder and CRISPRFinder to assess hypervariable sequences and their potential for strain discrimination.

Conclusions

Results show that the real-time PCR system is specific for the detection and quantification of L. parabuchneri in dairy products and that the multiplex PCR based on hypervariable sequences can discriminate strains.
The use of these methods is of interest since it can be employed to locate potential sources of *L. parabuchneri* contamination on farms and in dairy plants.
DISCRIMINATION OF VIABILITY STATUS OF SALMONELLA TYPHIMURIUM CELLS INDUCED BY DISTINCT INACTIVATION METHODS USING A COMBINED FLOW CYTOMETRY, MALDI-TOF-MS AND MDA APPROACH

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Background
Food chain contamination with Salmonella Typhimurium is one of the leading causes of foodborne illness worldwide. Rapid and reliable methods for the detection of foodborne pathogens and also discrimination profiles concerning its cells viability status are deeply required and of high importance to assure foodborne pathogens control in the food chain.

Objectives
The goal of this study is to ascertain the Salmonella Typhimurium viability status (viable, viable non-culturable and non-viable) induced by distinct inactivation methods through a combined Flow Cytometry, MALDI-TOF-MS and multivariate data analysis (MDA) approach.

Methods
Different inactivation treatments, such as batch pasteurization (65°C; 30 minutes), sodium hypochlorite (5% or 0.006% commercial bleach), commercial available disinfectant solution (AMUKINA), ethanol (100%) and methanol (100%) were tested in Salmonella Typhimurium LT2. Cell viability assessment was evaluated by CFU’s counts and Flow Citometry (Accuri C6) with LIVE(TO)/DEAD(PI) dyes. In addition, footprint patterns of different viability states were obtained by MALDI-TOF-MS using HCCA as a matrix. Data were analyzed by MDA.

Conclusions
All the inactivation methods tested resulted in a total death of Salmonella Typhimurium LT2 population, with the exception of bleach (0.006%) and pasteurization, the latter used in industrial facilities. These treatments induced cells viable non-culturable state, which is of concern, since under favorable conditions Salmonella Typhimurium could restore viability and eventually its pathogenicity potential. Moreover, we obtained discriminatory peptide/proteins profiles associated with different viability states of Salmonella Typhimurium, which combined with MDA could support the use of this approach to detect microbiological contaminants applicable for quality control purposes.
Background
The refrigerated storage of raw milk at the mill has advantages such as cost savings with the sample collection and reducing the loss of raw material. However, inadequate procedures during production and milk collection are considered the main sources of contamination by spoilage and pathogenic microorganisms, resulting in economic and public health problems.

Objectives
The work aims to enumerate the thermotolerant microbiota found in raw milk and in the milk after pasteurization, correlating it with the shelf life of the product.

Methods
Nine raw milk samples were collected directly from the extension tanks, nearby Castro – PR. After LTLT pasteurization, a portion of raw milk and pasteurized milk were diluted in peptone saline solution in the concentrations: integral, 10^-1, 10^-2 and 10^-3, sown in depth amid PCA inverted and incubated at 35°C for 48 hours. Petrifilm plates ECTM for the enumeration of total coliforms and E. coli were also used. An aliquot was plated in depth for thermoduric counting in the same as described for the days when milk was pasteurized. The milk used for plating thermoduric was also inoculated in Petrifilm ECTM for controlling the progression of coliforms and E. coli.

Conclusions
Four samples showed a reduction in the count of thermoduric microbiota between pasteurization and 25 days of experiment. Two samples obtained less than one log cycle of microbial growth, while three samples showed growth of thermoduric microorganisms, more than one log cycle in the same sampling period.
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FENOLIC ACIDS EFFECTS ON PROBIOTIC BACTERIA GROWTH AND SURVIVAL
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Background
The consumption of probiotics for colonization and maintenance of the intestinal microbiota, regulation of immune mechanisms and decrease of lactose intolerance is strongly credited through data already obtained by investigation. It is also a widespread antioxidative activity of phenolic acids the prevention of diseases related to oxidative stress such as cancer and cardiovascular diseases.

Objectives
Analise de groth behavior and tolerance of Lactobacillus plantarum and Lactobacillus casei when exposed to different fenolic acids.

Methods
We examined the behavior of two probiotic bacteria Lactobacillus plantarum and Lactobacillus casei when exposed to four different concentrations of six different phenolic acids: caffeic, ferulic, p-coumaric, hydroxybenzoic, vanillic and protocatechuich. Over 48h, analyzes were conducted by using a spectrophotometer.

Conclusions
In presence of acids hydroxybenzoic and protocatechuich, the absorbance increased slightly in the deceleration phase at a concentration of 50 mg/L and 100 mg/L, and decreases to a concentration of 200 mg/L compared to the values absorbance of the control. While for L. casei, was observed for all acids tested, the absorbance values at a concentration greater than 50 mg/L, and values approximately equal to a concentration of 200 mg/L, compared to the control absorbance values. For protocatechuich acid, ferulic acid and p-coumaric acid has been possible to verify that the absorbance values were higher than controls at a concentration of 100 mg/L. In all cases, it was found that 200 mg/L showed an effect similar to the control except hydroxybenzoic acid.
Background

NmlR, a transcription regulator in MerR family, is involved in oxidative and nitrosative stress responses in Neisseria gonorrhoeae and Haemophilus influenzae. A homolog is also found in a foodborne pathogen Listeria monocytogenes (lm).

Objectives

In this study, in order to study the role of NmlR\textsubscript{lm} in L. monocytogenes, a null nmlR\textsubscript{lm} strain was constructed. Transcriptomes of 10403S wild type (WT) and its isogenic null nmlR\textsubscript{lm} mutant (MT) strains grown to stationary phase were determined. Phenotypes of these strains in response to different stress conditions were also assessed.

Methods

RNAseq experiments were performed on Ion Torrent platform and the transcriptomes of WT and MT during stationary phase were identified. WT and MT strains were exposed to acid stress and oxidative stress conditions. Role of NmlR\textsubscript{lm} in virulence-associated phenotype was also evaluated in intracellular growth assay using U937 human macrophage-like cell line. Cell sizes of WT and MT were measured using transmission electron microscopy.

Conclusions

RNAseq result reveals that NmlR\textsubscript{lm} negatively regulates 46 genes and positively regulates 28 genes. NmlR\textsubscript{lm}-dependent genes are grouped into 28 operons, eight of which overlap HrcA regulon, another negative regulator in L. monocytogenes. Phenotypic characterization revealed that MT strain survived significantly less than WT under acid stress (pH 2.5) and oxidative stress (3% hydrogen peroxide). Null mutation in nmlR\textsubscript{lm} also resulted in significant decrease in cell length and impaired intracellular growth in U937. Our findings indicate that NmlR\textsubscript{lm} is not only involved in oxidative stress response but also contributes to acid persistence and intracellular growth via either direct regulation or co-regulation with other regulators.
Background

*L. kefiranofaciens* M1, a novel probiotic strain isolated from Taiwanese kefir grains, has been demonstrated to possess anti-allergic, anti-asthmatic, anti-colitis and immunomodulatory effects in our previous in vitro and in vivo studies. These findings support that this strain has the potential to be applied in probiotic products. *L. kefiranofaciens* M1 must survive after processing, production and gastric passage to exert beneficial effects. Thus, the stress tolerance of *L. kefiranofaciens* M1 is important for its survivability.

Objectives

This study was investigated the adaptation and tolerance of *L. kefiranofaciens* M1 to various environmental stresses.

Methods

*L. kefiranofaciens* M1 was adapted to sublethal heat (37°C), cold (25°C), acid (pH 5), bile salts (0.05%), salts (0.1 M), ethanol (4%) and hydrogen peroxide (100 ppm) for 1 h and then challenged with lethal heat (52°C), cold (-20°C), acid (pH 3), bile salts (0.2%), salts (3 M), ethanol (20%) and hydrogen peroxide (1000 ppm), respectively.

Conclusions

The results showed that adaptation to ethanol, hydrogen peroxide, salts and bile salts increased the tolerance of *L. kefiranofaciens* M1 to heat, cold, bile salts and ethanol stresses. Heat adaptation enhanced the tolerance to heat, bile salts and hydrogen peroxide. Acid adaptation caused an increased resistance to cold, acid, bile salts and hydrogen peroxide. Cold adaptation also induced cross-protection against cold, bile salts, salts and ethanol. However, ethanol- and cold-adapted cells decreased their survival after exposure to hydrogen peroxide. The occurrence of stress tolerance response in *L. kefiranofaciens* M1 varied depending on the type and condition of stress.
TRADITIONAL SOUR MILK FERMENTATION USING A POTENTIAL PROBIOTIC LACTOBACILLUS PLANTARUM STRAIN

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Background

Probiotic foods receive market interest as health-promoting, functional foods.

Objectives

To evaluate the performance of a Lactobacillus plantarum strain with probiotic potential as co-starter culture in sour milk fermentation and its ability to give a final product with desirable sensory characteristics.

Methods

UHT milk was inoculated with Streptococcus thermophilus and Lactobacillus delbrueckii subsp. bulgaricus without (control case) or with Lb. plantarum and incubated in appropriate conditions (42°C, 6h). Fermented samples were stored at 4 and 12°C until the end of shelf life of product. Microbiological analysis was performed in parallel with pH and titratable acidity measurements as well as with sensory analysis of the product. The presence of the potential probiotic strain was determined using PFGE.

Conclusions

On first day of storage, the population levels of all microbial groups exceeded 8 log cfu/ml for control and probiotic samples. During storage, the population levels were slightly elevated on probiotic samples compared to the control ones, whereas they were found to be above 8 log cfu/ml for all cases until the end of shelf life. The physicochemical properties (pH, titratable acidity) and the sensory characteristics of the potential probiotic product were similar to those of the control. According to PFGE the Lb. plantarum strain was recovered in high percentages during the products shelf life at both temperatures.

Acknowledgment: This work has been co-financed by the European Regional Development Fund (ERDF) of the EU and by National Resources under the Operational Program Competitiveness and Entrepreneurship (EPAN II), Action "COOPERATION 2011", Project "ProbioDairyMeat".
GREEK FUNCTIONAL YOGURT: ENHANCING THE PRODUCTION PROCESS USING A SELECTED LACTOBACILLUS STRAIN WITH PROBIOTIC POTENTIAL

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Background
Probiotic food products are in general fermented foods, with many studies reporting that the best matrices to deliver probiotics are dairy products, such as fermented milks and yogurt.

Objectives
To evaluate the performance of a Lactobacillus plantarum strain of dairy origin with probiotic potential as co-starter culture in Greek set-type yoghurt fermentation.

Methods
Low-pasteurized milk was inoculated with Streptococcus thermophilus and Lactobacillus delbrueckii subsp. bulgaricus (control) or with the former starters plus Lb. plantarum (probio) and incubated in 42°C for 5h. Subsequently, samples were stored at 4 and 12°C until the end of shelf life of the product. Microbiological analysis was performed in parallel with physicochemical and sensory analysis of the product. The presence of Lb. plantarum was determined using Pulsed Field Gel Electrophoresis (PFGE).

Conclusions
The initial levels of all microbial groups were above 7 log cfu/g for control and probiotic samples. During storage, the population levels of all counts were slightly elevated on probiotic samples compared to the control ones, whereas they exceeded the 7 log cfu/g at all cases. Probiotic yoghurt produced with Lb. plantarum strain exhibited a better sensory profile, with a rich traditional taste, but with similar physicochemical properties to the control. According to PFGE, the Lb. plantarum strain was recovered in high percentages during the products shelf life at both temperatures.

Acknowledgment: This work has been co-financed by the European Regional Development Fund (ERDF) of the EU and by National Resources under the Operational Program Competitiveness and Entrepreneurship (EPAN II), Action "COOPERATION 2011", Project "ProbioDairyMeat".
Background

The research of novel formulations with newly selected probiotic strains is important to satisfy the increasing request of the market and to obtain functional products with health-conferring properties. The most studied probiotic bacterial strains belong to the genus *Lactobacillus*, an important group of Lactic Acid Bacteria with functional and technological properties. Antagonistic activity toward potentially pathogenic microorganisms and adhesion to gut tissue are some of the main requirements when selecting successful probiotic strains.

Objectives

The aim of this work was to evaluate the antibacterial and anti-adhesion activity of *Lactobacillus plantarum* strains isolated from ewe’s raw milk and their ability to *in vitro* colonize the intestinal epithelium. These strains were previously selected as potential probiotics because of their ability to survive at low pH values and physiological bile concentrations.

Methods

For detection of antagonistic activity, agar spot test and a well diffusion assay were performed toward *Escherichia coli* O157:H7, *Salmonella* Typhimurium and *Listeria monocytogenes*. The *in vitro* anti-adhesion effect was measured by a competitive inhibition assay on Caco-2 cells.

Conclusions

The results showed our strains having an excellent antibacterial activity against foodborne bacterial pathogens and anti-adhesion effect toward *E. coli* O157:H7 and *L. monocytogenes*, under the condition tested. Therefore, they showed interesting
probiotic characteristics. An *in vivo* study would be useful for a deeper evaluation of their properties, and their potential use in the development of novel functional food. This work has been supported by Regione Autonoma Sardegna (RAS, L.R. 7/2007).
COOPERATION BETWEEN SPECIES IN THE YOGURT CONSORTIUM IS AFFECTED BY THE MODULATION OF INTRACELLULAR PH THAT IS DRIVEN BY STREPTOCOCCUS THERMOPHILUS UREASE ACTIVITY

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Background

The proto-cooperation between Streptococcus thermophilus and Lactobacillus delbrueckii subsp. bulgaricus in the yogurt consortium enhances the growth rate and size of each population. In contrast, the independent growth of the two species in milk leads to a slower growth rate and a smaller population size.

Objectives

In this study, we evaluated how urease activity of S. thermophilus might affect the intracellular pH of L. delbrueckii, modulating its bioenergetics during milk fermentation.

Methods

Urease-mediated intracellular alkalization of S. thermophilus and L. delbrueckii was evaluated by flow cytometry using cFSE as pH-dependent fluorescent probe. Lactose consumption and lactic acid production was followed in vivo by ¹³C-NMR analysis. D-L lactic acid production in milk was measured enzymatically. For the evaluation of the pH-dependent glycolysis efficiency, glucose and lactic acid were measured by HPLC.

Conclusions

We observed that intracellular alkalization caused by urea hydrolysis or the addition of ammonia to milk boosted lactic acid production in S. thermophilus and in L. delbrueckii when the species were grown separately or in combination. Therefore, we propose that urease activity acts as an altruistic cooperative trait, which is costly for urease-positive individuals but provides a local benefit because other individuals can take advantage of urease-modulated pH. Compared to the interactions that are known to occur between S. thermophilus and L. delbrueckii in the yogurt consortium, the modulation of the bioenergetic efficiency due to the intracellular pH alkalization represents a new type of cooperation that directly affects the kinetic parameters of enzymes involved in homolactic fermentation in both the species.
Background

*Escherichia coli* (*E. coli*) is commonly used as a surrogate for pathogens in research to identify sources of agricultural contamination and to characterize how pathogens persist on plant surfaces. However, *E. coli* strains are highly diverse, exhibiting differences in physical, chemical and biological properties which contribute to fitness in secondary habitats.

Objectives

An important step in selecting *E. coli* for use as pathogen surrogates is to identify strains with characteristics (including adhesion, motility and biofilm formation) that contribute to their occurrence and survival in a manner similar to that of pathogens.

Methods

To this end, strain-level differences in genotype (adhesions (*iha, agn43, eaeA* and *timH, kpsMTII*) and siderophores (*iroNe.coli, chuA*)) and phenotype (biofilm formation, curli expression, and growth rate) were evaluated for isolates from livestock (swine, poultry, dairy) and water sources. Selected *E.coli* isolates (n=18 of 1,300), a common *E. coli* control strain (ATCC 25922), and *Salmonella* (ATCC 13311) were used in soil studies to evaluate adhesion to plastic, sandy loam or clay soils.

Conclusions

Average adhesion of environmental isolates to plastic was two-fold higher than that of the control strain. *Salmonella* and one of the environmental isolates had adhesion rates seven-fold higher than the control strain. That isolate had extremely high adhesion to clay (91.3 ± 2.7%) and sandy loam (91.7 ± 0.1%) soils, but had none of the genes commonly associated with adherence or biofilm formation. These results suggest that *E. coli* strain choice should be an important consideration in its use as a pathogen surrogate.
BACTERIAL TRANSCRIPTIONAL ADAPTATION TO NICHES: MOLECULAR AND METABOLIC RESPONSES OF LACTOBACILLUS PLANTARUM C2 IN PLANT SUBSTRATES

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Background

Raw fruits and some vegetables possess intrinsic chemical and physical features that make them particularly hostile environments for bacteria. To cope with environmental conditions, microorganisms may adopt sophisticated adaptation mechanisms. The diversity of plant environments and of bacterial enzyme activities makes the microbial adaptation to plant niches markedly heterogeneous. \textit{Lactobacillus plantarum} is a highly heterogeneous and versatile lactic acid bacteria frequently found or used in vegetables and fruits fermentation.

Objectives

The aim of the study was to determine biological relevance of differentially expressed genes in \textit{L. plantarum} C2 during fermentation of plant substrates, aiming to provide new insights into transcriptional response and niche adaptation.

Methods

\textit{L. plantarum} C2 was grown and stored in carrot or pineapple juices to mimic the chemical composition of the respective raw matrices. MRS broth was used as the control medium for optimal growth. Whole-transcriptome analysis based on customized microarray profiles has been used to determine altered transcription patterns in \textit{L. plantarum} C2. These were compared with substrate utilization data gathered from high throughput phenotypic microarrays.

Conclusions

Plant substrates exerted a transcriptional pressure and induced specific molecular and metabolic responses in \textit{L. plantarum} C2. RNA and phenotypic microarray analyses revealed altered transcription patterns of genes encoding functions involved in primary metabolism, membrane transport, cofactors and vitamins metabolism, translation regulation, nucleotide metabolism, and fatty acid biosynthesis. Findings contribute to the description of bacterial transcriptional adaptation to niches, and provide a more solid basis for selection the most suitable starters for fermentation of targeted matrices.
IN VITRO ASSESSMENT OF ANTIBACTERIAL ACTIVITY OF RESINS FROM SOME WOODY PLANT BUDS AND SYNERGISTIC EFFECT BETWEEN STANDARD PHENOLIC COMPOUNDS

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Background
With emerging of new and reemerging of old infectious diseases, urgent need for new antimicrobial agents arises. Resins are complex mixtures of phenolic and isoprenoid compounds secreted by plants to provide protection against predators and pathogenic microorganisms.

Objectives
The present study was aimed at investigating the in vitro antimicrobial activity of resins from some woody plants on the selected bacterial strains. To elucidate origin of antimicrobial activity, potential synergistic effect between standard phenolic compounds was investigated, also.

Methods
The samples of resins were collected from different regions in Serbia and dissolved in methanol. Well diffusion and broth microdilution methods were implemented for assessing antimicrobial activity.

Conclusions
Gram-positive bacteria exhibited strong susceptibility to samples from Populus sp. while being resistant on samples from other origin. The most sensitive strain was B. subtilis, towards samples collected from Populus nigra and Prunus avium (MIC was 0.05 mg/ml). The strains of S. aureus, MRSA and L. monocytogenes showed similar sensitivity to poplar samples with MIC values mostly under 1 mg/ml. L. monocytogenes was also susceptible to cherry buds samples. Sample U19 from white poplar had significant antimicrobial activity on all tested strains, with lowest detected MIC values. Synergistic effect of gallic acid, quercetin and caffeic acid on B. subtilis; gallic, caffeic, p-coumaric acid and naringenin on E. faecalis; and gallic, caffeic acid and naringenin on S. flexneri was observed. In conclusion, antimicrobial activity of resins is probably outcome of action of phenolic compounds.
A COMPARISON OF LISTERIA SPP. PREVALENCE BETWEEN THE RAW FRESHWATER FISH FROM LAKE AND RETAIL MARKET IN LATVIA

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Background
Listeriosis is a foodborne infection caused by *Listeria monocytogenes*, which occurs due to ingestion of pathogen containing foodstuffs. *Listeria* spp. and *L. monocytogenes* are widespread in environment, but raw and processed fish products were found to be frequently contaminated with *Listeria* spp. Therefore fish are suspected to be a possible source for *Listeria* spp. including *L. monocytogenes* transmission to final food products.

Objectives

The objectives of this study were to determine and compare the prevalence of *Listeria* spp. in raw freshwater fish from retail market and fish captured from lake in Latvia.

Methods

A total of 51 raw freshwater fish samples were collected from lake (n = 11) and retail market (n = 39) in Latvia. A total amount of 25 g of fish skin, gut and muscle tissues of each fish was used for testing. Testing was performed in accordance to the International Organization for Standardization method (ISO 11290-1: 1996).

Conclusions

The fish samples from lake were found to be *Listeria* spp. negative. In contrast, 69% of retail fish were found to be contaminated with *Listeria* spp. There were no significant differences between the prevalence of *L. innocua* and *L. monocytogenes* (p>0.05), while the prevalence of *L. ivanovii* and *L. welshimeri* was significantly lower than the prevalence of *L. monocytogenes* and *L. innocua* (p<0.05). The results
indicate that fish contamination with *Listeria* spp. may occur during fish handling, and the high prevalence of *L. monocytogenes* in raw fish at retail level can pose a threat for public health.
GENOTIPIC DIVERSITY AND ANTIMICROBIAL RESISTANCE OF SALMONELLA TYPHIMURIUM STRAINS ISOLATED HUMANS AND FOOD BETWEEN 1983-2013

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Background

The foodborne disease caused by Salmonella Typhimurium is a major health problem worldwide.

Objectives

The aims of this study were to genotype S. Typhimurium isolated in Brazil and to verify its antimicrobial resistance profiles.

Methods

A total of 92 S. Typhimurium strains, isolated from humans (43) and food (49), between 1983 and 2013 in Brazil, were typed by PFGE, ERIC-PCR and MLVA. Additionally, their antimicrobial resistance was evaluated. Results: The 92 S. Typhimurium strains were grouped in two clusters, by PFGE, PFGE-A and PFGE-B (subdivided in PFGE-B1 and PFGE-B2); by MLVA in two clusters, MLVA-A and MLVA-B (subdivided in MLVA-B1 and MLVA-B2). By ERIC-PCR, in three clusters, ERIC-A, ERIC-B and ERIC-C. The strains isolated from humans before the mid-1990s were allocated in the PFGE-A, PFGE-B1, PFGE-B2, MLVA-A, MLVA-B1, MLVA-B2, ERIC-A and ERIC-B. The strains isolated from humans after mid-1990s were distributed in the PFGE-B1, MLVA-B1, MLVA-B2 and ERIC-A. The strains isolated from food were distributed in the PFGE-A, PFGE-B1, MLVA-A, MLVA-B1, MLVA-B2, ERIC-A, ERIC-B and ERIC-C. Twenty-three (25%) strains were multidrug resistance.

Conclusions

The results suggest that the studied strains isolated from humans before the mid-1990s were genetically more diverse, which might indicate that selection of a more adapted S. Typhimurium subtype occurred after S. Enteritidis became the most prevalent serovar in Brazil. Regarding, the food strains the results suggest the current circulation of more than one subtype. The occurrence of multi-drug resistant strains isolated from food is an alert for the possible risk for human consumption.
MOLECULAR CHARACTERIZATION AND RESISTANCE PROFILE OF CAMPYLOBACTER COLI STRAINS ISOLATED FROM DIFFERENT SOURCES IN BRAZIL

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Background
Campylobacter coli is an important causative agent of human diarrheal diseases worldwide. However, in Brazil is not frequently studied.

Objectives
This study aimed to genotype and to investigate the presence of virulence genes and the resistance profile in C. coli strains.

Methods
A total of 58 C. coli strains isolated from humans(10), animals(14), the environment(18) and food(9), between 1995-2011 in Brazil, were genotyped by PFGE. The presence of 16 virulence genes was searched by PCR. The resistance profile was obtained by MIC for erytromycin, ciprofloxacin, tetracycline and doxacycline. All the strains presented the genes flaA, cadF and sodB. The cdtB, flhA, dnaJ and pldA genes were observed in 20, 15, 10 and 6 strains, respectively. The ciaB, iamA, cdtA, cdtC, docA, virB11, wlaN, racR and crsA genes were not detected. PFGE grouped the strains in two main clusters with more than 45.4% of similarity. PFGE-A and PFGE-B clusters comprised 42 and 16 strains, respectively. In both clusters, strains of clinical and non-clinical sources were grouped in subclusters with a similarity value of more than 80%. The resistance profile showed that 19 strains, mainly of non-clinical sources, were resistant to at least one antimicrobial agent.

Conclusions
The PFGE results confirmed the heterogeneity of the C. coli strains studied. However, the high similarity (>80%) among some strains of different origins suggests a possible contamination between clinical and non-clinical sources in Brazil. Because of these, the existence of resistant strains becomes a concern. The presence of important virulence genes indicates the pathogenic potential of those strains.
Background

Staphylococcus spp. are common members of the normal human flora. However, some Staphylococcus strains are recognized as human pathogens, in particular due to the production of several virulence factors and enterotoxins particularly relevant in food intoxications.

Objectives

Since many of the food intoxications by Staphylococcus aureus are typically associated with cross-contamination, detection of Staphylococcus aureus was performed on food handlers.

Methods

Hand swabs from 167 food handlers were analysed for the presence of Staphylococcus aureus using Baird Parker agar and coagulase test, and genotypically confirmed by RT-PCR. A total of 26 strains were analysed using RT-PCR for the presence of virulence and enterotoxin genes, namely, sea, seb, sec, sed, seg, sei, tsst1 and pvl. The same strains were phenotypically characterized in terms of antibiotic resistance using the disk diffusion method and antimicrobial agents from 15 different categories.

Conclusions

More than 11% of the samples were positive for Staphylococcus aureus. A low prevalence of antibiotic resistant strains was found, with more than 55% of the strains being sensitive to all of the antimicrobial agents tested. However, a high prevalence of resistance to macrolides was found, with 42% of the strains showing resistance to erythromycin. At least one of the virulence or enterotoxin genes was detected in 54% of the strains and the seg gene was detected in 42% of the strains.
Potential genetic targets for controlling Campylobacter jejuni

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Background

Essential genes have been suggested to provide useful information for the development of new specific anti-infective agents in C. jejuni, as the major cause of bacterial gastroenteritis in humans worldwide (Stahl & Stintzi, 2011). To determine genetic regions that are present or highly prevalent in all C. jejuni strains could be also of interest as potential genetic targets for this purpose.

Objectives

The aim of this study was to identify by microarray comparative genomic hybridization (MCGH) potential genetic targets for controlling C. jejuni.

Methods

The prevalence of 1652 genes among 120 geographically diverse isolates obtained from different sources (human, chicken, cattle, water and wild birds) was determined by MCGH using the BµG@S CJv3.0.0 microarray. As MCGH does not confirm the absence or divergence of the tested genes, the results were compared with those obtained by both MCGH and genomic sequencing in previous studies (Stabler et al., 2013; van Tonder et al., 2014).

Conclusions

MCGH analysis detected 1366 genes (83%) with prevalence 90% in the C. jejuni population studied. These genes included 816 genes (60%) recently proposed as core (van Tonder et al., 2014), and 229 genes (28%) proposed as essential gene candidates (Stahl & Stintzi, 2011; Metris et al., 2011). A final list of 22 genes, including core and essential genes, was highlighted. All these genes encode products required for cell growth, and are therefore potential targets to consider for antimicrobial intervention strategies in C. jejuni.
Background

Osmotic dehydration is a water removal process, based on soaking food (fruit, vegetable, meat and fish) in a hypertonic solution. Recent research has shown that use of sugar beet molasses as a hypertonic solution improves osmotic dehydration process from technological, nutritional and microbiological aspect.

Objectives

Effect of osmotic dehydration process on microbiological profile of chicken meat was investigated in order to determine the usefulness of this technique as pre-treatment for further treatment of chicken meat.

Methods

Fresh breast chicken meat was cut into cubes of 1x1x1 cm and contaminated with *Esherichia coli*, *Listeria monocytogenes* and *Salmonella* spp in laboratory conditions prior the process of osmotic dehydration. The process was conducted during 5 hours at 44°C in sugar beet molasses as an osmotic medium. All microorganisms were tested according to ISO methodology: *Salmonella* spp EN ISO 6579:2002, *Listeria monocytogenes* EN ISO 11290-2:1998, *Escherichia coli* ISO 16649-2:2001, *Enterobacteriaceae* ISO 21528-2:2004 and total number of microorganisms EN ISO 4833-1:2013.

Conclusions

The results of the microbiological profile of fresh, contaminated and osmotically dehydrated chicken meat have shown that the numbers of all examined microorganisms have significantly statistically reduced after the process.

From presented results it can be concluded that process of osmotic dehydration in molasses significantly improves microbiological profile of treated chicken meat and it is suitable pretreatment for further chicken meat processing, providing microbiologically safe intermediate product.
USE OF COMPARATIVE GENOMICS AND THIN LAYER CHROMATOGRAPHY TO IDENTIFY SECONDARY METABOLITES RESPONSIBLE FOR ANTI-GRAM NEGATIVE ACTIVITY IN BACILLUS AMYLOLIQUEFACIENS SUBSP. PLANTARUM

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Background

Strong consumer demand for safe and high-quality products has continued to drive food and pet food producers to seek novel solutions targeting Gram-negative bacterial contamination, such as Escherichia coli and Salmonella spp.

Objectives

Since Bacillus spp. are known for their ability to produce antimicrobial compounds, culture supernatants of Bacillus amyloliquefaciens strains were evaluated and characterized for their antimicrobial properties. Furthermore, genome sequences were to be generated to provide insights to the mode of action.

Methods

Six B. amyloliquefaciens strains were used for the experiments. Titration of activity was conducted by measuring the halos in an agar well diffusion assay, employing an extensive range of bacteria and fungi. The effect of pH, heat and enzymes on the activity of the supernatants was expressed as a percentage of residual activity compared to non-treated samples. Draft genomes were generated by 454 FLX Titanium for each of the Bacillus strains and mined for the presence of secondary metabolites, which was furthermore confirmed with TLC-bioautography profiles of methanolic extracts.

Conclusions

All supernatants exhibited activity against important food related bacteria and fungi, including Gram negatives. Generally, considerable activity was retained after heating and at alkaline pH but not under acidic conditions. Whereas different enzymes had a variable effect on the antimicrobial activity, genomic data indicated that the Bacillus strains harboured gene clusters encoding proteins involved in the
production of an array of secondary metabolites, including several non-ribosomal peptide synthetases, polyketides, and ribosomally synthesized peptides, which could be responsible for the *in vitro* antimicrobial properties observed.
Background
According to the Register of Infectious Diseases, supplied by the National Public Health Institute (THL), in 2013 Campylobacter sp. was the leading cause of human bacterial infection in Finland with 4059 cases registered (incidence 76 cases/100000 inhabitants). However, the real number of cases is assumed to be considerably higher than the registered cases numbers. The prevalence of Campylobacter sp. in animal-derived foods is affected by its prevalence in the production farms and in the animals. The contamination of the final product is also dependent on procedures carried out in abattoirs and during the various stages of the food production chain. The handling of food contaminated with Campylobacter sp. in domestic or industrial kitchens may also lead to cross-contamination, either directly from raw meat to products that will not undergo further cooking or indirectly via work surfaces, hands or utensils.

Objectives
Estimating the prevalence and concentration of Campylobacter in fresh (chicken, turkey) meat (sampled at retail), as well as presenting a novel Bayesian model to assess the probability of infection and illness.

Methods
Bayesian modeling of prevalence and concentration using Openbugs software based on results from detection and quantification of campylobacters (chicken: 608 samples from 226 batches) and (turkey: 558 samples from 185 batches) using modified NMKL 119:2007 and ISO 10272-2:2006.

Conclusions
Risk of Campylobacter exposure from Finnish fresh poultry meat is mainly due to undercooking or cross-contamination especially during summertime. The model can be used to assess e.g. relative risks, with quantification of uncertainty based on the data.
EFFECT OF ELECTRON BEAM IRRADIATION IN MINAS FRESCAL CHEESE ARTIFICIALLY CONTAMINATED WITH ENTEROHAEMORRHAGIC ESCHERICHIA COLI (EHEC) O157:H7

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Background
The enterohemorrhagic Escherichia coli (EHEC) O157:H7 has been known to cause food-borne outbreaks. Cheeses are among the main food products incriminated in these outbreaks. Characterized as a perishable food, the Minas frescal cheese (MFC) has potential risk for O157:H7 transmission. As a result, the development of new technologies for its conservation has been studied. The use of irradiation has proved to be one of the most convenient options.

Objectives
The objective of the present study was to evaluate the effect of electron beam irradiation (doses of 1.0, 1.5 and 2.0 kGy) in MFC artificially contaminated with O157:H7.

Methods
Samples were produced to microbiological analyzes, one non-contaminated and non-irradiated, one contaminated and non-irradiated and three contaminated and irradiated. And to thiobarbituric acid reactive substances concentration (TBARS) analysis, one non-irradiated and non-contaminated and three non-contaminated and irradiated.

Conclusions
Artificially contaminated and irradiated MFC samples presented a shelf life of ten storage days at 4°C. The dose of 1.5 kGy reduced 2 log the number of O157:H7 and the doses of 1.5 and 2.0 kGy eliminated O157:H7 from the artificially contaminated samples. The TBARS assay was conducted to evaluate if the effect of irradiation could influence in the stability of the lipid fraction present in the cheese. All irradiated cheeses showed TBARS values, over the 40 day storage period, lower than in the non-irradiated sample. Results presented here demonstrate electron beam irradiation of MFC artificially contaminated with O157:H7 is a viable solution and a promising alternative to ensure the safety and quality of this food product.
BEHAVIOR OF SHIGA TOXIN-PRODUCING ESCHERICHIA COLI (STEC) O157:H7 AND O113:H21 UNDER THE EFFECT OF DIFFERENT LEVELS OF PH AND TEMPERATURE
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Background
Shiga toxin-producing Escherichia coli (STEC) cause hemorrhagic colitis, hemolytic uremic syndrome and bloody diarrhea.

Objectives
This work aimed to evaluate the effect of two individual factors (pH and temperature), and their interaction, on the multiplication and survival of STEC (O157:H7 and O113:H21) using statistics as a tool for development of the experimental design and the analysis of results.

Methods
Approximately 10⁸ CFU per ml were inoculated into Trypticase Soy Broth at pH conditions (4.0, 4.5, 5.5, 6.5 and 7.0) and temperature (6, 10, 20, 30 and 35°C). The experimental design was prepared by a central composite design, generated by the software Statistica 7 (StatSoft, OK, USA), and given 11 trials including three central points and four axial points. Assays were performed in 22 h.

Conclusions
Were generated equations that model the bacterial behavior from a multiple linear regression model. In the study with the RJ 581 strain (O157:H7), variables, individually and combined, showed linear significant effect on the response, the effect of the variable pH being the most relevant, with R² equal to 0.94. Explaining the optimal conditions for the multiplication of the RJ 581 strain values are closer to pH 7.0 and 35°C. In the study with the RJ 702 strain (O113:H21), we observed that the independent and combined variables had no significant effect on the response. The R² of the model was 0.4, showing a low suitability of experimental design to RJ 702 strain.
LACTIC ACID BACTERIA AS ANTAGONIST AGENTS TO FOODBORNE PATHOGENS ON FRESH-CUT ‘GOLDEN DELICIOUS’ APPLES

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Background

Food safety is a major concern in fresh-cut industry. Fresh-cut produce can become contaminated with foodborne pathogens during manipulation. Application of natural antimicrobial substances provides new opportunities for the control of pathogenic bacteria. Lactic acid bacteria (LAB) are known to have antagonistic effect against sensitive bacteria species/strains by the secretion of substances with antimicrobial activity.

Objectives

The main objective of this work was to do a screening of 37 LAB isolates for antimicrobial activity on minimally processed ‘Golden delicious’ apples against the foodborne pathogens *Escherichia coli* O157:H7 NCTC 12900, *Salmonella enterica* subsp. *enterica* Michigan ATCC BAA-709, *Cronobacter sakazakii* ATCC 51329 and *Staphylococcus aureus* ATCC 25923.

Methods

LAB studied were isolated from fresh-cut fruit purchased in the market. One g of fresh-cut apple was inoculated with the LAB followed by the inoculation of the pathogen at a concentration of $10^8$ cfu/mL. Samples were incubated at 30 °C for 48 h and after the recovery of the pathogens were made on selective/differential mediums. Fresh-cut apples inoculated with only the pathogen were used as control.

Conclusions

From the 37 LAB isolates 18 were capable to reduce all the pathogens more than 2 log cfu/g and 5 achieved a reduction of more than 3 log cfu/g for all the pathogens. *S. enterica* and *C. sakazakii* showed reduction with all the isolates studied and *E. coli* and *S. aureus* showed reduction with 36 of the 37 isolates. Results obtained support the potential use of LAB as biocontrol agents against foodborne pathogens in fresh-cut fruit.
FERMENTATION PROPERTIES AND POTENTIAL PREBIOTIC ACTIVITY OF A HIGH PURITY GOS ON IN VITRO GUT MICROBIOTA PARAMETERS IN HEALTHY INDIVIDUALS

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Background

Most in vitro and in vivo studies involving prebiotic oligosaccharides have been carried out using inulin and its fructooligosaccharide (IFS) derivatives, as well as various galactooligosaccharides (GOS). It has been shown that these food ingredients have the ability to improve selectively the growth of bifidobacteria and, consequently, lead to important changes in the gut microbiota composition that may confer health benefits to the host. Bimuno®, (Clasado Ltd., Buckinghamshire, UK) is a 50% mixture containing galactooligosaccharides (B-GOS) having multiple biological health activities within the colonic environment, beyond the stimulation of bifidobacteria and lactobacilli at genus level and it has been tested in different in vitro and in vivo studies.

Objectives

The present study aimed to determine the in vitro potential of a new high GOS content version of Bimuno (65% GOS content), comparing to B-GOS used as positive control, in a pH and volume controlled dose response batch culture experiments. Three different doses of the high purity GOS (2.75, 1.38 and 0.92 g) were tested.

Methods

Changes in the gut microbiota during a time course were identified by fluorescence in situ hybridisation (FISH), whilst the small-molecular weight metabolomic profiles and short chain fatty acids (SCFAs) were determined by NMR analysis and Gas Chromatography (GC), respectively.

Conclusions

The High purity GOS showed positive modulation of the microbiota composition after 4h fermentation at the lowest dose compared to B-GOS. Moreover, the administration of the specific GOS induced a significant increase in acetate as the major SCFAs synthesized compared to propionate and butyrate concentrations.
COMMON AND DIVERGENT FEATURES OF THE HEAT-RESISTANT SPOILAGE PROTEASE PRODUCING MICROBIOTA IN MILK FROM TWO DIFFERENT UHT-MILK PRODUCING COUNTRIES, BRAZIL AND BELGIUM

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Background

The majority of the bacteria from raw milk are inactivated by heat treatments commonly used in dairy industry (pasteurization or ultra-high temperature (UHT)). However, these treatments do not destroy heat-resistant spoilage enzymes such as proteases produced by certain psychrotrophic bacteria before processing.

Objectives

The first objective was to identify the main proteolytic micro-organisms in raw milk from Belgium and Brazil, two countries with a large UHT-milk production but under different primary production conditions which can influence the microbiota composition of raw milk. Secondly, the diversity of the protease enzymes was investigated in order to characterize the target of future spoilage detection assays.

Methods

Raw milk samples were stored at simulated conditions of farm storage and transport according to Brazilian and Belgian regulations, respectively. The highly proteolytic strains were polyphasically identified. Their protease enzymes were characterized for heat resistance, proteolytic activity in milk, casein zymogram analysis with MALDI-TOF identification of protein bands, and for gene sequence diversity.

Conclusions

The Pseudomonas fluorescens group together with Ps. fragi and Ps. lundensis were dominant in Belgian raw milk, while in Brazil, almost the same Pseudomonas spp. together with Serratia liquefaciens were dominant, the latter being an indication of less hygienic conditions during milking. Belgian isolates showed overall a significant higher proteolytic activity then most Brazilian isolates. The proteolytic activity from both sources was resistant to UHT conditions with the one from S. liquefaciens being the most heat resistant. It seems that the spoilage protease from S. liquefaciens is
unrelated to the heterogeneous *Pseudomonas AprX* protease.
POLYPHASIC IDENTIFICATION OF THE DOMINANT MICROBIOTA OF PEELED BROWN SHRIMP (CRANGON CRANGON) STORED UNDER MODIFIED ATMOSPHERE PACKAGING AT 4 °C WITHOUT PRESERVATIVES

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Background

Brown shrimp are, like many seafood products, prone to rapid spoilage. The main factor limiting shelf life is microbiological activity. Specifically for brown shrimp without preservatives the microbiota is dominated by the genera *Pseudoalteromonas* and *Psychrobacter* when stored under iced aerobic conditions, but the spoilage bacteria of shrimp when cool stored under modified atmosphere (MAP) conditions without preservatives, a product which is gaining popularity among consumers, are still unknown.

Objectives

The main objective was to evaluate the variation throughout the year and during storage in the microbial community of MAP packaged cooked and machine peeled shrimp (*Crangon crangon*) without preservatives and the description of the dominant spoilage microbiota after 7 days of shelf life at 4 °C.

Methods

Three series of shelf life experiments resulting from three sampling periods of cooked shrimps throughout the year were conducted with mechanically peeled shrimp stored in a modified atmosphere (40% CO₂ and 60% N₂). Culture-dependent and – independent methods, notably (GTG)₅ clustering followed by 16S rRNA gene sequencing of isolates and 16S rRNA gene based DGGE in combination with metagenomics, were used to monitor the bacterial population in brown shrimp without preservatives.

Conclusions

According to culture dependent identification the microbiota of brown shrimp was dominated by *Arthrobacter bergerei*, *Shewanella putrefaciens*, *Aliivibrio* spp., *Psychrobacter* spp., *Brochothrix thermosphacta* and *Vagococcus salmoninarum*. Using the culture independent approach the dominating genus after 7 days of storage was *Carnobacterium*. Most of the micro-organisms identified through the complementary approach are known to contribute to spoilage, but for some (e.g. *V. 
*salmoninarum* the spoilage potential is yet unknown.
DNAJ GENE AS A MOLECULAR DISCRIMINATOR WITHIN THE LACTOBACILLUS CASEI GROUP
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Background
The current taxonomy of the Lactobacillus casei group is comprises three closely related species: L. casei, L. paracasei and L. rhamnosus. Some strains of these species have probiotic features and are now widely used in the food industries. However, neither phenotypic nor the most frequently applied genotypic marker (16S ribosomal DNA) provides sufficient resolution for accurate identification of L. casei group.

Objectives
The aim of this study was to use the dnaJ gene as a molecular marker for species-level discrimination and identification within the L. casei group.

Methods
The degenerate primers (LcasDnaj-F1/R1) were designed for dnaJ gene amplification and sequencing. The relationships between the members of L. casei group were conducted based on sequence similarity and phylogenetic analysis. The species-specific primers were designed by analysing the highly variable regions of the dnaJ gene.

Conclusions
Within the L. casei group, the nucleotide sequence similarity of dnaJ gene was significantly lower than that of 16S rRNA gene, and all examined strains could be clearly distinguished. In addition, the species-specific primers were developed, which were then employed for two-plex minisequencing analysis, and were shown to be specific for L. paracasei and L. rhamnosus. Our data indicate that the phylogenetic relationships in the L. casei group can be resolved using dnaJ gene sequencing, and the species of L. paracasei and L. rhamnosus can be identified using novel species-specific minisequencing assay.
INFLUENCE OF pH ON GENE EXPRESSION IN LACTOBACILLUS PARABUCHNERI FAM21731

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Background

The formation of biogenic amines in cheese can occur as a result of the presence of bacteria possessing amino acid decarboxylase activity. One of the most important biogenic amines in cheese is histamine, since it can cause serious human health problems upon ingestion.

Objectives

To better understand the factors contributing to histamine accumulation, we isolated histidine decarboxylase positive lactic acid bacteria from various raw-milk cheeses with elevated histamine levels. All isolates belonged to the species Lactobacillus parabuchneri. Various factors, such as starter culture, proteolysis during ripening, storage temperature, salt concentration and pH can affect histamine formation in cheese. For cheeses made from raw milk, a thermal treatment of the milk prior to cheese making to reduce the numbers of histamine producing L. parabuchneri is not an option. Other strategies could be developed to prevent accumulation of biogenic amines if the metabolism of these bacteria is known in more detail.

Methods

RNA from L. parabuchneri FAM21731 grown at various pH was isolated and sequenced on an Ion Torrent PGM sequencer.

Conclusions

Whole transcriptome analysis of L. parabuchneri was established using next generation sequencing technology. The methodology helps us to better understand how cheese making technology and choice of starter culture influence the metabolism of histamine producing L. parabuchneri strains.
Background

Proteolysis and amino acid catabolism are the most important events taking place during cheese ripening. Microorganisms play a crucial role in these biochemical changes. Among various lactic acid bacteria *Pediococcus acidilactici* is often found in cheese at the end of ripening. Little is known if this species contributes to protein break down and flavor development in cheese. When *Pediococcus acidilactici* FAM18098 was used as adjunct culture in cheese making, we observed that it degraded arginine, serine and threonine and synthesized ornithine, alanine and 2-aminobutyrate during cheese ripening.

Objectives

*P. acidilactici* FAM18098 did not degrade serine and threonine under laboratory conditions using MRS broth, a broth usually used for lactic acid bacteria in the laboratory. By changing the composition of media, we found a medium in which this strain showed an amino acid metabolism similar to the one observed in cheese. By analyzing the transcriptome of *P. acidilactici* FAM18098 we want to better understand the transcriptional regulation of genes involved in amino acid metabolism and to find experimental conditions that are similar to the ones of cheese.

Methods

RNA from *P. acidilactici* FAM18098 grown in various media was isolated and sequenced on an Ion Torrent PGM sequencer.

Conclusions

We established a protocol to analyze and compare whole transcriptomes of *P. acidilactici* with next generation sequencing technology. With this methodology we studied genes that are involved in amino acid metabolism under conditions that probably mimic a cheese environment.
TRANSFER OF L. MONOCYTOGENES FROM DIFFERENT FOOD CONTACT SURFACES TO CHEESES

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Background

The food contact surfaces are subject to contamination by pathogens that could lead to cross-contamination by transfer events to other food products. However, the European regulation n° 1935/2004 of 27th October 2004 (Anonymous, 2004) specifies that materials intended for safe food contact must not interfere with foodstuff characteristics.

As a traditional and natural material, wooden boards are traditionally used as a “technological tool” during cheeses ripening process. In France, wood is authorized for food contact by the French Arrêté November 1945 (Anonymous, 1945) and the information note of DGCCRF 2012-93(Anonymous, 2012).

Objectives

The aim of this study was to determine the behavior of wooden surfaces in direct food contact, contaminated by a well-known risk along the production chain of dairy products: Listeria monocytogenes.

Methods

For this purpose, a protocol was defined and new spruce boards were inoculated by L. monocytogenes solution at a concentration of $10^5$ CFU/cm² and the microbial transfer to pressed non-cooked cheeses was studied. Factors such as cheese time contact, wood moisture content and age of the cheeses were tested.

A comparison of transfer quantification, with other materials surfaces: glass plates and plastic sheets with inclined meshes, used in cheese production, was also realized among the same conditions.

Conclusions

The results showed for all tested surfaces a transfer yield below 3% (CFU/cm²) in the first hour. No differences were found for older cheeses or for higher wood moisture
content. In conclusion, wooden shelves are, as much as, safety for food contact than plastic or glass surfaces.
Background

Manipulation or consumption of raw or poorly cooked foods of animal origin is an important source of human infection with the emerging food and waterborne pathogen *Arcobacter butzleri*. The ability of this bacterium to form biofilm favors the transmission and its adherence to food and food-contact surfaces possibly favors cross-contamination. Preventing cross-contamination is a key factor in Food Safety. Besides, studying the virulence properties of any potentially pathogenic foodborne bacteria is essential for the consumer safety.

Objectives

The aim of this study was to determine the biofilm formation ability of 42 *A. butzleri* isolates obtained from different food products and to study the prevalence of ten putative virulence genes among adherent isolates.

Methods

Biofilm formation was assayed by microtiter plate method (Teh *et al.*, 2010) and categorized according to Naves *et al.* (2008). Putative virulence genes were studied by PCR (Douidah *et al.*, 2012; Karadas *et al.*, 2013).

Conclusions

Eight isolates were categorized as weakly adherent and one isolate as strongly, under the tested conditions. Adherence was not related with a specific food product. The genes *cadF, ciaB, cj1349, mviN, pldA* and *tya* were detected in all adherent isolates; whilst *iroE* (4/9), *hecA* (1/9) and *hecB* (2/9) were detected only in few. *irgA* was not detected.

The occurrence of adherent *A. butzleri* isolates possessing virulence markers in food products of animal origin is an important factor to consider for the foodborne illness risk assessment.
SOURCE TRACKING GENETIC MARKERS IN CAMPYLOBACTER JEJUNI BY MLST AND MCGH
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Background
Campylobacter jejuni is a major cause of food-borne diarrheal illness in humans worldwide. Despite being a well known food and water-borne zoonosis, establishing the source of campylobacteriosis is still complicated (Hepworth et al., 2011).

Objectives
The aim of this study was to find genetic markers for clonal complex (CC) and source attribution in C. jejuni.

Methods
Multilocus sequence typing (MLST) and microarray-based genomic hybridization (MCGH) were applied to 69 C. jejuni isolates from different sources (wild birds, surface and wastewater, and human and bovine feces). Oligonucleotide DNA microarrays (BμG@S CJv3.0.0) were used to compare C. jejuni isolates, with C. jejuni NCTC 11168 and RM1221 as reference strains.

Conclusions
The population was genetically diverse. MLST differentiated the isolates into 38 sequence types (STs), of which 29 were included in seven different CCs. Ten new alleles were identified, which resulted in the assignment of 12 new STs.

MCGH confirmed most of the previously described variable and/or plasticity regions (Parker et al., 2006; Hepworth et al., 2011; Stabler et al., 2013). Combining MLST and MCGH results allowed us to detect several potential genetic markers associated with both, specific CCs and certain sources of isolation. More accurately, the genes cas2, cas1, Cj0659c-Cj0660c, Cj0887c, Cj1041-Cj1042, hsdM and hsdR related with the CCs ST-21, ST-42, ST-177 and ST-677; and the genes glpT, ldh and Cj1562 related with isolates from wild birds.
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GENETIC DIVERSITY AND HORIZONTAL GENE TRANSFER IN FOODBORNE ARCObACTER CRYAEROPHILUS ISOLATES
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Background

Arcobacter species have been recognized as potential food- and waterborne pathogens, and are implicated in human enteritis. The wide genetic diversity reported among Arcobacter species reflects the continuous evolving nature of these pathogens.

Objectives

The aim of this work was to determine the genetic diversity and virulence gene content of 10 Arcobacter cryaerophilus isolates obtained from clams, mussels and raw cow milk obtained in Vitoria, Spain.

Methods

Genetic diversity was studied by multilocus sequence typing (MLST). The obtained genotypes were analyzed using the MEGA 5.1 software, and possible recombination events were evaluated in silico using the RDP3 software. Recombination events were considered when detected by at least 3 methods. The presence of nine putative virulence genes was determined by PCR.

Conclusions

A total of 9 sequence types (ST) not previously described were identified among the 10 isolates. All the isolates harbored virulence genes. The most frequent were ciaB and mviN (100%), followed by tlyA (40%) and hecA (30%). Due to the incongruent phylogenetic relationships observed for glnA and glyA genes, recombination analysis was performed. This analysis detected potential recombination events in the ST413, ST415 and ST416 isolates, probably derived from interspecies recombination between A. cryaerophilus and A. skirrowii.
From these results, we conclude that MLST typing showed high strain diversity and that recombination should be considered as a potentially relevant mechanism generating genetic diversity in *A. cryaerophilus*. 
Background
Staphylococcal food poisoning represents the most prevalent foodborne intoxication worldwide. It is caused by oral intake of enterotoxins preformed by *Staphylococcus aureus* in food. The relevance of enterotoxins SEG and SEI in outbreaks of staphylococcal food poisoning is controversially discussed. Although SEG and SEI elicit emesis in a monkey feeding assay, there has been no conclusive proof of their emetic activity in humans.

Objectives
In this study, we present novel evidence suggesting that SEG and SEI can cause staphylococcal food poisoning and describe the special challenges associated with investigating outbreaks linked to SEG and SEI.

Methods
We analyzed all outbreaks registered with the Swiss Federal Office of Public Health, in which only *Staphylococcus aureus* strains harboring *seg* and *sei* linked to typical signs of staphylococcal food poisoning were isolated (n = 3).

Conclusions
The outbreaks were caused by consumption of raw goat cheese, potato salad, and semi-hard goat cheese, and were linked to strains assigned to CC45 (*agr* type I), CC30 (*agr* type III), and CC9 (*agr* type II), respectively. Investigation of the outbreaks was particularly challenging as, in contrast to classical staphylococcal enterotoxins, there are currently no suitable methods for detection of SEG and SEI in food and feces. Our data provides strong evidence suggesting that SEG and SEI can cause staphylococcal food poisoning in humans. Further research will be necessary to determine the individual role of SEG and SEI in outbreaks.
FOOD POISONING OUTBREAK AMONG CHILDREN AND STAFF AT SWISS BOARDING SCHOOL DUE TO RAW MILK CHEESE CONTAMINATED WITH STAPHYLOCOCCUS AUREUS OF GENOTYPE B

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Background

In 2014, an outbreak of staphylococcal food poisoning occurred at a Swiss boarding school due to consumption of Tomme, a soft cheese produced from raw cow milk. All 14 persons that had ingested the cheese fell ill, among them ten children.

Objectives

We present a food poisoning outbreak caused by raw milk cheese contaminated with a Staphylococcus (S.) aureus strain of genotype B.

Methods

The Tomme cheese was tested for presence of staphylococcal enterotoxins and S. aureus strains isolated from the cheese were characterized by spa typing and a DNA microarray.

Conclusions

The cheese exhibited low levels of staphylococcal enterotoxin A (> 6 ng SEA/g cheese) and high levels of staphylococcal enterotoxin D (> 200 ng SED/g cheese). A total of $10^7$ CFU coagulase-positive Staphylococci per gram cheese were detected, with three different S. aureus strains being present at levels higher than $10^6$ CFU/g. A strain exhibiting sea and sed was identified as the source of the outbreak. The strain was assigned to spa type t711 and CC8 and exhibited genetic criteria consistent with the characteristics of a genotype B strain. This genotype comprises bovine S. aureus exclusively associated with very high within-herd prevalence of mastitis and has been described as a major contaminant in Swiss raw milk cheese. It is highly likely that the raw milk used for production of the Tomme was strongly contaminated with S. aureus and that levels further increased due to growth of the organism and physical concentration effects during the cheese making process.
TEMPORAL EXPRESSION OF THE STAPHYLOCOCCAL ENTEROTOXIN D GENE UNDER NACl STRESS CONDITIONS ENCOUNTERED DURING FOOD PRODUCTION AND PRESERVATION

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Background
Staphylococcus aureus represents one of most osmotolerant food-borne pathogens. While its growth is repressed by competing bacteria, the organism exhibits a growth advantage at increased salt concentrations. Staphylococcal enterotoxin D (SED) leads to vomiting and diarrhea upon consumption. To date, the effect of NaCl on both sed expression and its regulatory control are unclear.

Objectives
We aimed to determine the impact of NaCl stress on sed expression and the influence of agr, sarA, and sigB on sed expression under NaCl stress.

Methods
Using qPCR, temporal expression of sed in LB and LB with 4.5% NaCl was compared, as well as sed expression between wild type strains and isogenic Δagr, ΔsarA, and ΔsigB mutants.

Conclusions
In general, NaCl stress led to decreased sed expression. However, one strain exhibited a trend towards increased sed expression under NaCl stress. One ΔsarA mutant each showed decreased sed expression in the early stationary and increased sed expression in the stationary growth phase under NaCl stress. No significant effect of agr on sed expression was detected, and only one ΔsigB mutant showed a significant decrease in sed expression in the early stationary phase under NaCl stress. These findings suggest high strain-specific variation in sed expression and its regulation under NaCl stress.
Background
In vitro digestion models are simplified artificial systems to simulate digestion processes in humans/animals. Our digestion system aimed to study structural changes, digestibility and bioavailability of food components and behaviour of microbes. The model consisted of four stages, imitating the environment of mouth, stomach, small intestine and large intestine.

Objectives
Selective agar plates ensured reliable enumeration of bacterial colonies, but cultivation took 2-3 days, especially in case of the slowly growing species (e.g. *Bifidobacterium*), therefore, qPCR technique was applied for enumeration of bacteria.

Methods
Microbial activity was investigated in the large intestinal phase. Bacterial community consisted of two probiotic (*Bifidobacterium* and *Lactobacillus*) and four neutral/potential pathogenic species (*Clostridium, Bacteroides, Enterococcus, Escherichia*), found frequently in the human large intestine. The effect of food components (including prebiotic carbohydrates) on proliferation and survival of bacteria was evaluated on selective agar plates and by qPCR, using newly designed species-specific primers in the latter case.

Conclusions
The use of species specific primers for qPCR-based quantitation was successful, the number of each bacteria was determined precisely. Differences between the plate count and qPCR techniques were within an order of magnitude range in most cases. However, selective agar plates are still widely used for enumeration of bacteria in laboratories, other methods like qPCR are used more and more frequently, due to their speed and reliability. Our experiments showed that carefully selected primers were able to ensure the specificity for quantitation of bacteria from mixed cultures, in the presence of interfering compounds, such as digestion juice.
EFFECT OF OZONATED AND CHLORINATED WATER ON MICROBIOLOGICAL LOAD AND SENSORY PROPERTIES OF FRESH LETTUCE (LACTUCA SATIVA) AND BROCCOLI (BRASSICA)

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Background

Ozonated and chlorinated water used as potential antimicrobials agents in food industry. Since 2001, when US Food and Drug Administration (FDA) announced the permission of Ozon's usage as a suitable antimicrobial agent for food, many investigations have been applied (Singh et al. 2002, Trinetta et al. 2011).

Objectives

The objectives of this experimental work are to determine the effectiveness of ozonated (33.3 mg/l) and chlorinated (50 mg/l) water on the reduction of microbiological load on two vegetable species. The project's target is to estimate the load of four pathogen microorganisms (Echerichia coli, Listeria monocytogenes, Salmonella spp, Enterobacteriaceae) at the different market points and highlight if those specific vegetables are safe for the consumers. One more target is to remark any changes at the qualitative and sensory characteristics. Furthermore, was examined the microbiological load of vegetables when they have been washed with tap, chlorinated and ozonated water.

Methods

The methods which used for the bacteria are Health Protection Agency (HPA), ISO 11290-1:1996/Amd 1:2004, ELOT EN ISO 6579:2003/TC1:2004, Statutory Instrument SI 2383, 1989 and BS 4285:3.7. The part of the sensory analysis consists of a specific panel with ten people, who tasted and evaluated vegetables, after treated and washed with three treatments.

Conclusions
According to the results was proved that the region of origin probably is important for the surcharge of the microbiological load, but this is not affected by the sampling point. Ozonated water has the best performance for all samples for the microbiological control and sensory analysis. In general, broccoli have less Enterobacteriaceae load than lettuce.
DETERMINATION OF KILLER ACTIVITIES OF THE YEAST STRAINS ISOLATED FROM NATURAL FERMENTATION MEDIA OF GEMLIK CULTIVAR BLACK OLIVES

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Background

Yeasts are common microorganisms present in spontaneous table olive fermentation. Killer yeasts secrete protein toxins that are lethal to sensitive strains. *Wickerhamomyces anomalus*, *Kluyveromyces marxianus*, *Pichia guilliermondii*, and *Saccharomyces cerevisiae* strains isolated from olive brines have been shown to produce killer toxins with a broad spectrum of activity against spoilage microorganisms of table olives. The killer characteristics of the yeasts have been applied to combat undesirable yeasts during the production of table olive.

Objectives

The aim of this study was to determine killer activities of the yeast strains isolated during natural fermentation process of “Gemlik” cultivar olives grown in Akhisar and Iznik regions of Turkey. The effect of regional differences on killer yeast profile of Gemlik olives was also investigated.

Methods

A total of 54 endogenic yeast strains were investigated for killer activity. These were belonging to 7 genera as; *Candida*, *Aureobasidium*, *Debaryomyces*, *Cryptococcus*, *Kloeckera*, *Pichia* and *Clavispora*. Killer activity was detected by agar diffusion assay. In the method, two killers; *S.cerevisiae*NCYC 232 (K₁), *S.cerevisiae*NCYC 738 (K₂) and a sensitive strain; *S.cerevisiae*NCYC 1006 were used as reference yeasts.

Conclusions

From 54 strains, 40 were found as killer and 6 were killer-sensitive strains. Additionally, 8 strains were found to have neutral character. All *Debaryomyces hansenii*, *Pichia anomala*, *Clavispora lusitaniae* and *Kloeckera apiculata* strains were found as killer. No sensitive strain was found among the tested isolates. In conclusion, killer behavior of the endogenic yeast strains of Gemlik cultivar olives of Akhisar and Iznik regions was found as common characteristics.
INVESTIGATION OF BRUCELLA SPP. AND BRUCELLA DNA IN RAW MILK OBTAINED FROM TRAKYA REGION, EDIRNE, TURKEY

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Background

Brucellosis is an important zoonotic disease transmitted to human by consuming contaminated milk and milk products. The prevention from Brucellosis is provided by live-attenuated vaccines for animals. Although Trakya is the pilot region in Turkey, for vaccination of animals, Brucellosis is endemoepidemic in this region. Thus, many dairy companies supply their milk from the cows grown in this region; screening of milks for Brucella contamination becomes very important.

Objectives

Our aim was to investigate Brucella spp. in milk, procured from Edirne and compare the conventional and molecular methods in terms of screening Brucella spp. in milk.

Methods

Milk samples from 99 cows at 12 different barns in 5 different villages of Edirne were collected. Bacteriological analyses and Q-PCR were applied to all samples. For samples that were culture negative and Q-PCR positive, a Q-PCR based method was evolved to differentiate the virulent and the vaccine strains.

Conclusions

In 2 of the 99 milk samples, Brucella spp. was isolated by bacteriological methods. After Q-PCR, B. ovis and B. melitensis were not detected, however, in 16 of the 99 samples B. abortus was detected. After the Q-PCR based method evolved to differentiate virulent and vaccine strains, only the 2 bacteriologically positive samples were detected to be positive.

As a result, it is reported that; 2.02% of the samples were detected to contain Brucella, both with the bacteriological method and PCR. It is also determined that, to obtain true positive results in Brucella spp. screening studies for milk, differentiating the virulent and vaccine strain should not be disregarded.
PRODUCTION OF ACE-INHIBITORY PEPTIDES BY LACTIC ACID BACTERIA ISOLATED FROM TRADITIONAL GREEK DAIRY PRODUCTS.

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Background

Fermented dairy products are generally considered to be beneficial for human health as they contain live lactic acid bacteria frequently exhibiting biofunctional features. Here we present 55 lactic acid bacteria isolated from three traditional Greek yogurt samples and five fermented milk samples where their ability to produce bioactive peptides with inhibitory activity against the angiotensin-converting enzyme (ACE) was evaluated.

Objectives

The aim of this study is the potential of several lactic acid bacteria strains to be used as health promoting starter cultures with interesting technological features.

Methods

Strain typing was performed by rep-PCR, while 16S rDNA sequencing was used for the identification at the species level. The angiotensin-converting enzyme inhibitory (ACE-I) activity of the isolates was evaluated using a spectrophotometric assay with N-Hippuryl-His-Leu hydrate as substrate. Finally, a semi-preparative HPLC analysis was conducted using an ACE-5C18 column and positive fractions were subjected to MS analysis.

Conclusions

During this work 55 microorganisms have been isolated from traditional Greek yogurt and fermented milk samples. Strain typing by rep-PCR showed that the 33 isolated bacilli corresponded to 10 and the 22 cocci to 12 strain groups, respectively. The
strains did not show neither strong acidification capacity, nor peptidolytic activity and citrate metabolism, so their use as starter cultures does not seem suitable. Among them, many strains possessed interesting technological features as proteolysis and lipolysis and few exhibited high ACE-I activity. Purification of the ACE-I peptides was performed by RP-HPLC. These results indicate their potential use as adjunct or probiotic cultures.
EVALUATION OF CINNAMON AND MARJORAM ESSENTIAL OILS FOR THE CONTROL OF MYCOTOXIGENIC FUSARIUM SPECIES

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Background

*F. culmorum* and *F. graminearum* are the most common causes of Fusarium ear blight of corn, wheat and other grains. Both of them produce the trichothecenes mycotoxins and *F. graminearum* also produces zearalenone.

Objectives

The aim of this study was to evaluate the efficacy of cinnamon and marjoram essential oil (EOs) to control the growth of Fusarium species in vitro and to observe the changes of morphological characteristics of the fungi by SEM images after treatment with the oils.

Methods

Colony growth inhibition effect of the EO vapors was investigated on the *Fusarium* species by the reversed Petri dish method using 0.070 mg/l, 0.14 mg/l, 0.28 mg/l and 0.56 mg/l EO/dish. The antifungal effect was measured by determination of growth-rate (mm/day) and antifungal index (%). MIC (minimal inhibitory concentration) and MFC (minimal fungicidal concentration) values were determined by agar dilution method. Morphological characteristics of the hyphae and micro conidia of *F. culmorum* were observed using a scanning electron microscope.

Conclusions

Cinnamon EO vapor at 0.28 mg/l concentration caused total growth inhibition of all investigated species. Using marjoram EO vapor total inhibition was not detected, but growth-rates and colony diameters were reduced in each case. Agar dilution method was more effective to inhibit the growth of *F. culmorum* (MIC value ≥2.5 mg/ml). SEM imaging of *F. culmorum* showed that EO treated hyphae were thinner. Rupture of the cell wall and leakage of the cytoplasm contents were also observed. This study showed that EOs have potential antifungal effect against mycotoxigenic Fusarium species.
Background
In fermented foods, the presence of biogenic amines (BAs), which were produced by microbial decarboxylase activity of microorganism, might serve as a useful indicator of food poisoning. Although BAs are also frequently found in ganjang, representative Korean traditional soy-sauce, there is no information about the BAs-producing microorganism in ganjang.

Objectives
In this study, to investigate microorganism responsible for BAs production in ganjang, metabolite and bacterial community analysis were performed during entire fermentation process of ganjang.

Methods
Ganjang with a NaCl concentration of approximately 18% (w/v) was prepared in one batch using traditional manufacturing method. The same amounts of ganjang were periodically sampled and their pH, NaCl concentration, and cell number were measured. $^1$H NMR and PCR-DGGE were applied for analysis of metabolites and bacterial succession during the ganjang fermentation, respectively. In order to investigate the correlations among the ganjang samples, bacterial communities, and metabolites, a multivariate statistical redundancy analysis was performed using the vegan package of the R programming environment.

Conclusions
Metabolite profiling and bacterial community analysis using a $^1$H NMR and PCR-DGGE showed that a large amount of putrescine detected after 74 days of fermentation, which time begins to predominate the members of Chromohalobacter. A statistical analysis based on metabolite profiling and bacterial succession data clearly shows that members of Chromohalobacter are involved in the production of putrescine during ganjang fermentation. This study will allow for the successful understanding of the BAs-producing microorganism in ganjang.
CHARACTERISTICS OF CONTAMINANT MICROBES IN COMPOUND FEEDS AND THE GROWTH INHIBITION EFFECT BY PROBIOTIC B. SUBTILIS

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Background

During long term storage of animal compound feed, contaminated microbes can grow with providing a problem of feed safety.

Objectives

The aim of this study was to investigate contaminant microbes and the effects of probiotic B. subtilis to inhibit them in animal feeds.

Methods

During storage of feed at room temperature, the changes of pH and titrable acidity were monitored with total microbial counts. Bacterial and fungal strains were isolated and tested for antibiotics resistance. Contaminated microbes were further identified by the sequencing of 16S and 18S rDNAs.

Probiotic B. subtilis was inoculated into both non-sterile and sterile feeds and tested for the growth inhibition of contaminated microbes during storage of feed at room temperature with pH change and probiotic viability.

Conclusions

Most contaminated microbes were originated from soil and plants. Sterile and non-sterile compound feeds displayed a different pattern of pH change. B. subtilis showed a normal growth and inhibited the cell growth of contaminant microbes.
EVALUATION OF MICROBIAL CONTAMINATION LEVELS AT THE PRE- AND POST-GERMINATION STAGES ON SPROUTS.

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Background
The consumption of raw sprouted seeds has led to a large number of outbreaks of foodborne illness in a great number of countries.

Objectives
The objective of this study was to investigate and evaluate microbial contamination levels of several kinds of sprouts in pre- and post-germination stage.

Methods
Fourteen kind of sprout seeds were purchased and analyzed. Quantitative analyses comprised aerobic plate counts (APCs) and the measurement of coliforms and Bacillus cereus levels, whereas qualitative analyses involved assessing the levels of Escherichia coli and major foodborne pathogens (E. coli O157:H7, Listeria monocytogenes, Salmonella spp. and Staphylococcus aureus).

Conclusions
The APCs (2.3-4.07 log CFU g⁻¹) for sprouts seeds increased by approximately 3-4 log CFU g⁻¹ during germinating, reaching 5.61-8.94 log CFU g⁻¹. Similarly, increasing trends were noted in the level of B. cereus (N.D.-2.62 log CFU g⁻¹ at the seed stage, increasing to N.D.-3.14 log CFU g⁻¹ by the germinated stage). E. coli, E. coli O157:H7, Salmonella spp. and L. monocytogenes were not detected in the sprouts analyzed in this study. The present study provides comprehensive information regarding the microbiological safety of seeds and sprouts during manufacturing.
Background

Antioxidants are compounds with role in food preservation, as well as in the prevention and treatment of some degenerative diseases. Despite their high amounts in plants, there is a growing interest for less conventional sources. Cyanobacteria, as a particular producers of bioactive compounds, represent an important source of antioxidants, which in terms of chemical structure include diverse compounds.

Objectives

In the present study, the antioxidant capacity of three filamentous cyanobacterial strains belonging to the *Anabaena*, *Nostoc* and *Spirulina* genera have been tested. Furthermore, the content of C-phycocyanin, a pigment known as antioxidant compound was determined.

Methods

The antioxidant capacity was tested using DPPH and FRAP assays. In DPPH assay, strains *Spirulina* S1 and *Anabaena* C5 showed similar IC\textsubscript{50} values of 118.47 µg/ml and 120.26 µg/ml respectively. In FRAP assay, the highest antioxidant activity was observed in *Spirulina* strain S1, ranging from 452.41 to 637.38 mg of ascorbic acid (AA) equivalent/gram of dry extract. The lowest antioxidant capacity expressed strain *Nostoc* 2S9B in both assays (IC\textsubscript{25} of 21.93 µg/ml, and 142.21-350.18 mg of AA equivalent/gram of dry extract).

The C-phycocyanin content was determined spectrophotometrically. The highest value of 38.63 µg/ml was detected in strain *Spirulina* S1, while lower contents were obtained in strains *Nostoc* 2S9B and *Anabaena* C5 (6.57 µg/ml and 9.34 µg/ml respectively).

Conclusions
Since *Spirulina* strain S1 has expressed the highest antioxidant capacity in both assays, it could be further investigated as a potential source of antioxidants. High C-phycocyanin content found in this strain could be associated with its antioxidant activity.
Background
Securing the food chains from primary production to consumer-ready food against major deliberate, accidental or natural contaminations is directly related to the safety of food products. Spices and herbs are natural products of almost all processed food that can be contaminated with several microorganisms and toxins. The identification of those condiments would be difficult, as consumers and experts often focus on major food ingredients instead of minor components. Securing of spices and herbs commodity chains is the key issue within the EU FP7 project SPICED.

Objectives
Characterization of the heterogeneous matrices and their respective production and supply chains in context with relevant biological hazards and the improvement of the knowledge on biological hazard properties as well as on-site and high throughput diagnostic methods for appropriate detection are main tasks.

Methods
Within the SPICED project we mainly focusses on the toxins Ricin, which is the toxic component from castor beans of *Ricinus communis*, and the Enterotoxin-type B produced by the gram-positive bacteria *Staphylococcus aureus* (SEB). Their qualitative and quantitative detection in spices and herbs including the development of sample preparation methods were performed. For this purpose different types of spices and herbs were spiked with the toxins, stored at room temperature and analyzed e.g. with immunological based methods. Fieldable technologies and methods for the rapid on-site detection of toxins in the different were also examined.

Conclusions
The intention is to present the recent results on the detection and sample preparation of the toxins from the spices and herbs with laboratory and field methods.
EFFECT OF THERMALLY PROCESSED RICE KOJI EXTRACTS ON 
CHRONOLOGICAL LIFE SPAN OF SACCHAROMYCSES CEREVISIAE ON HIGH-
SUGAR FERMENTATION
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Background

Yeast cell growth and viability on high ethanol concentrations become limiting factors in fermentation productivity and ethanol yield. That is why measures to improve ethanol tolerance in yeast have been expected.

Objectives
This study focused on koji as a material for sake brewing which provides nutrients for yeast cell growth. Our goal is to develop strategies for extending the life span of yeast by revealing the influence of heat-processed koji extracts on the survival of yeast under ethanol stress.

**Methods**

Rice koji extracts were prepared by water extracts of dried rice koji heated at 60°C for 5 hrs. The koji extract was autoclaved for different times at 121°C. The availability of thermally processed rice koji extracts was investigated by analysis of ethanol tolerance and survivals on high-sugar fermentation in *S. cerevisiae* K-701 while making a comparison between heated and non-heated koji extracts.

**Conclusions**
Addition of heat-processed *koji* extracts to high-sugar medium led to an improvement of chronological life span in *S. cerevisiae* K-701. The longer the heating time of *koji* extracts was, the higher the population of *S. cerevisiae* was during the late stationary phase. The Maillard reaction products which occurred in the heat-processed *koji* extracts had an explicit effect on the survivals of yeast under lethal ethanol stress. These show that heat-processed rice *koji* extracts were served as protectants of both ethanol and oxidative stresses when yeast cells became more oxidized in the stationary phase at higher ethanol concentrations.
Introduction
Oxidative stress arising from an imbalance in the human antioxidant status, reactive oxygen species vs. defense and repair mechanisms, is mainly responsible for the development of pathological disease. Phenolic compounds in safflower seeds are receiving much attention as potential therapeutic agents against several pathological diseases. The antioxidant properties of safflower seeds is matairesinol 4‘-o-β-D-glucose, 8‘-hydroxyarctigenin 4‘-o-β-D-glucose, matairesinol, 8‘-hydroxyarctigenin, N-feruloylserotonin 5-o-β-D glucoside, N-(p-coumaroyl)-serotonin-5-o-β-D-glucoside, N-feruloylserotonin, N-(p-coumaroyl)serotonin, luteolin 7-o-β-D-glucoside, luteolin, acacetin 7-o-β-D-glucuronide, and acacetin.

Objective.
In our experiment, bacteria isolated from ginger fermented food were used to make phenolic compound from aglycon. It makes to increase antioxidant activity of safflower seeds.

Methods
Fermented samples were got from one of the member of Korea royal cuisine institute. Fermented sample was inoculated in tryptic soy broth media and incubated 28°C for 48h. After bacteria grow, bacteria were centrifuged 6000rpm at 4°C for 20 min. Bacteria were dissolved 60ml mineral salt media (MSM) (20g/L) and mixed with 30g safflower seed. Each bottle was incubated 28°C for one week. For phenolic compound extraction, each safflower seed mixed with bacteria was sonicated in hexane for 30min twice and filtered. The defatted seed residue was extracted twice with 70% EtOH under sonication, filtered and evaporated under reduced pressure. The extract was used for HPLC and GC-MS and DPPH activity.

Conclusion
This study was to evaluate the antioxidant activity in safflower seeds after fermentation. Bacteria that improved antioxidant activity in safflower seeds after fermentation was identified Sphingomonas sp using 16s rRNA sequencing. Safflower seed was improved antioxidant activity from 7.11% to 51% after 5 days. In HPLC analysis, Major peak of safflower seed extracts after fermentation was changed after 5 days under aerobic condition at 30°C. Furthermore analysis, we will conduct to investigate which compound in the safflower seed will be changed to phenolic compound for increasing antioxidant activity.
EFFECTS OF SEED TREATMENT BY COLD ATMOSPHERIC PLASMA ON CUCUMBER SEEDLING GROWTH

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Background
Cold atmospheric plasma can have different effects on seed germination and seedling growth depending on the plasma frequency, the power and its environmental conditions.

Objectives
Cold plasmas are already well-known for their improvement effect on plant growth. However, the mechanisms of this ability are unclear.

Methods
Non-thermal atmospheric-pressure DBD plasma in He was applied with different exposure times on cucumber seed. Changes in the seed surface with different exposure time were examined by scanning electron microscope (SEM). Direct and indirect influence on cucumber seed of the cold plasma is compared in terms of seedling growth.

Conclusions
Germination and early seedling growth of cucumber were improved by the plasma treatment for 6 second. In contrast, the seeds exposed for 10 minutes were not germinated. In 6 sec treatment and no treatment control, gold nanoparticles were observed along the cell wall. There were no particles observed in the 10 min treatment. It suggests that no tissue damage is observed visually or microscopically following 6 sec and control, but the rough surface of seeds were smooth and damaged at long exposure time (10 min). We observed that the indirect method had more enhanced effect on growth than direct method. This infers to an important conclusion that the effect of plasma on improved growth of seedlings is due to various plasma components that can exist inside or outside the plasma region not mechanical power such as an abrasion of seed surface.
METABOLICALLY ACTIVE BACTERIA IN LYMPHATIC TISSUES OF PIGS AND ITS SPREAD DURING SLAUGHTER
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Background
The occurrence of potential food-borne pathogenic organisms in lymphatic tissues of farm animals has practical relevance for carcass contamination. A transmission of pathogens, which can occur from harvesting until the last step in the meat-processing chain, remains the preeminent difficulty in slaughter house hygiene.

Objectives
The aims of this study were i.) to evaluate the contamination degree of porcine cervical musculature during slaughter, ii.) to compare the bacterial microbial community detected by DNA-based amplicon sequencing with RNA-based amplicon sequencing and thus measuring the fraction of metabolically active bacteria in lymphatic tissues.

Methods
Mandibular lymph nodes, tonsils and scrapings from cervical musculature were sampled from eight healthy slaughter pigs derived from different farmers. RNA was isolated from lymph node cortex regions and tonsils (n=16) and DNA was isolated from cervical musculature (n=8). Total RNA was transcribed into cDNA and all samples were sequenced with Roche/454 using 16S rRNA gene amplicons.

Conclusions
Organisms relevant for food safety, e.g. Erysipelothrix, Listeria, Escherichia and Pseudomonas, could be detected in lymphatic tissues. RNA-based amplicon sequencing revealed a highly diverse, metabolically active bacterial microbiome in mandibular lymph nodes and tonsils; consisting of 348 and 390 operational taxonomic units (OTUs) respectively (30% shared OTUs). The functional metagenome prediction (PICRUST) and subsequent assignments of orthologues to COG categories revealed amino acid transport and metabolism-related orthologues being increased in lymph nodes compared to tonsils. Both, mandibular lymph nodes and tonsils of slaughter pigs harbor a replication-competent bacterial microbiome indicating a possible contamination source during and after slaughter.
ISOLATION AND CHARACTERIZATION OF THE MICROFLORA PRESENT IN A MINIMALLY PROCESSED VEGETABLES PLANT

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Background
Food safety is an area of growing concern due to the increasing demand for microbiologically safe products. Additionally, disinfection techniques must evolve to cope with the development of antimicrobial resistance. An efficient disinfection strategy must consider the type of microbial contaminant. Therefore, the knowledge on the microorganisms present in an industrial process is crucial to define the best strategy for their control.

Objectives
The aims of the present work were to isolate and characterize the resident microflora present in a minimally processed vegetables (MPV) plant.

Methods

The microorganisms were isolated from a MPV plant process surfaces by cotton swabbing, from the air and from the vegetable surfaces. The isolates were identified by 16S rRNA gene sequencing with 4 primers: 27F, 518F, 800R e 1492R. The isolates were also characterized for their production of proteases, gelatinases and siderophores, quorum-sensing inhibition and biofilm formation ability.

Conclusions
From 50 different isolates 46% belong to the Pseudomonas genera and 22 were from different species. Some microorganisms persisted along the food chain, which means that the cleaning process should be targeted for the more resistant microorganisms. Most of the microbial isolates are capable of producing virulence and food spoilage molecules. Moreover, Pseudomonas were the genera with a higher biofilm formation ability, being the predominant microflora and showing recalcitrant properties along the process chain.
Biogenic amines (BA) are low-molecular nitrogenous organic bases, which are formed in foodstuffs by microbial decarboxylation of the precursor amino acids and are potentially toxic to human health. Cheeses are among the foods most commonly associated with the presence of BA (1). Concentration of BA results from a balance between amino acid decarboxylating and amine oxidising activities.

Objectives
To correlate the content of BA of twenty traditional Apulian or Sicilian (Southern Italy) cheeses with several technology and microbiological features such as time of ripening, pH, concentration of precursor free amino acids (FAA), and occurrence of decarboxylase-positive lactic acid bacteria.

Methods
Cheeses were analysed through plate counting and HPLC, and lactic acid bacteria isolated from cheeses were assayed (by decarboxylase medium) for their capacity to generate BA. Principal Component Analysis was performed to find the effect of different parameters on the distribution of the cheeses.

Conclusions
Although short-ripened cheeses did not show significant BA concentrations, the only BA showing high positive correlation with time of ripening was histamine. Concentration of histidine and, especially, percentage of histidine-decarboxylase bacteria presumably affected histamine concentration. High pH values were negatively correlated to the concentration of tyramine, putrescine, and cadaverine (Fig. 1). Fifty percent of the cheeses contained at least one BA at potentially toxic concentrations. Unambiguous and ever-valid relations among parameters and BA are difficult to determine, because BA are the result of combined and varied factors.
DYNAMICS OF BACTERIAL COMMUNITIES DURING THE RIPENING PROCESS OF CROATIAN RAW EWE’S MILK CHEESE

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Background

Croatian raw ewe’s milk cheeses are hard cheeses that are produced by traditional techniques without pasteurization and application of starter cultures, and they are characterized by an aging process of 90-120 days. This aging time is rather long and the low pH and water activity (aw) of ripened cheese usually does not support the growth of pathogens. However, due to the commercial pressure, the ripening time has been shortened and it has become market practice to sell the cheese as soon as possible.

Objectives

A close monitoring of bacterial communities and a reliable identification of indigenous microflora is crucial in order to maintain the quality and safety of artisan cheese.

Methods

Cheese samples of three cheese types (0, 45 and 90 days of ripening) were collected from 6 cheese makers. Total DNA was extracted and microbial diversity was investigated based on fingerprinting in combination with next generation sequencing of 16S rRNA gene amplicons.

Conclusions

Overall up to 213 OTU97 could be assigned. Twenty of the major OTUs were present in all samples and include mostly LAB, mainly Lactococcus and Enterococcus species. Abundance and diversity of these genera differed to a large extent between the 3 investigated cheese types and in response to the ripening process. Also a large number of non LAB genera could be identified based on phylogenetic alignments including mainly Enterobacteriaceae and Staphylococcaceae. Some species belonging to these two families could be clearly assigned to species which are known as potential human pathogens. However, during cheese ripening their abundance was reduced.
Background
Sausages from game meat produced by traditional procedures by small scale producers in Croatia do not include the application of starter cultures. Such artisan products are increasingly appreciated because of their sensory properties and authenticity, although considerations regarding their microbiological safety are existent.

Objectives
An in depth analysis and characterization of the indigenous microflora of traditionally produced wild boar meat sausages from Croatia is needed in order to estimate microbiological hazards and to preserve an indigenous microbial pool.

Methods
Traditionally produced wild boar sausages were collected from three farms during ripening (0, 2, 4, 7, 10, 20 and 40 days) and analyzed for spoilage and pathogenic microflora (L. monocytogenes, Salmonella spp., total Enterobacteriaceae, coliforms, yeasts and molds, B. cereus group and coagulase-positive staphylococci) as well as for beneficial microbes e.g. lactic acid bacteria (LAB) and coagulase-negative staphylococci (CNS). Genomic DNA was extracted from 720 LAB and CNS isolates. After RAPD and rep-PCR based grouping identification was done by PCR and partly 16S rRNA gene sequencing.

Conclusions
A remarkable diversity was found among the investigated LAB and CNS populations of which several isolates relevant for sausage ripening and product quality were identified as e.g. Lactobacillus sakei, Leuconostoc mesenteroides, Staphylococcus xylosus, Staphylococcus saprophyticus and Staphylococcus warneri. Although some spoilage and pathogenic microflora were present at the initial stages of sausages’ ripening, they were absent in the ripened “ready to eat” sausage.
REDUCING THE RISK OF BABY LEAF SURFACE BACTERIA WITH IRRIGATION RESTRICTIONS

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Background

Baby leaf crops like Spinach (*Spinacia oleracea*), Chard (*Beta vulgaris*) and Rocket (*Diplotaxis tenuifolia*) are grown on huge open space flat fields in Sweden. It is hard to stop birds and wild animals to visit the field with related risks of E. coli infections. But as a grower, one can make sure the irrigation water doesn’t act as a bacteria vector.

Objectives

The main objective of this study was to reduce the risk of E. coli bacteria reaching baby leaf crops during crop growth in field, thereby reducing the risk of food borne diseases further down the delivery chain.

Methods

Leaves were grown with more or less constant optimal wet conditions in the root zone (irrigation water given up to field capacity every other day) or with restricted irrigation (drought conditions) in the final part of the growth period. 24 hours prior to harvest plants were sprayed with a known quantity of E. coli. Harvested leaves were put in a stomacher and the leaf rinsates plated on agar and the colonies counted and compared to non-sprayed leaves.

Conclusions

Irrigation restriction only had minor impact on E. coli colonization on the leaf surface. The yield and quality was significantly reduced and the content of bioactive compounds affected. The main conclusion is that drought stress during the final part of baby leaf production is not to recommend and as a way to reduce the risk of food borne diseases caused by baby leaf salads.
IMPROVED DETECTION OF ESCHERICHIA COLI AND COLIFORM BACTERIA
BY MULTIPLEX PCR

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Background
The presence of coliform bacteria is routinely assessed to establish the microbiological safety of water supplies and raw or processed foods. Coliforms are a group of lactose-fermenting Enterobacteriaceae, which most likely acquired the lacZ gene by horizontal transfer and therefore constitute a polyphyletic group. Among this group of bacteria is Escherichia coli, the pathogen that is most frequently associated with foodborne disease outbreaks and is often identified by β-glucuronidase enzymatic activity or by the redundant detection of uidA by PCR.

Objectives
Because a significant fraction of essential E. coli genes are preserved throughout the bacterial kingdom, alternative oligonucleotide primers for specific E. coli detection are not easily identified.

Methods
Here two strategies were used to design oligonucleotide primers with differing levels of specificity for the simultaneous detection of total coliforms and E. coli by multiplex PCR. A lacZ sequence and an orphan gene were chosen as targets for amplification.

Conclusions
A comparison with previously described primers indicates an increase in identification efficacy when tested with laboratory collection and lactose-fermenting strains isolated from dairy samples. While lacZ amplicons were found in a wide range of lactose-fermenting strains, amplification using the selected orphan gene was highly specific for E. coli. Additionally, the detection of this target sequence is non-redundant with enzymatic methods.
EFFECT OF OXYGEN CONCENTRATION AND GAS PRESSURE ON THE GROWTH OF ARTHROSPIRA SP. PCC 8005

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Background
The MELiSSA project of the European Space Agency, ESA, aims to create a bio-regenerative life support system for long term space travels. One of its bioreactors contains *Arthrospira* sp. PCC 8005, an edible cyanobacteria that can remove CO₂ and provide O₂. Normal operating conditions will likely enhance oxygen concentration and gas pressure buildup.

Objectives
The objective of this study is to elucidate the impact of oxygen concentration and gas pressure on the duration of the lag phase, the specific growth rate and the maximum number of attained generations in a batch culture of *Arthrospira* sp. PCC 8005.

Methods
Cultures of *Arthrospira* sp. PCC 8005 were grown in 100 ml closed septum vials with Zarrouk-UBP medium, at 30 °C with 32 µE m⁻² s⁻¹ of irradiance and shaken at 130 rpm. Growth was monitored measuring OD₇₅₀nm. Produced oxygen was accumulated. Oxygen tests were done with an initial 100% oxygen gas phase, in comparison to 100% argon, both at initial 1 bar. Pressure tests were done at 1, 2 and 3 bars of starting absolute pressure in 100% argon. Growth parameters were calculated using a modified Gompertz function. Statistical analysis was performed with 95% confidence interval T-Test.

Conclusions
The oxygen concentration has only a limited but statistically significant effect on the growth rate. Growth tests at elevated pressures of up to 3 bar had no effect on neither growth parameter. Test results at elevated pressures with 100% oxygen pressures, are expected to prove and further quantify the inhibitory effect of oxygen on the growth.
DETECTION OF ITURIN AND FENGYCIN BY MALDI-TOF IN FOOD MATRIX OF PUBA, A STAPLE BRAZILIAN FERMENTED FOOD

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Background
Puba or carimã is a Brazilian staple food obtained by spontaneous submerged fermentation of cassava (Manihot esculenta, Crantz) roots. Traditional fermentation of cassava is predominantly performed by lactic acid bacteria, but yeasts and Bacillus spp. were also found in our sample.

Objectives
To investigate the production of lipopeptides by Bacillus sp. P5 isolated from puba in the culture medium and in puba matrix by MALDI-TOF.

Methods
In a previous study, 16S rRNA of *Bacillus* sp. P5, isolated from *puba*, was sequenced. The phylogenetic analyses showed recovery in a node with 92% of support with *B. amyloliquefaciens*. The culture supernatant was subjected to extraction with n-butanol (Landy medium) purification by reversed-phase HPLC (C18 column) and MALDI-TOF mass spectrometry analysis. Surfactins, iturins and fengycins were detected. To investigate the production of lipopeptides by *Bacillus* sp. P5 isolated from *puba* in the culture medium and in *puba* matrix by MALDI-TOF. An aqueous suspension of *puba* was extracted with butanol and concentrated in Zip Tip® C18. Molecular mass was determined by MALDI-TOF. Data were acquired in the mass range m/z 1,000-4,000 for the butanol extract. The results are shown in Figure 1. Two clusters were found: iturin and fengycin (Fgy) (1A). Expansion of the spectrogram of Fgy indicated protonated ion and sodium and potassium adducts (1B), and in (C) MS/MS spectrum of precursor ion of m/z 1478.5 (Fgy C_{17}) which generated fingerprints of fgy A.

**Conclusions**

In conclusion, *puba* is manufactured in an unhygienic environment, and iturin and fengycin exhibit biocontrol properties that are probably important for the safety of this product.
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BACTERIAL ASSESSMENT OF FOOD - HANDLERS IN SARI CITY AND ITS SUBURB, MAZANDARAN PROVINCE, NORTH OF IRAN. 
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Background
Diarrheal diseases, mostly caused by food borne or waterborne microbial pathogens, are leading causes of illness and deaths in developing countries, killing an estimated 1.9 million people annually at the global level.

Objectives
This study was designed to determine the prevalence of carriers of pathogenic bacteria among the food-handlers employed at food service facilities in Sari city and its suburbs, Mazandaran Province, Iran

Methods
Of the total subjects examined, 62.2 % were found to be carriers of some pathogenic bacteria.

The pathogens isolated and identified were the bacteria Staphylococcus aureus, shigella sonnei and pseudomonas aeroginosa.

Fast food makers were the greatest prevalence of bacteria in respect of their fingernail contents (86.6 %, p=0.04) followed by butchers (76.4%), storekeepers (73%), bakers (58.3%) and restaurant workers (51.5 %). Staphylococcus aureus was the predominant bacteria isolated from nail-washing samples (45.9%), followed by Escherichia coli (28.4%), coliform (18.2%) and Pseudomonas aeroginosa (6.5%),

Illiterate food–handlers had the highest percentage of positive cultures from nail-washing specimens (86.8%, p= 0.0005) and the lowest rate of infestation was among university educated subjects (33.3 %).

The rate of of Staphylococcus aureus nasal carrier among food-handlers was 18.6 % with the highest rate among store-keepers (31.7%) followed by bakers (24.3%), restaurant workers (19.5%), butchers (14.6%) and fast-food makers (9.7%). Shigella sonneii were isolated from stool samples of two food-handlers (0.9%).

Conclusions
The finding emphasized the importance of food-handlers as potential sources of infections. A medical check –up program with health education could improve worker’s health status.
ISOLATION AND MOLECULAR IDENTIFICATION OF LACTIC ACID BACTERIA FROM DONKEY MILK
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Background

Lactic acid bacteria (LAB) are living organisms that play a major role in many food and dairy products as a probiotics. Donkey’s milk is being used as a source of human nutrition, especially for infants with allergy to cow milk proteins, due to its antimicrobial components, many defense factors and probiotic properties. Few studies have so far dealt with the microbiota in donkey’s milk, specifically lactic acid bacteria (LAB) diversity.

Objectives
This study was undertaken to identify the microflora of LAB in donkey's milk in IRAN.

Methods
The culture-dependent microbial and 16S rRNA sequencing techniques were used to identify lactic acid bacteria species in donkey milk. A total of 250 LABs from donkey milk were assessed by culture-dependent microbial tests, including culture at different temperatures, pH and Multiple antibiotic resistance indexing.

Conclusions

\textit{Enterococcus faecalis} (55\%) and \textit{Streptococcus devriesei} (45\%) were the most prevalent species in donkey milk. No other LAB belonging to the most technically important genera was identified. Biochemical and sugar fermenting analyses confirmed the results of 16S rRNA sequencing. Both species identified were sensitive to nearly all antibiotic tested, including vancomycin. Few strains were found to have >99\% sequence similarity with \textit{Streptococcus devriesei}, indicating the presence of this LAB in donkey milk. This is the first study to investigate the LAB microbiota present in donkey’s milk in IRAN.
BIOGEOGRAPHY OF NON-SACCHAROMYCES YEASTS ASSOCIATED WITH SPONTANEOUSLY FERMENTED MUSTS FROM DIFFERENT VINEYARDS AND WINE PRODUCING REGIONS IN GREECE

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Background

Non-Saccharomyces (wild) yeasts may confer diverse chemical composition and improved sensory properties to wines. Yet, our knowledge on population structure of these species is limited.

Objectives

Species diversity and population structure of 486 wild yeasts associated with spontaneously fermented musts from two geographically separated regions in Greece, Peza and Nemea, were assessed. The efficiency of different fingerprinting methods in resolving yeast populations was evaluated.

Methods

Species identification was conducted by PCR-RFLP analysis. For molecular typing the tandem repeat-tRNA (TRtRNA)-PCR and RAPD analysis with primers (GTG)$_3$, R$_5$ or RF$_2$ were applied. Molecular patterns and vineyard populations were clustered by Principal Coordinates Analysis and UPGMA, respectively. Differences of populations were evaluated by analysis of molecular variance (AMOVA).

Conclusions

Ten populations of wild species were identified and further genotyped by TRtRNA-PCR with the primer ISSR-MB (Lachancea thermotolerans, Metschnikowia pulcherrima and Hanseniaspora osmophila), and RAPD with the primer RF$_2$ (Candida zemplinina), (GTG)$_3$ (Hanseniaspora uvarum and Hyphopichia pseudooburtonii) or R$_5$ (Issatchenkia orientalis, Pichia anomala, Torulaspora delbrueckii, and Hanseniaspora guilliermondii). Except for I. orientalis, the percentage of biodiversity of yeast populations was higher in Nemea than in Peza. No particular grouping of banding patterns according to the sampling point or to the vineyard of origin was observed.
However, AMOVA showed that populations of *C. zemplinina* and *L. thermotolerans* differed significantly between the two regions. Results point to the presence of region specific sub-populations of non-*Saccharomyces* yeasts species that could enhance the Greek wines distinctiveness.

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SELECTION OF INDIGENOUS YEASTS FOR PRODUCTION OF QUALITY TERROIR-DRIVEN GREEK WINES

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Background
According to the microbial terroir concept, the use of indigenous yeasts as starters in winemaking may add to the uniqueness of regional wines.

Objectives
Genetically different yeast strains from spontaneously fermented musts from Santorini island, Mantinea plateau and Nemea region were characterized phenotypically in order to select strains with desirable enological properties to be used as starters.

Methods
Strains belonging to Candida zemplinina, Issatchenkia orientalis, Hanseniaspora quilliermondii, H. opuntiae, Lachancea thermotolerans, Saccharomyces cerevisiae and Torulaspora delbrueckii were evaluated for tolerance to ethanol, SO₂ and different temperatures, H₂S and biogenic amines production, maximum population size, foam generation, killer and flocculation phenotype.

Conclusions
Most of S. cerevisiae strains tolerated the selective pressures assayed, exhibiting growth at 16% ethanol, 250 ppm SO₂ or 37 °C. Lower tolerance limits were observed for non-Saccharomyces yeasts, with strains of C. zemplinina, I. orientalis and T. delbrueckii growing well at concentrations of 14% ethanol or 200 ppm SO₂. Most strains did not produce histamine, tyramine, putrescine, cadaverine phenylethylamine, tryptamine and isoamylamine, presented low foaming, no-flocculation or killer phenotype and were resistant to killer toxins. Substantial intra- and inter-species differences were observed in H₂S production. Present results highlight the phenotypic diversity among indigenous yeasts and reveal strains with prominent oenological traits that could be exploited in local wine production.

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DEVELOPMENT AND EVALUATION OF YOGHURT FLAVOURED WITH TAMARIND (TAMARIDUS INDICA)

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Background

Yoghurt is produced by bacterial fermentation of milk sugar (lactose), using Streptococcus thermophilus and Lactobacillus bulgaricus, producing lactic acid give yoghurt its texture and its characteristic tang. To modify certain properties of yoghurt, the fermented milk could be blended with ingredients fruits (for flavor and color) and sucrose or aspartame (sweetness). Exotic natural or artificial flavors to boost yoghurt like vanilla, orange, pineapple, strawberry, raspberry, cinnamon among others. However, tropical underutilized flavours like tamarind (Tamarindus indica) have not been exploited.

Objectives

To develop and evaluate yoghurt flavored with Tamarind chemically, microbiologically and organoleptically.

Methods

Tamarind fruit was pulped to juice and treated in three ways (sweetened with sugar, honey or without sweetener/ Skimmed milk and water are homogenized, pasteurized for 15 mins at 85°C, and cooled to 42 - 44°C. The starter culture inoculated, mixed and incubated for 12 hour fermentation, flavoured with graded levels (0, 2, 4, 6, 8, 10, 20, 30, 40, 50 %.) of tamarind juice and stored for 0 to 21 days with weekly monitoring. The sample was then subjected to proximate, micro-nutrient, microbiological and sensory analysis.

Conclusions

Micro-nutrient content (Mg, Na, K, C and Vitamin C) increased with the addition of the fruit juice. Total solids increased with increase concentration of the sweetened fruit flavour in yoghurt. There was no mould or coliform growth but the total viable count ranged from $6.3 \times 10^2$ to $2.34x 10^7$ cfu/ml after 21 days. The sugar sweetened samples were most acceptable over the honey and fruit juice without sweetener.
HIGH-THROUGHPUT SEQUENCING REVEALS BIOGENIC AMINE FORMING COMMUNITIES IN A VARIETY OF ARTISANAL CHEESES

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Background

Biogenic amines are nitrogenous compounds with biological activity produced, primarily, by a variety of lactic acid bacteria. Biogenic amines, such as histamine and tyramine, are commonly found in fermented food products. In cheese, high concentrations can result in adverse health effects for the consumer.

Objectives

The aim of this study was to screen a range of artisanal cheeses (n=10) for the presence of microbial populations capable of producing biogenic amines.

Methods

Segments of the histidine (hdc) and tyrosine (tdc) decarboxylase genes were amplified using previously published PCR primer pairs. PCR amplicons were then TOPO-cloned and subjected to Sanger sequencing. In addition, the Ion Torrent PGM sequencer was used to provide a novel, complementary, in depth analysis of amine forming communities.

Conclusions

Analysis of both Sanger and Ion PGM sequence data revealed hdc and tdc positive bacterial populations both within cheeses and across different cheese varieties. Lactobacillus curvatus, Lb. brevis, Enterococcus faecium and E. faecalis were identified as the predominant species capable of producing tyramine while Lb. buchneri, Lb. sakei and Lactococcus lactis were among the species found to harbour histaminogenic potential. High-throughput Ion sequencing alone revealed the presence of sub-dominant genera, including, among others, Staphylococcus saprophyticus and Streptococcus thermophilus. Actual levels of histamine and tyramine in the respective cheeses were quantified by HPLC.
BACTERIOLOGICAL QUALITY OF COMMERCIALY PREPARED FERMENTED OGII (AKAMU) SOLD IN SOME PARTS OF SOUTH EASTERN NIGERIA

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Background
Food poisoning and infection by bacteria are of public health significance to both developing and developed countries.

Objectives
This study compared the bacteriological qualities of commercially and laboratory based fermented ogi (akamu) prepared from maize.

Methods
Samples of ogi (akamu) prepared from white and yellow variety of maize sold in Uturu and Okigwe were analyzed together with the laboratory prepared ogi for microbial quality using the standard microbiological methods. The analyses showed that both white and yellow variety had total bacterial counts (cfu/g) of $4.0 \times 10^7$ and $3.9 \times 10^7$ for the laboratory prepared ogi while the commercial ogi had $5.2 \times 10^7$ and $4.9 \times 10^7$, $4.9 \times 10^7$ and $4.5 \times 10^7$, $5.4 \times 10^7$ and $5.0 \times 10^7$ for Eke-Okigwe, Up-gate and Nkwo-Achara market respectively. The Staphylococcal counts ranged from $2.0 \times 10^2$ to $5.0 \times 10^2$ and $1.0 \times 10^2$ to $4.0 \times 10^2$ for the white and yellow variety from the different markets while Staphylococcal growth was not recorded on the laboratory prepared ogi. The laboratory prepared ogi had no Coliform growth while the commercially prepared ogi had counts of $0.5 \times 10^3$ to $1.6 \times 10^3$ for white variety and $0.3 \times 10^3$ to $1.1 \times 10^3$ for yellow variety respectively. Lactic acid bacterial count of $3.5 \times 10^6$ and $3.0 \times 10^6$ was recorded for the laboratory ogi while the commercially prepared ogi ranged from $3.2 \times 10^6$ to $4.2 \times 10^6$ (white variety) and $3.0 \times 10^6$ to $3.9 \times 10^6$ (yellow).

Conclusions
There are chances of contracting food borne diseases from commercially prepared ogi. Therefore, there is the need for sanitary measures in the production of fermented cereals so as to minimize the rate of food borne pathogens during processing and storage.
ISOLATION AND MYCOTOXIGENIC ACTIVITIES OF SOME MYCOFLORA OF WEST AFRICAN CHEESE
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Background
Cheese (wara) is a local dairy product produced and consumed in many African countries. It is produced locally by the Fulanis who use Calotropis procera leaves as coagulating agent for the milk.

Objectives
The isolation and identification of the fungi species present in wara and their ability to produce mycotoxins was carried out to determine the safety level associated with consumption of the cheese.

Methods
Mycotoxin production was determined with the use of the enzyme-linked immunosorbent serological assay (ELISA) technique

Conclusions
Some of the isolated mycoflora were found to be mycotoxigenic producing aflatoxins at various levels.
INVITRO PROTEIN DIGESTIBILITY OF FERMENTED BAMBARA NUT (VOANDZEIA SUBTERRANEAN L. THOUARS) USING DIFFERENT SPECIES OF RHIZOPUS

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Background
Bambara nut as legume contains sufficient quantities of carbohydrate and protein high in lysine and methionine. Processing methods, such as soaking, cooking or fermentation can improve the quality of legume protein.

Objectives
Therefore, this research considered the effect of fermentation on in vitro protein digestibility of bambara nut

Methods
The bambara nut was obtained from a local market in Ogbomoso, Nigeria. Different species of Rhizopus for the fermentation were obtained from Ladoke Akintola University of Technology, Ogbomoso, and the Institute of Agricultural Research and Training, Ibadan to produce fermented bambara nut flour. The In vitro protein digestibility of the flour sample was consequently evaluated.

Conclusions
Fermentation process using Rhizopus species alone, and with their combinations showed improvement in the nutritive value of the fermented bambara nut in relation to in-vitro protein digestibility, and the highest rate of protein digestibility was obtained using R. oligosporus alone
Background
Various Lactobacillus species are responsible for the production of fermented food products. Prior to the use of a certain isolate for industrial purpose, it is advisable to classify it at least on species level. Nevertheless, closely related species are not easy to distinguish but molecular methods might be able to reveal the differences even on subspecies level.

Objectives
The aim of our experiments was to verify the usability of several molecular methods in discrimination of Lactobacillus species and compare the efficiency of these tools.

Methods
DNA of lactobacilli were subjected to investigations by PCR (general bacterial and species-specific primers) and qPCR (including newly developed primers and High Resolution Melting - HRM), RFLP and sequencing in order to compare the efficiency of these methods in separation of species.

Conclusions
The HRM had proved its power as a one-step tool for preliminary classification of lactobacilli. With this method it might be much easier to separate closely related species, though, additional techniques, e.g. traditional PCR (species-specific primers), sequencing and RFLP pattern analysis might be necessary to use in order to confirm the results.
GENOMICS OF STREPTOCOCCUS MACEDONICUS: MOVING FROM PATHOGENICITY TO ADAPTATION TO THE DAIRY ENVIRONMENT
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Background
Lactic acid bacteria (LAB) constitute a significant group of microorganisms for food fermentations and for human health. The *Streptococcus bovis/Streptococcus equinus* complex (SBSEC) within LAB includes members that have been implicated in human disease, but are also found in foods.

Objectives
In this study we compared the three available genome sequences of *S. macedonicus* strains isolated from dairy products. Only one strain has its genome complete sequenced and previous analysis showed diminished pathogenic potential and adaptation to the milk environment. Here we present the *in silico* analysis of these strains, in order to better understand the *S. macedonicus* species.

Methods
Chromosomal maps were constructed using DNAPlotter and whole genome sequence alignments were performed by progressiveMAUVE and Webact in order to visualize conserved genomic regions or chromosomal rearrangements. Genomic islands were identified and visualized by IslandViewer, potential bacteriocins were predicted by BAGEL3 and CRISPRs were analyzed by the tools available in the CRISPRcompar web-service.

Conclusions
The analysis revealed that *S. macedonicus* strains have lost genes involved in the catabolism of complex plant carbohydrates, in the adhesion to the host’s cells and in haemolysis that are present in pathogenic SBSEC. In addition, *S. macedonicus* carries two lactose operons and a proteolytic system characteristic of dairy LAB. Our whole genome analysis of *S. macedonicus* shows adaptation traits to the nutrient-rich milk environment.
Background
Brochothrix thermosphacta is considered one of the predominant food spoilage organisms in meat. Atmospheric cold plasma technology (ACP) is proposed as a potential technology for elimination of microbial contamination in food products.

Objectives
The aim of the study was to investigate the antimicrobial efficacy of dielectric barrier discharge atmospheric cold plasma (DBD ACP) for inactivation of B. thermosphacta.

Methods
A large gap DBD ACP system was used to investigate the inactivation of B. thermosphacta (10⁷-8 CFU/ml) in PBS, liquid meat model medium (beef extract broth) and raw lamb chops. Samples were placed inside sealed rigid polypropylene containers and were treated with cold plasma using modified atmosphere as the inducer gas. The effect of ACP critical control parameters, such as treatment time, voltage level, media composition and post treatment temperature storage conditions against both planktonic bacteria and biofilms were evaluated. The surviving bacterial populations were estimated by colony count assay.

Conclusions
ACP showed substantial reductions of bacteria depending on the growth media. ACP treatment (80kV) for 30s completely inactivated the bacterial population in PBS, while 5 min treatment showed 2 Log unit bacterial reduction with the meat model medium. The antimicrobial efficacy of plasma against bacteria on lamb chops showed a reduction around 0.5 log units over a 10 day storage period. The results indicate ACP could be effective for inactivation of B. thermosphacta if treated with sufficient treatment periods providing a prolonged shelf life for meats and meat products. Further studies will investigate the effect of extended ACP treatment times for complete inactivation.
LARGE-SCALE GENE DELETION IN LACTOCOCCUS LACTIS USING THE CRE-LOXP RECOMBINATION SYSTEM

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Background
Minimum genome factories (MGFs) can be defined as recombinant strains whose metabolic pathways have been optimized for special applications. The plasticity of genomes gives us hints that it is probable to streamline physiological pathways through genome rearrangements as strains possess a number of non-essential genes. Lactic acid bacteria (LAB) are low G-C content Gram-positive bacteria and have relatively small genome. Lactococcus lactis could be served as the starting platform for value-added metabolites biosynthesis. Construction of a set of chassis cells provides the opportunity to obtain specific strains with desired performance.

Objectives
To delete the large-scale genes in Lactococcus lactis and construct the set of chassis cells for further synthetic biology research.

Methods
The Cre-loxP recombination system was used to construct a large scale region (19.0 kb) deficient strain N8ΔL1. Two loxP sites were integrated into the genome at target sites, and the large scale region (L1) was replaced by a cat-cassette. The recombination between two loxP sites was achieved when the Cre recombinase was expressed by constructed pNZTS-Cre plasmid, and then the L1-null strains were isolated. Despite the loss of 19.0-kb regions, the strain N8ΔL1 still exhibited normal growth and similar growth profile was also observed compared to the parental strain.

Conclusions
The Cre-loxP recombination system can be used for large-scale genome deletion and generate sequential chromosomal modification mutants of L. lactis efficiently. Based on the precision of its deletion, further genome streamlining and advance the regulatory network study about synthetic biology could be easily achieved.
EFFECT OF RESPIRATIVE METABOLISM OF LACTOBACILLUS CASEI ON MODEL SOURDOUGH FERMENTATION

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Background

*Lactobacillus casei* is a lactic acid bacterium (LAB) used in the production of many fermented food and feed products. Furthermore, this specie comprises strains commercially exploited as probiotic cultures. Thus, novel strategies to improve their adaptation to adverse conditions could enhance their safety, functional and technological properties. Recently, it has been demonstrated that respiratory metabolism results in this species in the expression of a phenotype with enhanced technological and stress response properties (increase in biomass, synthesis of antioxidant enzymes, robustness to stress conditions).

Objectives

The aim of the study was evaluating the impact of respirative/anaerobic metabolism of *L. casei* N87 strain on model sourdough fermentation.

Methods

The strain *L. casei* N87, grown both in anaerobic and respirative conditions, was inoculated as starter in wheat flour in combination with a commercial baker’s yeast culture. After 0, 6 and 24 h of fermentation LAB count, pH and Titratable acidity values, free amino acid (FAA), volatile compounds (VOCs) (by SPME-GC-MS), antioxidant activity (by DPPH assay) and albumins/globulins, gliadins and glutenins protein fractions (by SDS-PAGE) were evaluated.

Conclusions

A major increase in biomass for the strain *L. casei* N87 grown in respirative condition was registered after 6h of fermentation. Results obtained by FAA and VOCs evaluation indicated that respirative metabolism of *L. casei* N87 influences the flavour of sourdough. Furthermore, SDS-PAGE highlighted a different wheat protein degradation in anaerobic/respirative cultures. In conclusion, respirative *L. casei* N87 performs well in sourdough definition and a better explanation of its role in sourdough production is considered opportune.
Background

Chenopodium quinoa and Amaranthus spp. are considered pseudocereals characterized by high nutritional and functional values which are associated with the quality and quantity of their proteins, fats and antioxidant compounds. In the recent years, these pseudocereals have attracted the interest of research for their exploitation in the production of gluten-free products and food products with improved nutritional and health benefits.

Objectives

To characterize quinoa and amaranth seeds (produced in Campania Region) for microbiological, technological and safety aspects with the aim to use them in the production of “functional foods”

Methods

Three samples of Amaranthus (accessions A1, A2 and A3) and three samples of Chenopodium quinoa seeds (2 Titicaca and 1 Puno varieties) were obtained by experimental fields of Vitulazio (Campania Region). Total mesophilic count, lactic acid bacteria (LAB), yeasts, moulds, Enterobacteriaceae, enterococci, total and faecal coliforms were assessed by standard pour-plate technique by using selective media. Total polyphenols content was determined using a method based on Folin-Ciocalteu reagent, while antioxidant activity was carried out spectrophotometrically using DPPH (2,2-diphenyl-1-picrylhydrazyl) radical scavenging.

Conclusions

Results evidenced a high LAB concentration in quinoa and amaranth seeds ranged between 2.5 and 7.5 log CFU/g and between 2.8 and 5.0 log CFU/g, respectively. Quinoa seeds showed higher level of polyphenol content compared with amaranth, whereas no significant differences were observed in antioxidant activity, registered in high concentration for both the seeds. In conclusion, the characteristics of these seeds are highly desirable for the production of both gluten-free products and food with high nutritional value.
UNDERSTANDING DIVERSITY OF BACTERIAL COMMUNITIES FROM POULTRY MEAT TO IMPROVE FOOD QUALITY AND SAFETY

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Background

Meat products can host variable microbial communities according to seasonal changes and production processes. These may encompass pathogenic or spoilage bacteria which must be controlled to ensure safety and quality of the products.

Objectives

Our aim was to describe the microbial community of chicken legs packaged under modified atmosphere.

Methods

Whole bacterial populations were collected and stored frozen at -80°C. Their ability to regrow on meat was checked. Bacterial diversity was determined by cultural methods (23 samples) and by 16S rDNA pyrosequencing (10 samples).

Conclusions

Total viable counts varied with samples ($10^3$ - $10^8$ CFU/g) and plating methods showed that lactic acid bacteria, *Brochothrix thermosphacta*, and *Pseudomonas* spp. were the main bacterial. The characterization of the bacterial diversity by 16S rDNA pyrosequencing confirmed the presence of *B. thermosphacta*, and revealed that *Pseudomonas* was mainly represented by *P. extremiaustralis* and *P. cedrina*. As well the main LAB species were *Carnobacterium* and *Shewanella* species. The predominance of *Pseudomonas* was correlated to meat packaging under high oxygen concentration, except when *B. thermosphacta* was dominant, suggesting a competition between these species. No clear cut correlation could be observed between farming practices or meat processing and bacterial communities. However, our results showed similar microbial profiles of samples issued from the same slaughterhouse suggesting the main contamination may occur during this processing. The frozen communities will now be used to evaluate the abiotic and biotic (e.g. the *Brochothrix-Pseudomonas* competition) factors in reproducible challenge tests on meat to finally improve the storage life-time.
Background
Fungal growth is the main cause of fruit decay, and is usually controlled by the application of synthetic fungicides, however, currently their use has been restricted. Therefore, it is urgent to find alternative antifungal substances. Phenolic compounds are secondary metabolites of plants and naturally present in fruit which have been associated with antimicrobial effect. Citrus peel contains high amounts of flavonoids and is the main byproduct of processing industry.

Objectives
Then, the aim was to investigate the inhibitory activity of phenolic extracts obtained from citrus peel against fungal fruit spoilage.

Methods
The antifungal activity of extracts from orange and lemon peels were tested in-vitro against 6 yeast strains (Cryptococcus spp., Torulaspora spp, Aerobasidium pullulans, Rhodoturula spp., Hanseniaspora uvarum, Meyerozyma caribbica) and 8 molds (Botrytis cinerea, Monilia laxa, Alternaria spp., Penicillium glabrum, Penicillium expansum, Penicillium corylophilum, Cladosporium uredinicola, Cladosporium cladosporioides), by following the ability to grow in a medium containing different concentrations of extracts.

Conclusions
Overall, the inhibitory effect of orange extracts (OE) against yeast strains was around 90% at concentrations higher than 0.75 g/L, whereas the effect of lemon extracts was more strain dependent and powerful, with values closed to 90% against four strains at 0.15 g/L. On the other hand, although inhibitory effect against molds was lower, OE presented a remarkable activity against B. cinerea, M. laxa and Alternaria spp. and moderated against Penicillium and Cladosporium strains. In conclusion, these results suggest the possibility of using citrus polyphenolic extracts as a safer alternative to synthetic fungicides to control postharvest decay of fruit.
LISTERIA MONOCYTOGENES OF SEQUENCE TYPE 121 HARBOR SPECIFIC ADAPTATIONS SUPPORTING PERSISTENCE IN FOOD PRODUCTION PLANTS.

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Background

The foodborne pathogen Listeria monocytogenes is able to survive for months and even years in food production environments. Among a great strain diversity particularly strains belonging to sequence type (ST)121 are persistent in food production plants.

Objectives

To elucidate the molecular determinants responsible for persistence of L. monocytogenes.

Methods

We analysed the genomes of two L. monocytogenes ST121 strains, which persisted for up to eight years in food production plants in Ireland and Austria. Additionally we characterized two ST121 strain-specific genetic features: Tn6188 and lin0464/lin0465.

Conclusions

All ST121 genomes are highly similar and show a tremendously high degree of conservation among prophages and particularly among their plasmids, which are usually variable parts of genetic information in bacteria. This remarkably high level of conservation suggests a strong selective pressure.

In addition, all ST121 strains share adaptations related to persistence in food production environments such as the presence of Tn6188, a transposon responsible for increased tolerance against quaternary ammonium compounds, and the presence of homologues of the L. innocua genes lin0464 and lin0465, a transcriptional regulator and a putative pfpl protease. Deletion of lin0465 resulted in reduced survival under oxidative and alkaline conditions suggesting a role in stress response. Furthermore all ST121 strains reveal a yet undescribed insertion harboring recombination hotspot (RHS) repeat proteins, which are most likely involved in...
competition against other bacteria. In conclusion we show that \textit{L. monocytogenes} ST121 strains are highly similar to each other harboring conserved regions which provide fitness adaptations to survival in food production environments.
NOVEL VIRULENCE FEATURES OF LISTERIA MONOCYTOGENES ISOLATED FROM FOOD SAMPLES SOLD AT A ROMANIAN BLACK MARKET.

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Background

Listeria monocytogenes is a facultative intracellular foodborne pathogen responsible for listeriosis. Investigated neglected exogenous routes of transmission of foodborne pathogens into the EU, we have isolated 15 L. monocytogenes strains in food products illegally sold at a Romanian black market.

Objectives

The aim of this study was to characterize the subtype and the virulence features of these L. monocytogenes strains.

Methods

We determined invasion efficiency and intracellular growth in human intestinal epithelial Caco2 and macrophage-like THP1 cells; and analysed the sequence of three main virulence factors: PrfA, internalin A (InlA) and listeriolysin O (LLO).

Conclusions

Multilocus sequence typing revealed that these L. monocytogenes strains belong to six different sequence types (ST). In vitro virulence assays showed a high strain variability regarding the invasion efficiency in Caco2 cells and the intracellular growth rate in both cell types. In parallel we revealed a high diversity in the InlA and LLO amino acid sequences, however strains belonging to the same ST harbour identical sequences. We detected in total 30 different amino acid substitutions, resulting in seven different InlA variants, two of which have not yet been described. All ST121 and ST9 strains, harbouring a premature stop codon resulting in truncated InlA, were unable to invade Caco2 cells. In addition the number of LLO mutations correlated negatively with intracellular growth in both cell types; and all ST155 strains showed no proliferation inside macrophages. In conclusion, we show a high diversity in the amino acid sequence of main virulence factors in L. monocytogenes resulting in distinct virulence profiles.
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Background

Interactions of bacteria living in the same microenvironment influence the behaviour of pathogens like Listeria monocytogenes. Therefore inter-strain competition between different L. monocytogenes strains could influence fitness and pathogenicity.

Objectives

This study investigated the impact of co-culture on growth and in vitro virulence of L. monocytogenes strains and the effect of cell-contact on the observed growth and virulence competition.

Methods

Growth of eight L. monocytogenes strains was determined in single and two-strain mixed cultures in tryptic soy broth at 10°C for 10 days. The effect of strain competition (24h, 10°C) on invasion efficiency and intracellular growth was investigated using human intestinal epithelial Caco2 cells. Two selected strains were inoculated singly, mixed or separated by 0.4μm PET-membrane to analyzed whether the observed effects are cell-contact dependent.

Conclusions

Significant differences in growth between single and mixed cultures were observed for certain strains e.g. ScottA and 6179 showed reduced growth rates when co-cultivated with other strains. Cell-contact was essential for growth competition.

High virulent strain e.g. PL25 showed either increased or unchanged invasion efficiency when co-cultured with low virulent strains, whereas the invasiveness of low virulent strains like ScottA and 6179 was attenuated. These results suggest a competitive advantage for the strains displaying higher invasion efficiency. In parallel to growth the effect of strain competition on virulence is cell-contact dependent.

In conclusion competition between L. monocytogenes strains has a strain-dependent effect on fitness and virulence.
NEW INSIGHTS OF A NATURAL GUM: FOCUSING ON GUT ECOLOGY.

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Background

Gum odina obtained from bark of Odina wodier (Anacardiaceae) is evaluated as prebiotic in this study. The natural gum (polysaccharide) is chemically galactopyranose associated with arabinose side chain.¹

Objectives

The use of prebiotic to fortify gut flora is current aspect of research for the control and prevention of colon cancer as well as boosting of immune system.

Methods

In vivo prebiotic potential of gum odina was performed on Swiss albino mice by deliberate challenging of S.typhi 62 focusing on immunological parameters² and in vitro studies was investigated by Simulator for Human Intestinal Microbial Ecosystem (SHIME).³

Conclusions

The non-degradability of gum odina by simulated gastric fluid (SGF) and simulated intestinal Fluid (SIF) attributes to one of the major characteristics of prebiotics i.e. indigestible carbohydrate.Microbial analysis of SHIME studies shows increase in colonization of various probiotic organisms such as Lactobacillus sp. and Bifidobacterium sp. upon administration of odina. In vivo study revealed natural gum selectively stimulates probiotic strains and eliminates enteric pathogens and also boosts immune system by increasing slgA in gut. Moreover increase in levels of IFN γ signifies additional protection against various pathogen induced primary and secondary infections.This gum is also believed to possess colon carcinoma preventing properties as it liberates short chain fatty acids (SCFA) upon fermentation which we are planning to focus in our next studies.
USE OF BACTERIAPHAGE-BASED SOLUTION AND ELECTROLYZED OXIDIZING WATER AS ALTERNATIVE SANITIZING SOLUTIONS TO INACTIVATE LISTERIA MONOCYTOGENES ON FOOD CONTACT SURFACES FOR THE SEAFOOD PROCESSING FACILITIES

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Background
Listeria monocytogenes is an important foodborne pathogen that can cause serious public health-related issues worldwide. L. monocytogenes could be isolated from food contact surfaces (FCS) from the seafood processing facilities, e.g., slicers, utensils, knives, sieves, conveyors, towels, employer’s gloves, which could then link to the occurrence of cross-contamination of seafood products.

Objectives
Effective sanitizing agents are thus needed to reduce the incidence of Listeria contamination on FCS. In this study.

Methods
Phage-based solution LP-124 (LPS; broad host-range listeriaplague) and electrolyzed oxidizing water (EOW; pH 2.5 and ORP 1150 mV) were evaluated for the ability to inactivate L. monocytogenes cells that were artificially contaminated on four different FCSs (plastic cutting board, towel, stainless steel spoon, and sieve) at 4 logCFU/cm² on a given FCS.

Conclusions
Results showed that EOW and LPS led to significantly (p<0.05) decrease of L. monocytogenes counts, representing about 2 logCFU/cm² on a given FCS, [KV1] compared with the counts recovered from washing with water. L. monocytogenes counts were not detected in the previously used EOW solution stored at 4 °C during 7 days after used, while about 2.8 logCFU/ml was observed in used phage-based solution after 7 days of storage. Both EOW and LPS can effectively inactive L. monocytogenes on different food contact surfaces, and can further be used as alternative sanitizing solution for the seafood processing facilities.
Background
Parasitic infection is highly prevalent throughout the developing countries of the world. Food handlers are potential source of infection of many intestinal parasites and other enteropathogenic infections as well

Objectives
The aim of this study was to determine the prevalence of intestinal parasites carriers among food handlers attending the public healthcenter laboratory in Sari, northern Iran for annual check-up

Methods
The study was performed from August 2011 through February 2012. Stool samples were collected from 1041 male and female food handlers of different Jobs aged between 18 to 63 years and were examined following standard procedures. Sociodemographic, environmental and behavioral data of the food handlers were recorded in a separate questionnaire. Intestinal parasites were found in 161 (15.5%) of the studied samples. Seven species of protozoan orhelminth infections were detected. Most of the participants were infected with Giardia lamblia (53.9%) followed by Blastocystis hominis (18%), Entamoeba coli (15.5%), Entamoeba histolytica/dispar (5.5%), Cryptosporidium sp. (3.1%), Iodamoeba butschlii (3.1%) and Hymenolepis nana (1.9%) as an only helminth infection.

Conclusions
The finding emphasized that food handlers with different pathogenic organisms may predispose significant risk on the consumers. Routine screening and treatment of food handlers is a proper tool in preventing the food-borne infections.
Background

Accurate identification and assessment of antibiotic resistance of bifidobacteria are essential to secure quality and safety of probiotics and food products, containing these microorganisms.

Objectives

The objective of the work was identification and evaluation of antibiotic resistance of 25 probiotic strains of bifidobacteria, isolated from feces of healthy adults, therapeutic preparations and dairy products, marketed in Belarus.

Methods

Sequencing of 16S rDNA and transaldolase gene, MALDI-TOF MS protein profiling, biochemical testing, analysis of fatty acid methyl esters (FAMEs) were used for bifidobacteria identification. PCR-analysis, disc-diffusion and broth micro-dilution methods were applied for determination of antibiotic resistance of bifidobacteria.

Conclusions

Results of molecular-genetic identification of bifidobacteria indicated that 7 strains belong to species *Bifidobacterium animalis* subsp. *lactis*, 7 strains – *Bifidobacterium longum*, 5 strains – *Bifidobacterium bifidum*, 5 strains – *Bifidobacterium adolescentis*, 1 strain – *Bifidobacterium breve*. MALDI-TOF MS analysis confirmed taxonomic affiliation of tested strains and proved to be a rapid and reproducible technique for bifidobacteria identification. Protein biomarkers suitable for species and strain discrimination were defined. Both biochemical testing, based on evaluation of enzymatic activity and carbohydrates fermentation, and FAMEs analysis did not provide accurate species identification of bifidobacteria, but were useful for strain differentiation and characterization. Tested strains of bifidobacteria were susceptible to chloramphenicol (100%), penicillin G (96%), amoxicillin (96%), ampicillin (88%), streptomycin (80%), vancomycin (80%), gentamycin (76%), resistant to erythromycin (40%), clindamycin (40%), tetracycline (52%), and harbor genes *tet(W)* (36%), *tet(M)* (16%), *erm(X)* (32%), *erm(B)* (8%). Gene *tet(W)* was detected in all *B. animalis* subsp. *lactis* strains, gene *erm(X)* – all *B. longum* strains.
INFLUENCE OF TEMPERATURE AND INOCULUM SIZE ON SPORES GERMINATION PERCENT OF CLOSTRIDIUM ALGIDICARNIS AND CLOSTRIDIUM ESTERTHETICUM
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Background
'Blown pack' spoilage causes high economic losses for Brazilian meat industry.

Objectives
This research aimed to determine the influence of temperature and inoculum size on spore germination percent of *C.algidicarnis*, isolated from spoiled vacuum meat and *C.estertheticum* DSM8809, both recognized for their package blowing ability.

Methods
A central composite design - 2 factors x five levels - was applied: storage temperature (-2,3,7,11, 15°C) and inoculum size(10¹,10²,10³,10⁴ and 10⁵ spores/mL, previously prepared). Experiments were conducted using RCM media, in anaerobic conditions. Counts were performed daily, during 30 days, using NMP method. Spores germination percent (SGP) for each day was calculated as: SGP (%)=(count of germinated cells/initial spore inoculum)*100).

Conclusions
In all assays, for both microorganisms, germination was observed. *C.estertheticum* germinated faster than *C.algidicarnis* at lower temperatures: at 3°C/10⁵ spores of inoculum, after 1 day all spores, initially inoculated, were germinated and for *C.algidicarnis*, 100% germination was reached after 4 days. In addition, at 11°C/10³ sp/mL, germination of *C.estertheticum* was slower than *C.algidicarnis*, achieving 100% after 11 days. For inoculum of 10⁵ spores/mL, *C.algidicarnis* reached high germination percent quickly at 15°C (52.4% after 2 days) while *C.estertheticum*, at 7°C (39% at 2 days). Increasing initial inoculum size from 10² to 10⁵ spores/mL/3°C, increased the time for total germination from 4 to 10 days for *C.algidicarnis* and from 1 to 15 days for *C.estertheticum*. These results clearly shows that tested temperatures were no enough to inhibit spores germination of both microorganisms, even if initial population is 10⁵ spores/ml, emphasizing the need to avoid the vacuum packed red meat clostridial spore contamination before packaging.
THE EFFECTS OF ANTIBIOTICS AND DIFFERENT HOSTILE CONDITIONS ON SURVIVAL AND ADHESION OF PROBIOTIC LACTIC ACID BACTERIA IN VITRO IN MODEL OF HUMAN GASTROINTESTINAL TRACT

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Background

The probiotic microorganisms must be able to survive during passing through the hostile conditions of the gastrointestinal tract and have an ability to adhere to the intestinal epithelial cells. Nowadays there are many physical and chemical factors affecting these abilities of probiotics.

Objectives

In this work was studied the effects of electromagnetic irradiation at the frequencies 51.8 and 53 GHz (1h) and antibiotic ceftazidime (20 µM) on survival of \textit{Lactobacillus acidophilus} VKMB-1660 \textit{in vitro} in model of gastrointestinal tract.

Methods

Bacteria developed in artificial gastric juice (treatment time 20 min) with pepsin (0.3\%) and different pHs (2.0, 3.0, 4.0, 5.0) and artificial intestine juice (treatment time 1 h) with bile salts (0.45\%) and pancreatin (0.1\%) at pH 8.0. Moreover it was studied the effects of antibiotics and different pHs (5.5, 6.5 and 8.0) on adhesiveness of bacteria.

Conclusions

The results obtained shown that the colony forming unit number significantly decreased, in artificial gastric juice than in intestine juice in comparison with untreated control samples. The irradiation at both frequencies and ceftazidime significantly decreased the viability of bacterial samples in both gastric and intestine juices. Moreover, ceftazidime suppressed the adhesion ability of bacteria at all pHs used in comparison with untreated samples and adhesiveness was much weaker at pHs 5.5 and 8.0 than at pH 6.5.

Thus, lactic acid bacteria \textit{L. acidophilus} can survive in viable state in the gastrointestinal tract even after irradiation and treatment with antibiotics, but antibiotics can significantly decrease an adhesion of lactic acid bacteria to intestine mucosa.
HYGIENE BARRIERS IN FOOD PROCESSING FACILITIES MAY SERVE AS “TROJAN HORSES” FOR L. MONOCYTOGENES

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Background
Generally, hygiene sluices are efficient barriers against the transmission of spoilage and foodborne pathogen bacteria.

Objectives
The aim of the study was to determine the efficiency of hygienesluices on the reduction of hygiene indicator bacteria and zoonotic agents in food processing companies

Methods
We included five food processing facilities located in Upper- and Lower Austria. Samples were taken before and after sanitation comprising personal fingerprints, swab samples from shoes and the hygiene lock, and water residues and investigated according to microbiological standard methods. L. monocytogenes were subtyped by Serogroup PCR and pulsed-field gelelectrophoresis (PFGE).

Conclusions
One shoe sample each was found positive for E. coli (rfbE positive) and Salmonella spp. after sanitation in slaughterhouse B and RTE-food producer C, respectively.

Coliform bacteria and L. monocytogenes on shoes could not be reduced after passing the hygiene locks. In detail, 29% of the shoe samples were detected L. monocytogenes positive before and after the foot bath. Most of L. monocytogenes isolates were found on shoes of two slaughterhouses and one RTE-food company working staff. Subtyping revealed that the slaughterhouse A harbored L. monocytogenes serogroup 1/2a, 3a and 4b, 4d, 4e (n=21) resulting in five PFGE pulsotypes. In slaughterhouse B L. monocytogenes serogroup 1/2a, 3a, 1/2c, 3c and 4b, 4d, 4e (n=11) was present, representing five PFGE pulsotypes. In one RTE-food producing company genetic lineage II (1/2a, 3a; 1/2c, 3c; n=10) were over represented, including persistent strains. Hygiene locks, if not efficiently working, could be a reservoir for human pathogen bacteria and widely distributed L. monocytogenes clones.
DIVERSITY AND ANTI-LISTERIAL POTENTIAL OF BACTERIAL FLORA ON HARD-CHEESE RINDS

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Background

Vorarlberger Mountain cheese Protected Designation of Origin (PDO) owes its special characteristics from the raw milk of extensively farmed cows grazing wild herbs on alpine pastures during the summer months.

Objectives

The aim of the study was to determine the diversity of bacterial flora on hard-cheese rinds isolated from artisan Vorarlberger Mountain cheese. A further goal was to test their antilisterial capability.

Methods

The bacterial rind flora of seven cheese samples, including the variety of three ripening cellars (A, B, C) and two different ripening stadiums, were characterized. Furthermore, five cheeses were investigated during a six month period, to determine the variations in bacterial groups on cheese surfaces due to failures or instabilities in cheese-curing. Bacterial cheese flora isolates were enumerated and confirmed applying standard agar isolation methods and 16S rRNA sequencing. The antilisterial potential of 56 cheese rind bacteria was tested in an agar-overlay model against \textit{Listeria} spp. Additionally, cheese rind bacteria with antilisterial potential and a brine-adapted \textit{L. monocytogenes} isolate were introduced in a challenge experiment on cheese during ripening.

Conclusions

The majority of isolates comprised following genera and species: \textit{Brevibacterium} spp. (30.9%), \textit{Staphylococcus equorum} (20.9%), \textit{Corynebacterium casei/variabilis} (11.4%) and \textit{Brachybacterium alimentarium} (10.9%). Antilisterial potential in the agar model was observed from 30.4% of test strains. In the challenge experiment higher \textit{L. monocytogenes} spike levels ($10^4$ cfu) were positive after enumeration and enrichment until the end of smearing at day 14. Applying a brine with higher salt concentration (>20%) resulted in higher growth rates during ripening (average $3.98 \times 10^3$ cfu/g).
MICROBIAL QUALITY OF EDIBLE INSECTS

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Background

Current animal protein sources will not be sufficient for the growing population. Therefore, alternative solutions need to be found. Edible insects are considered as a sustainable and nutritionally equivalent alternative to meat. Nowadays, there is a lack of profound insight into the microbiological quality of edible insects.

Objectives

The purpose of this research was to investigate the microflora of edible insects in a quantitative and a qualitative way, in particular of the yellow mealworm (Tenebrio molitor) and the grasshopper (Locusta migratoria migratorioides).

Methods

Living insects for human consumption were purchased from an eco-shop (Belgium). Both insect species were subjected to culture-dependent analyses (classical microbial counts) to determine the microbial load (total aerobic count (TAC), Enterobacteriaceae, lactic acid bacteria (LAB), yeasts and moulds (Y&M) and aerobic bacterial spores). In addition, culture-independent analyses (454 pyrosequencing) were performed to gain insight into the bacterial community composition.

Conclusions

Average microbial counts (3 batches, log cfu/g) in mealworms were 8,0 TAC, 7,3 Enterobacteriaceae, 7,4 LAB and 5,4 Y&M. Lower counts were observed for spore-forming bacteria ranging between <1,0-3,5 log cfu/g. Grasshoppers showed average counts (2 batches, log cfu/g) of 7,2 TAC, 5,6 Enterobacteriaceae, 7,0 LAB, 4,2 Y&M and 4,2 bacterial spores. The culture-independent analyses demonstrated a high bacterial diversity in mealworms and grasshoppers. Most bacteria belonged to the Enterobacteriaceae and LAB, however, the dominant bacterial species differed between both insect species. Hence, raw edible insects contain a high microbial load and diversity, including spoilage organisms and potential pathogens, a decontamination step before consumption is required.
A TEN-YEAR SPOT-SAMPLING (2005-2014) OF FISH, WATER AND SURFACES ALONG THE PRODUCTION CHAIN IN THE NORWEGIAN PELAGIC FISH SECTOR.

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Background

Microorganisms are of great importance in post-harvest spoilage of marine resources. Both quality-reducing and potential pathogenic bacteria may reduce the first-hand value, the export potential, and the reputation of the seafood industry.

Objectives

To examine microbiological conditions of the commercially most important pelagic fish species harvested in Norway, with respect to quality, hygiene and food safety.

Methods

Microbial fish quality was evaluated by heterotrophic plate counts (HPC) and counts of H₂S-producing bacteria, whereas hygienic- and safety conditions were assessed by analysing for faecal indicator organisms, e.g. presumptive E. coli and enterococci, and the pathogenes Listeria monocytogenes, Salmonella and staphylococci. Examined contact points in the production environment at vessels and factories, included surface samples of equipment and water samples associated with fish catch, -storage, and -production. Contact points were examined for HPC, Enterobacteriaceae, presumptive E.coli, enterococci, Salmonella, and L. monocytogenes. For evaluation, an assessment scheme with recommended microbiological limits suitable for fresh pelagic fish products are proposed based on contemporary EU- and Norwegian guidelines.

Conclusions

Quality-, hygienic- and food safety conditions were according to the recommended assessment scheme in all samples from 23 samplings. However, 18 samplings had samples that were not optimal and contamination of fish, the pump nozzle, sift box, sorting chambers, storage tanks, landing tanks and conveyor belts were most often recurring. Two samplings revealed that contamination early in the production chain followed the fish throughout the process, thus an increased focus on clean equipment during capture and storage is highly suggested.
Background

In food processing lines or in complex equipment, microorganisms are exposed to varying hydrodynamic conditions caused by the flow of liquid food, and biofilms grown under a wide distribution of hydrodynamic strengths.

Objectives

The aim of the present work was to investigate the in situ thickness, architecture, and rheological properties of yeast biofilms growing on stainless steel under turbulent flow.

Methods

The yeast species used (Rhodototula mucilaginosa, Candida krusei, Candida kefyr and Candida tropicalis) were isolated from an apple juice industry. Biofilm formation in turbulent flow were performed in a Rotating Disk System (RDS) already described (Brugnoni et al., 2011). Viscoelastic properties of biofilms were determined by small deformation dynamic oscillatory measurements in a Paar Physica rheometer MCR301 (Anton Paar GmbH, Graz, Austria), using parallel plates.

Conclusions

Results show yeasts biofilms formed on stainless steel at Reynolds (Re) numbers ranging from $294,000$ to $1.2 \times 10^6$. These growth phases transform adherent blastospores to well-defined cellular communities. Biofilm formation increases with Re and time. Flow conditions impacted biofilm composition, with a predominance of C. krusei. Under turbulent flow in biofilm thickness increased >100 μm and cell morphology was governed by hyphal structures and rounded cells. Biofilms resulted viscoelastic materials with a solid-like behavior. Rheological values were not significantly affected by flow conditions or growth time. At large deformations their weak structure collapsed beyond a critical strain.

ROLE OF THE COLD SHOCK DOMAIN FAMILY PROTEINS (CSPS) IN NISIN AND BENZALKONIUM CHLORIDE STRESS TOLERANCE OF LISTERIA MONOCYTOGENES

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Background

Listeria monocytogenes is an important foodborne pathogen that causes rare but serious illness (listeriosis) and high mortality in those with diminished immunity. Proteins of the cold shock domain family (Csps) are global gene expression regulators that promote various stress adaptation responses in bacteria. Nisin and Benzalkonium chloride (BC) are widely used antimicrobials for food preservation and disinfection of food processing environments, respectively.

Objectives

To investigate the functional role of Csps in the innate resistance of L. monocytogenes to nisin and BC stresses.

Methods

L. monocytogenes EGDe wild type and csp deletion mutant strains were phenotypically compared under nisin and BC stress in growth assays. Impact of nisin and BC stress on gene expression was determined using quantitative reverse transcription PCR (RT-qPCR).

Conclusions

In absence of Csp functions the sensitivity of L. monocytogenes EGDe to nisin and BC stress exposure was increased. Both growth and survival capacities of the L. monocytogenes EGDe strain lacking Csps were significantly diminished compared to the wild type strain when exposed to nisin and BC stress in BHI cultures. In addition the Csp-lacking EGDe strain also showed increased susceptibility to cell envelope-targeting antibiotics. Induction of csp genes upon exposure to nisin but not BC was also detected using RT-qPCR. A comparative gene expression analysis revealed an altered expression in genes encoding peptidoglycan-binding proteins in absence of Csp functions. Our results suggest that Csp-dependent gene expression regulation
contributes to optimal intrinsic nisin and BC resistance mechanisms in the foodborne pathogen *L. monocytogenes*. 
GENOTYPIC AND PHENOTYPIC CHARACTERISTICS OF LISTERIA MONOCYTOGENES STRAINS ISOLATED DURING 2011-2014 FROM DIFFERENT FOOD MATRICES IN SWITZERLAND

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Background

*Listeria monocytogenes* is a food-borne pathogen of public health and food safety concern responsible for highly invasive disease (listeriosis) and high mortality in those with diminished immunity.

Objectives

The objective of this study was to characterize *L. monocytogenes* strains recovered from different food matrices in Switzerland between 2011 and 2014 with respect to their genotypic and phenotypic properties.

Methods

Genotypes were determined through DNA sequencing (MLST) and PCR (stress survival islet (SSI-1), *qacH* and *brcABC*) methods. BC resistance (BC') was determined based on minimum inhibitory concentrations on MH agar and biofilm formation by the crystal violet staining method.

Conclusions

One hundred and forty two strains isolated from various food products between 2011 and 2014 in Switzerland were examined. The strains comprised serotypes 1/2a (64%), 1/2b (7%), 1/2c (12%), 3c (2%) and 4b (15%). There were 61 MLST sequence types (ST) determined including 24 new ST that had not been previously described. Fifty percent of the strains were SSI-1 PCR positive whereas 31% were negative. Remaining strains either harbored smaller than expected PCR amplicon (15%) or the SSI-1 PCR primers failed (4%) to amplify. BC' was detected in 25 (18%) strains including 20 (80%) and 3 (12%) that possessed *qacH* and *bcrABC* BC' genetic determinants, respectively. Ninety one percent of the strains were (129/142) were classified as weak, 8% (11/142) as moderate and only 1% (2/142) as strong biofilm formers. Our results indicate a high genetic and phenotypic diversity among *L.*
monocytogenes strains that contaminate different types of food products in Switzerland.
Background

Freshwater fishes are commonly used for fish preparations in Latvia and the hygienic status of a lake from there the freshwater fishes were originated may alter microbiological quality of fresh caught fish.

Objectives

The aim of the present study was to detect the microbiological quality of freshly caught freshwater fish in Latvia.

Methods

Altogether, 36 fishes were collected, among them 25, four and seven were European eel (Anguilla anguilla), silver bream (Blicca bjoerkna) and European perch (Perca fluvialitis), respectively. Samples were collected between September and November, 2014 from fishermen from lakes in Latvia. Surface swabs of a 5x5 cm² of freshly caught fish skin were examinated according to ISO methods for total bacterial count (TBC), Enterobacteriaceae, fecal coliforms, psychrotrophic microorganisms, Salmonella spp., Listeria spp. and Yersinia spp.

Conclusions

Counts of TBC, Enterobacteriaceae, fecal coliforms and psychrotrophic microorganisms were found to be significantly lower on eels than on perch and silver bream (p>0.05). Eels in Latvia are introduced to lakes in elder or glass eel stage in the frame of eel breeding program and this could influence the contamination rates of eel microflora as they are mostly not a native part of freshwater environment. Observed significant differences (p>0.05) between the counts of TBC,
Enterobacteriaceae, fecal coliforms and psychrotrophic microorganisms may be explained with the hygienic status of lake of origin of freshwater fish. Absence of Salmonella spp., Listeria spp. and Yersinia spp. in tested samples indicates that they do not share pathogens of public health significance.
Background: The stable and safe supply of plant-derived food is endangered by unusual weather, environmental pollution, and shortages of water, fossil fuel and plant biomass. When leaf vegetables are grown in the open field, their quality and productivity tend to vary with the local climate, weather conditions and soil fertility. On the other hand, when plants are grown in the greenhouse, their quality and productivity are generally improved. Thus, the plant factory is a useful system for providing the food supply. When hydroponic nutrient solution in the plant factory is contaminated by the pathogens, the pathogens will propagate and spread, quickly. And the importance of the disinfection system for hydroponic nutrient solution is pointed out in recent years.

Objectives: The aim of this study was to develop the disinfection system with UVA-LED and estimate the ability of disinfection of nutrient solution.

Methods: We developed a circulating disinfection device including a cylindrical disinfection tank of 1L volume. Escherichia coli strain (ATCC25922) was used as model microorganism for this study. Inactivation ability was determined by a colony-forming assay.

Conclusions: The disinfection system with UVA-LED was able to sterilize the hydroponic nutrient solution in the plant factory. The equation which indicated the relation between log survival ration, fluence of UVA, and energy consumption was lead by the experiments. The disinfection system with UVA-LED was excellent in not using a chemical substance and low damage to crops and hydroponic nutrient solution.
THE CHANGING BACTERIAL POPULATION OF FRESH-CUT ICEBERG
LETTUCE DURING STORAGE

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Background
Minimally processed (fresh-cut) iceberg lettuce is prone to spoilage after being harvested and processed. As the produce is not sterile, bacterial contaminants may proliferate and add to the organoleptic changes.

Objectives
The objective of this study is to describe changes in the bacterial composition of modified atmospheric (MA) packed fresh-cut produce during the storage period at refrigerated temperature.

Methods
Lettuce samples were taken daily during the storage period for microbiological analysis. Plate culturing was used to analyze changes in the total viable count. Application of pyrosequencing of 16S rRNA gene amplicons was used to monitor the changes in the bacterial population. Dedicated quantitative PCR (qPCR) protocols are developed to monitor the dynamics of specific bacterial entities covering Leuconostoc, Lactococcus, and Pseudomonas.

Conclusions
The mesophilic aerobic bacterial counts increased within 8 days from about $10^5$ to $10^8$ colony forming units per gram of lettuce. Pyrosequencing showed at the genus level, Pseudomonas, Leuconostoc, and Lactococcus, being most dominant directly after packaging while Leuconostoc becomes most dominant while storage time is elapsing. Application of these qPCR protocols confirmed the typical change of the bacterial population of MA packed fresh-cut iceberg lettuce towards high dominant numbers of Leuconostoc species at day 8 after packaging. In several experiments, Leuconostoc spp. was found to dominate the bacterial composition at the later time points during storage, suggesting a role for Leuconostoc-derived metabolites in lettuce quality attributes.
ASSOCIATION BETWEEN VIRULENCE PROFILE, SEQUENCE TYPE AND SEROTYPE OF SALMONELLA ENTERICA STRAINS ISOLATED FROM FOODSTUFFS IN MICHOACÁN, MEXICO
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Background

The products of the genes invA, rmbA, ssaQ, sopB, sopE, and sip4-F are involved in the invasion and infection mechanisms of Salmonella enterica. Virulotyping is a new scheme of genotyping based on the detection of these virulence associated genes (Virulence profile or VP) that has been proven to be an efficient tool for identification and clustering of strains/clones of S. enterica.

Objectives

Evaluate the association between VP with Sequence Type (ST), serotype, and the geographic region of provenance of Salmonella enterica strains.

Methods

Ninety strains of S. enterica isolated from meat and dairy products collected in the State of Michoacán, Mexico with known ST, serotype and provenance were analyzed. PCR products of the genes invA, rmbA, ssaQ, sopB, sopE and sip4-F were included in the study.

Conclusions

Eleven different VPs were observed and named alphabetically as A through K. VP-A contained all of the virulence genes analyzed and is widely distributed throughout the state, followed by VP-B and C with the largest number of virulence-associated genes, except for genes invA and sopE, respectively. The highest diversity of VP was observed in the port of Lázaro Cárdenas, and in the regions of Morelia and Apatzingán. VP was unrelated to ST and serotype of tested strains. Because in all cases at least three virulence-associated genes were found, our results show that the studied strains of S. enterica have a high pathogenic potential, which is relevant for
public health given the presence of these strains in foodstuffs of daily consumption in the state of Michoacán.
POTENTIAL USE OF ESSENTIAL OILS FOR SURFACE DISINFECTION

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Background
Bacteria can attach to different surfaces and form biofilm causing hygiene problems. Microbes in biofilms are more resistant to antimicrobials than planktonic cells. The organic food industry can use only a limited number of disinfectants. Customers prefer "green technologies" in food production and processing thus there is a continuous need for the use of new natural compounds. Essential oils (EOs) have antimicrobial effect, are easy to produce and eco-friendly. Essential oils are hydrophobic and make the membrane more permeable and more hydrophobic on low pH.

Objectives
In the present study we have investigated cinnamon, marjoram and thyme EOs (on the basis of our pre-study) against bacterial biofilm formation on different surfaces.

Methods
MIC/MBC values of the EOs were determined against food spoilage bacteria: E. coli, P. putida and pathogens: L. monocytogenes, S. aureus, MRSA, B. cereus and B. subtilis. Disinfection time depending on pH and EO concentration was established. Biofilm elimination capacity of the EOs (24 and 168 hours old) from stainless steel and polypropylene surfaces (frequently used in food industry) was tested.

Conclusions
Biofilm formation on plastic was stronger, than on stainless steel probably due to the roughness of surfaces. All investigated EOs had good biofilm removing effects. Best result was achieved with marjoram. In most cases bacterial reduction was 99.99%. EOs had also antibacterial effect on the tested isolates of MRSA and L. monocytogenes.
EFFECT OF DRYING TREATMENT WITH POTASSIUM CARBONATE (K2CO3) ON SAFETY QUALITY OF DRIED FIG (FICUS CARICA L.), AS AN ALTERNATIVE TO TRADITIONAL PROCESS.

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Background

Sun drying process involves microbial risks such as fungal spoilage which could also produce mycotoxin.

Objectives

Therefore, the aim of this work was to study the effect of drying treatment with potassium carbonate (K2CO3) combined with osmotic solution on safety quality of fig from ‘Calabacita’ cultivar.

Methods

Two treatments of drying were used, firstly figs were dipped into K2CO3 solution (10%) for 30 min, then in a sugar solution (50%) for 24h and finally drying at 65°C in a drier (TP). A second batch, with figs traditionally sun drying, was used as control (ND). Both batches of dried fig were packed in polyethylene punnets with a macroperforated film and stored at 20 ºC for 90 days. Moisture loss was monitored during treatment and storage. Moreover, mold counts and mycotoxin presence were study during storage. Finally, molds were isolated and identified by sequence of the ITS region.

Conclusions

The moisture level in TP was 22% after two days of treatment, whereas similar value was obtained in ND after 12 days. Then, the used of potassium carbonate combined with sugar solution reduced significantly the time of drying. During storage, mold reached counts of 1.82 and 2.22 log CFU/g for TP and SD, respectively, at 30 days. Cladosporium cladosporioides was the main specie followed by Aspergillus fumigates in both batches, although they were found in greater number in ND. No mycotoxins were detected in both batches, however, the higher level of mold found in control batch, as well as longer drying process could suppose greater risk of mycotoxin production.
PROFICIENCY TESTS FOR CAMPYLOBACTER ORGANIZED BY THE NATIONAL REFERENCE LABORATORY FOR FOOD OF ANIMAL ORIGIN IN POLAND DURING 2008-2014
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Background
In accordance with the EU regulation No. 882/2004 National Reference Laboratories (NRL) are responsible for organization of proficiency tests (PT) for official food control laboratories. In Poland, NRL for Campylobacter was nominated at the National Veterinary Research Institute in Pulawy.

Objectives
The aim of the study was to evaluate the proficiency of laboratories by their participation in PTs towards determination of presence and number of Campylobacter in samples of food of animal origin.

Methods
The PTs were organized between 2008–2014 according to ISO/IEC 17043:2001, ISO/TS 22117:2010, and ISO 13528:2005 standards. In each year, two rounds of PT, i.e. one for detection and one for enumeration of Campylobacter in fresh poultry or pork meat, were performed. The samples were contaminated with the target C. jejuni or C. coli microorganisms at the levels between $10^2 - 10^3$ cfu for 10 g or 25 g (detection) and $<10^0 - 10^5$ cfu/g (enumeration). The samples were sent to the laboratories by the courier post in a cooled conditions which were obliged to begin the analyses within 24 h (detection) or immediately upon receiving the samples (enumeration). The number of official laboratories participated in each PT round varied from 7 (enumeration; year 2010) to 20 (detection; year 2012).

Conclusions
Most of the laboratories obtained satisfactory results (70 – 100% participants depending on the PT round) which indicated that food safety is under appropriate supervision. The PTs were the effective tool used by NRL to supervise the official food control laboratories in Poland.
A NEW MULTIPLEX REAL-TIME PCR METHOD FOR DETECTION OF SALMONELLA SPP., LISTERIA MONOCYTOGENES AND VEROTOXIGENIC ESCHERICHIA COLI IN CARCASSES OF SLAUGHTERED ANIMALS

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Background

Salmonella spp., Listeria monocytogenes and verotoxigenic Escherichia coli (VTEC) are among the most dangerous food-borne bacterial pathogens. Therefore, the availability of rapid and specific methods, especially to simultaneous detection of these pathogens, is needed.

Objectives

The objective of the study was to develop and evaluate a new TaqMan multiplex real-time PCR method for detection of Salmonella spp., L. monocytogenes and VTEC in slaughtered animals.

Methods

The method includes the enrichment step in Tryptone Soya Broth, DNA extraction and two real-time PCR reactions, one for detection of invA and hly genes of Salmonella spp. and L. monocytogenes, respectively, and another for detection of vtx1, vtx2 and eae of VTEC. The results of validation showed a relative sensitivity, specificity and accuracy of 100% compared to the reference methods. The limit of detection (LOD) was established for 1 cfu per swab for Salmonella spp. and L. monocytogenes and 2 cfu for VTEC. The method developed was used to analyze 145 swabs collected at the slaughterhouse level from cattle, pigs and poultry carcasses. Among the 75 samples recovered from cattle 24 were positive for VTEC and two for Salmonella spp., whereas swabs from pigs (n=45), 3 and 2 samples, were positive for these pathogens, respectively. None of microorganisms tested were detected in 25 samples from poultry.

Conclusions

The method developed allows for rapid and specific detection of three food-borne pathogens in carcasses of slaughtered animals.
PROPHAGE-ENCODED STAPHYLOCOCCAL ENTEROTOXIN A: REGULATION OF PRODUCTION IN DIFFERENT STAPHYLOCOCCUS AUREUS STRAINS

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Background
Staphylococcal Food Poisoning (SFP) is one of the most common food-borne intoxication diseases, caused by enterotoxins produced by Staphylococcus aureus. 21 staphylococcal enterotoxins (SEs) or enterotoxin-like proteins (SEls) have been recognized until now with enterotoxin A (SEA) being most frequently implicated in SFP outbreaks. SEA is encoded by the sea gene, located on the genome of Siphoviridae bacteriophages and unlike most other enterotoxins is not regulated by the assessor gene regulator (agr). Previous studies have shown that the regulation of the sea gene in the different SEA-producing strains is partially linked to the lifecycle of the prophages carrying the gene. Differences have been observed on the levels of toxin produced among the strains, and are hypothesized to originate from the Siphoviridae phages.

Objectives
With this study, we aim to get a better insight on the regulation of the sea gene by understanding the level on which the phage life-cycle affects SEA production.

Methods
We study five S. aureus strains under optimal and chemically induced conditions and monitor the replication of the phage in connection with sea gene copies and mRNA transcript levels. Furthermore, we follow the extracellular toxin levels and estimate phage titters.

Conclusions
The data obtained have showed a distinct correlation between replicating phage copy levels and sea gene copy levels in both growth conditions, linking phage induction with the increase observed on SEA levels. The significance of these findings lays in the realization that any treatment potentially inducing replication of the sea-carrying prophages, will enhance the risk for SFP.
Background

Smoked salmon as a ready-to-eat (RTE) food can be typically contaminated with *Listeria monocytogenes*, albeit in low numbers. The legal safety limits for RTE are set in the European Commission Regulation No. 2073/2005 and are particularly exceeded by RTE fishery products and cheeses.

Objectives

The aim of the present study was to investigate the *L. monocytogenes* contamination rates in cold-smoked fishery products in Austria over a one-year period (2013-2014).

Methods

A total of 250 RTE samples were collected in the region of Vienna and analysed at the end of shelf-life. The influence of physico-chemical properties (aw, pH and phenolic content) on the prevalence was also assessed. Isolates were characterized by multiplex PCR serogrouping and rep-PCR. The biofilm forming ability of *L. monocytogenes* isolates was assessed by the crystal violet method and a modified fluorescence microscopy-based technique.

Conclusions

Among the tested fishery products 23% were contaminated with *L. monocytogenes* which correlates with international literature data. Enumeration results ranged from 1 to 760 CFU/g. Serogrouping achieved that 87% of isolates have been classified to 1/2a, 3a serotype and others mainly belonged to groups 4b, 4d, 4e and 1/2b, 3b, 7. Biofilm forming ability could be determined for some isolates when compared with *L. monocytogenes* reference strains representing different serotypes and sources. The physico-chemical data revealed a strong variability within the smoked fishery product category. These discrepancies show the need for factory specific analysis of *L. monocytogenes* growth potential in smoked fishery products in order not to misinterpret the risk-categorization of such products.
FEMS-0709
Fungal bacterial interactions

ACTINOMYCETES AS BIOCONTROL AGENTS IN GRAPEVINE WOOD DISEASES: A GOOD TENANT IN THE ROOT?
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Background
One of the most important problems in wine production regions is the premature death of young grapevines. It is due mainly by the infection of fungus that produces wood diseases, like Petri disease and black-food disease. There are two principal ways to infection. One is the 'industrial process' in order to produce the commercial grafted plants. Another possibility is associated to 'natural processes' when the infection is produced on the field mainly by the pruning wounds or by natural infection through the root system. In fact, most of pathogens associated to these diseases are able to infect the grapevine through the root system.

Objectives
The main objective of this study is the use of actinomycetes isolated from the rhizosphere of young grapevines as biocontrol agents (BCAs) in order to reduce or avoid the infection through the root system by pathogenic fungus.

Methods
Several actinomycetes were isolated from rhizosphere of young plants of Vitis vinifera. These were unequivocally identified by molecular methods. Their capability as biocontrol agent were bioassayed in vitro against several pathogenic fungus. Those showing the best capabilities were selected for 'field studies' on grafted plants in order to know the real behaviour as BCAs.

Conclusions
Several actinomycetes have shown a good capability to reduce the infection rate through the root system of several fungal pathogens causing decline of young grapevines. They could be a good alternative in order to avoid chemical products in the establishment of new vineyards.
FEMS-3002
Fungal bacterial interactions

EFFECT OF SOME LIGNICOLOUS EDIBLE AND MEDICINAL FUNGAL SPECIES AGAINST VIRAL (BACTERIOPHAGE) AND BACTERIAL DNA

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Background

Nowadays, when high resistance against common antibiotics is evident, a great attention has been focused on some new natural sources of antimicrobial agents, including mushrooms. Medicinal mushroom species mostly belong to lignicolous (wood-decaying) fungi and represent particularly promising organisms with bioactive compounds acting as antimicrobials.

Objectives

In the present study, the genotoxic effect of 80% ethanol and 70% methanol, extracts of some lignicolous edible and medicinal fungal species: Agrocybe aegerita, Meripilus giganteus, Coriolus versicolor, Daedaleopsis confragosa, and Stereum fuscum were investigated. Ganoderma applanatum extract was prepared in ethanol after hydrodistillation procedure. They have been tested against bacteriophage DNA (δ - Podoviridae and σ-1 - Siphoviridae), as well as against DNA extracted from Escherichia coli (ATCC 25922) and Micrococcus flavus ATCC 40240.

Methods

The extracted DNAs from viruses and bacteria were analyzed after treatment with fungal extracts (initial concentrations 200 mg/ml and 500 mg/ml) after 24 h in broth medium. Activity of the analyzed extracts on DNA fragmentation was visually determined on agarose gel after electrophoresis.

Conclusions

After 24h, extracts of C. versicolor and G. applanatum lead to the disintegration of E. coli DNA, while the extract of M. giganteus showed slight effect. All extracts showed disintegration effect against both phage DNAs, except extract of edible species A. aegerita. The other extracts give effect in concentration 1:100, except S. fuscum and C. versicolor against σ-1 phage DNA. The phage DNAs were not disintegrated with
any extract in concentration 1:10 000. The results showed that some higher concentrations of extracts possess genotoxic activity.
Background

Since fungi share common microorganisms such as *E. coli*, *S. aureus* and *P. aeruginosa* with animals and humans, it is believed that the humanity will meet more benefits from fungi and mushrooms, as antibiotics producers, than from plants. Many lignicolous species have been detected to act as antimicrobials until now, but activities are mostly influenced by fungal strain specificities and habitat characteristics, including geographic position.

Objectives

Since antibiotic mechanisms of fungi are still unclear, the aim of this investigation was to find connection between antibacterial and DNA genotoxic activity. Two kinds of extracts were prepared from fresh samples of autochthonous fungal species: *Fomes fomentarius* and *Pleurotus ostreatus* after simple extraction and after pouring with liquid nitrogen in order to disintegrate sporoderm.

Methods

Antibacterial testing was done with 70% MeOH extracts using macro and micro-dilution methods. Genotoxic effect against two bacterial strains *E. coli* and *M. luteus* was checked with 20%, 10% for *P. ostreatus* and 10%, 2.5% for *F. fomentarius* extract concentration. After 24h incubation, isolated DNA was applied on agarose gel electrophoresis.

Conclusions

The best antibacterial activity was obtained for *F. fomentarius* extracts mostly against Gram positive strains: *M. flavus* ATCC40240, *S. aureus* ATCC25923, *C. perfringens* ATCC19404 and *B. subtilis* ATCC6633 and gram negative *E. coli* ATCC25922. Significant increase in the size of bacterial cells was detected.

The MIC showed significant *E. coli* DNA fragmentation, indicating that this can be a possible mechanism of antibacterial activity, but also that these extracts can have a potential genotoxic activity which should be further examined.
ENHANCED ACCESS TO ORGANIC NITROGEN AS A DIRECT BENEFIT IN THE INTERACTION OF THE FUNGUS MORCHELLA CRASSIPES WITH SOIL BACTERIA

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Background

Fungi and bacteria have a long co-evolutionary history in terrestrial ecosystems. In soils, they are the basis for nutrient recycling. Therefore, not only do they co-exist, but they are also potential competitors for resources. In our laboratory, it was shown that the bacterium *Pseudomonas putida* benefits from interaction with fungi by using the so-called “Fungal Highways” to disperse in porous unsaturated soil-like media. Dispersal allows the bacteria to reach nutrient sources, as their mobility is limited in these conditions. However, the direct fitness benefit for the transporting fungi remains unknown. Until now, a long-term benefit was shown for the ectomycorrhizal and saprophytic fungus *Morchella crassipes*, which acted as a farmer of the dispersing bacteria and storing bacterial carbon into resting structures.

Objectives

We focus on the investigation of the benefit obtained by fungi during dispersal of bacteria.

Methods

Experiments were all undertaken on skimmed-milk agar medium (SMA; protein as N source) to see how extracellular proteolytic activity is used to access organic nitrogen. In addition, the fungal enzymes responsible were characterized according to their activity.

Conclusions

We have determined that fungal proteolysis can be enhanced by the presence of *P. putida* even when fungal cells are removed. The enhanced activity requires living bacterial cells, which is also observed for other bacteria dispersing in the fungal
mycelia. In conclusion, enhancement of proteolytic activity seems to be a direct fitness benefit for the interaction of *M. crassipes* with soil bacteria.
ENDOPHYTIC BACTERIA OF TOMATO: A SOURCE OF ANTIFUNGAL AND ANTIBACTERIAL COMPOUNDS
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Background

Many microorganisms are found inside various plant tissues. There, they not only compete with the plant immune system, but also with other endophytes and pathogenic microorganisms. We study endophytes of two underexplored niches inside the tomato plant, the xylem and the leaf apoplast. Xylem vessels are the habitat of vascular wilt causing pathogens such as the bacterium \textit{Clavibacter michiganensis} and the fungus \textit{Verticillium dahliae}, whereas the leaf apoplast harbours the leaf mould \textit{Cladosporium fulvum} and the late blight oomycete \textit{Phytophthora infestans}.

Objectives

Study the mechanism of \textit{in planta} inter-microbe competition. Identify biocontrol species

Methods

Endophytic bacteria were isolated from tomato xylem and leaf apoplastic space. Colonies were picked after incubation on TSA for 2-10 days at 25°C. Isolated strains were inoculated on a lawn of fungal, bacterial or oomycete plant pathogens and the release of antimicrobial compounds was scored. Garden cress was inoculated with selected strains. Seedling development was followed, and after 20 days endophytic bacteria were re-isolated from the stems.

Conclusions

We isolated >120 tomato associated bacteria. More than half have antifungal, antibacterial and/or anti-oomycete properties during \textit{in vitro} co-culture. >75\% of species isolated from above ground parts of tomato were gram-positive. Garden cress was inoculated with strains with strong antifungal properties. Most strains promoted plant growth, and could be re-isolated from the stems, indicating a true endophytic lifestyle. Currently we are elucidating the antagonistic mechanisms and testing their potential for inhibiting pathogens \textit{in planta}. In conclusion, the above
ground tomato endophytic bacteria are a promising source of antimicrobial compounds.
FEMS-2937
Fungal bacterial interactions

EFFECT OF BACTERIAL DISPERSAL ON THE BIOAVAILABILITY OF SEMIVOLATILE ORGANIC CONTAMINANTS

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Background
Contaminants bound to biogeochemical interfaces (BGI) are only hazardous if they become bioavailable. Therefore, the most effective remediation technique should lead to an optimal coverage of BGI with degrading microorganisms in order to prevent contaminant release to the environment. Simultaneously they should reduce the contaminant concentration at its source.

Objectives
Here, we investigated the impact of bacterial dispersal on BGI on the outgassing of semivolatile phenanthrene (PHE). We tested the hypothesis whether the presence of previously described fungal transport networks (‘fungal highways’) of bacteria leads to: (i) bacterial distribution along the transport network, (ii) efficient bacterial distribution on the surface, and (iii) an increased biomass production fostering the degradation of PHE, that are released from the system.

Methods
We designed a laboratory microcosm that mimicked a continuous PHE release to a model surface (agar) in the presence and absence of model dispersal networks (glass fibers) that facilitated the transport of the poorly motile PHE-degrading Pseudomonas fluorescens on agar surfaces.

Conclusions
We observed that the presence of the dispersal network resulted in (i) an increased spatio-temporal spreading of bacteria, (ii) an increased bacterial coverage and growth on the agar surface, and (iii) a subsequent effective degradation of outgassing PHE and effective reduction of PHE contamination beyond the PHE hotspot. Our data suggest that fungal mycelia may promote the formation of an adapted microbial population that will degrade hazardous molecules that desorb from the contaminant source. Potentially, such an activity produce no emission of contaminants to the pore and groundwater, and hence, to higher organisms.
FEMS-3088
Fungal bacterial interactions

STUDYING INITIAL FUSARIUM GRAMINEARUM-BIOCONTROL PSEUDOMONAS FLUORESCENS INTERACTIONS IN AN ESTABLISHING MICROTITER PLATE MICRO COSM SYSTEM
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Background
The hyphosphere constitutes a microenvironment in which enzymes, metabolites and antibiotics secreted by the fungi are abundant which could attract bacteria.

Objectives
We here describe the first steps in establishing a microcosm system to study fungal-bacterial interaction under low-nutrient conditions emulating natural habitats like soil or the rhizosphere, then study fungal-bacterial interactions. Fusarium graminearum head blight in cereals was used as a model fungus, whereas Pseudomonas fluorescens strains P13, UTPF105, UTPF127, UTPF125 and B4 identified as biocontrol agents from Iranian soils were used as model bacteria.

Methods
For establishing this microcosm, 1000 spores of F. graminearum with 100 µl diluted medium was added to each microtiter well and incubated for 38 h at 20°C then the media was removed and inoculated with 100µl of 10^4/ml bacterial suspension. After 24 h further incubation the microcosm system was stained by SYBR Green (SG) and Calcofluor white (CW) fluorescent stains. SG binds DNA and absorbs blue light and emits green light (λ = 520nm). CW binds to chitin. Some photos of each CW (under UV light) and SG (under blue light) wells were took. Both blue hyphae and green DNA signals were analyzed by ImageJ software. The experiment was arranged as randomized complete design with 4 replications. A Duncan test with a probability of 0.01 was used to show significant differences between treatments.

Conclusions
CW result showed that the biocontrol strains decreased fungal growth. In this reduction, UTPF125 was more successful. SG results showed that in a same hyphosphere, growth of different bacterial strains were significantly different.
A MECHANISM OF ALTERNATIVE SPlicing IN FILAMENTOUS FUNGI

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Background

Alternative intron splicing could lead to frame-shifted ORFs so that peptides with different terminal extensions can be produced. Alternative splicing of introns bordered by coding exons may yield the presence or absence of additional domains. We have found a Penicillium chrysogenum gene that yields two transcripts of different size.

Objectives

An alternative intron gives rise to the two mature mRNAs using alternative, coupled lariat-branchpoint domain/acceptor sites. In P. chrysogenum, the short variant is 213 nt, the long variant is 891 nt long. Both transcripts yield a full-length protein. We found the presence of one canonical donor and one canonical lariat-branchpoint domain/acceptor couple within the short variant of the alternative intron, potentially defining an “internal” intron of 64 nt. We investigated whether this “internal” intron produces splicing intermediates.

Methods

By sequencing cloned RT-PCR amplification products, we have elucidated the sequences of the mature mRNAs and that of splicing intermediates in P. chrysogenum and Aspergillus nidulans.

Conclusions

We showed that contrary to “intron definition theory”, the “external” (5’) donor was not used with the most 5′ (“internal”) acceptor site. The “internal” (3′) donor was not used with the most distal (most 3′) acceptor site. It is the “internal” intron that is spliced alternatively using the 3′ donor and either of the closely neighbouring lariat-domain/acceptor couples. The identity of the (more outward) acceptor used for the subsequent removal of the “external” intron and hence, the peptide ultimately produced, are thus primarily determined by the outcome of the first, alternative splicing reaction.
Background

Cutinases play a role in plant pathogenesis as they degrade cutin, a structural lipid component of plant cuticle. They also act on short- and long-chain fatty acids, a feature that makes them suitable for biotechnology applications (Carvalho C, 1998). In Aspergillus nidulans genome there are four putative genes of cutinases (ancut1,2,3, and 4) and neither their role in cutin degradation or the expression conditions and their regulation are well understood, although it may be related to lipid metabolism (Hynes MJ, 2006).

Objectives

- Determine the expression levels of the cutinases encoded in the A. nidulans genome in response to different carbon sources that may act as repressors or inducers of the enzymes.
- Study the effect of two global TF (farA, farB) involved in lipid metabolism over the cutinase expression.
- Propose a regulation model of the cutinolitic system.

Methods

Aspergillus nidulans was grown in minimal medium and different carbon sources were used. Mycelium was used for RNA’s extraction with TRIzol. cDNA was synthetized using dT primers and Reverse Transcriptase. Finally the expression level of the genes was measured in Multiplex qRT-PCR assays and normalized with an endogenous gene. These expression levels were compared with the enzyme activity found in the supernatants in all the conditions studied.
Conclusions

ancut1, ancut2 and farA were upregulated with cutin and cutin monomers and showed CCR in presence of glucose. ancut3 and farB were expressed constitutively and ancut4 wasn’t detected in any condition. Cutinase expression in A. nidulans resembles that found for F. solani (Li D, 1997).
Background
The fungus *Penicillium roqueforti* has been poorly studied. Currently, we are studying a strain of this fungus that contains a dominant active Gαi protein. This strain shows several phenotypic alterations compared with the wild-type strain. In a preliminary study, we observed a cDNA sequence differentially expressed in the wild-type strain. This sequence matched an unnamed ORF from *P. roqueforti* that encodes a putative Zn(II)$_2$Cys$_6$ protein of unknown function. Proteins containing Zn(II)$_2$Cys$_6$ domains are exclusively found in fungi, but few of them have been functionally characterized.

Objectives
To characterize a gene from the filamentous fungus *Penicillium roqueforti* that encodes a Zn(II)$_2$Cys$_6$ protein, whose function to date remains unknown. We have named this gene *pcz1*.

Methods
*pcz1* was silenced using RNAi-silencing technology. Two transformants (named M9 and M11) showed dramatic reductions in the levels of *pcz1* transcript, confirming the successful knockdown of the gene. In these transformants, the apical extension rates, the conidiation, and the conidial germination were measured by standard methodologies.

Conclusions
The silencing of *pcz1* in the transformants altered the normal phenotype of *P. roqueforti*. Specifically, RNA-mediated silencing of *pcz1* resulted in the strong repression of conidiation, decreased apical growth, and the promotion of conidial germination in *P. roqueforti*, even in the absence of a carbon source. Our results suggest that *pcz1* may be involved in fungal development. Specifically, it may be a positive regulator of conidiation and apical growth, but it may be a repressor of conidial germination in *P. roqueforti*.

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Background

Coprinellus congregatus, generates mushrooms from dikaryotic mycelia, and this fungus produces several laccases: hyphal tip laccase and primordial laccase and they are not secreted outside of cells. When the dikaryon was grown in an acidic medium (pH 4.0-4.5), a new laccase was secreted into the culture supernatant. We have also cloned its genomic and cDNA genes, and also confirmed its expression only under acid condition by Northern blot analysis.

Objectives

There must be two responsive regions in the promoter region which are related to the signals that the mycelia is a dikaryon and is under acid stress respectively. In order to determine the responsive domain in the promoter, a recombinant construct having the acidic laccase gene promoter and green fluorescent protein (GFP) gene was generated.

Methods

The construct was introduced into C. congregtus monokaryones (a1 and a2), and the transformants were selected. Heterozygotic and homzygotic transformants were generated by mating proper partners. These transformants were transferred to the acid liquid media. Fungal cells were collected and they were used for the analysis of the promoter expression under confocal microscope and/or real time RT-PCR.

Conclusions

Homozygotic transformants showed more than 25 times higher expression than the negative control: this system helped the promoter domain analysis. Different lengths of acid laccase gene promoter-GFP will be constructed, and these will be introduce to the monokaryons. When the transformants carrying these various lengths of promoter
were analyzed at the acid culture conditions, the responsible doamins for dikaryon and acid stress will be determined.
Background
Transcription start site (TSS) sequencing gives a high resolution map of transcription initiation region for identifying promoters and fine-tuned transcriptional regulation mechanism. Although TSSs of various eukaryotic model organisms have been revealed, no species belonging to Basidiomycota has been known yet.

Objectives
In order to understand transcriptional regulation in terms of promoters and transcriptional regulation on growth conditions, we report the first TSS analysis of Basidiomycota, the lignin-degrading fungus Schizophyllum commune

Methods
Total 8.5 million RNA reads were generated by rapid amplification of cDNA ends (RACE) from two different growth conditions (cellulose or glucose as a sole carbon source) and mapped into the genome to allocate the transcription start sites.

Conclusions
The positions of transcript head for the 4,959 genes were retrieved from both growth conditions. Comparing to the features of promoters found in eukaryotic model organisms, TSS distribution of S. commune were categorized into distinct groups such as single start sites (‘sharp’ promoter), multiple start sites (‘broad’ promoter) or mixed. To associate these features with DNA motifs in promoters, we examined the upstream 500 bp region of all genes. We found 16 motifs in the ‘sharp’ promoter upstream regions including typical TATA-box. The TSS analysis provides not only a basis for understanding of promoter structures but also a principle to build synthetic promoter construction for desired gene expression in S. commune.
THE PREVALENCE OF PNEUMOCYSTIS JIROVECII IS CHANGED?
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Background
Pneumocystis pneumonia (PCP) that is caused by Pneumocystis jirovecii is prevalent in HIV-infected patients. Pneumocystis the colonization of Pneumocystis is associated with airway inflammation and obstruction.

Objectives
The current study was conducted to identify the prevalence of P. jirovecii in suspected tuberculosis (TB) patients and four HIV positive patients in Ahvaz, Iran.

Methods
111 suspected TB positive patients and four HIV patients were investigated for P. jirovecii identification from Bronchoalveolar lavage specimens. The specimens were subjected for Grocott's methenamine silver (GMS) staining and then DNA extraction following Nested-PCR with specific primers. The PCR product then subjected for sequencing.

Conclusions
The findings demonstrated 50% of HIV positive patients were contaminated with P. jirovecii and 2 remaining patients showed negative results. In 111 suspected TB positive patients 25.2% were positive for TB, which 9 samples were positive for PCP by P. jirovecii. In negative TB patients, 19 positive P. jirovecii were identified. The analysis by Chi-square demonstrated that there was no significant difference between sex and presence of P. jirovecii. The current statistical analysis also presented no significant difference between TB and P. jirovecii. Finally, the GMS only detected Pneumocystis jirovecii in three patients. Our findings demonstrated high prevalence of P. jirovecii in Ahvaz which was more than the previous reports in Iran. Our investigation revealed that the Pneumocystis jirovecii is important in all respiratory tract infected patient not only in HIV or TB patients.
AXL2, THE HOMOLOGUE OF THE YEAST BUD SITE SELECTION PROTEIN AXL2P AFFECTS ARTHROSPORE FORMATION IN ACREMONIUM CHRYSOGENUM

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Background

Acremonium chrysogenum is the natural producer of the β-lactam antibiotic cephalosporin C. Due to its antibacterial activity against gram-positive and gram-negative bacteria, this β-lactam antibiotic is of great biotechnological and medical relevance. To optimize cephalosporin C production efficiently, continuous and directed improvement of industrial strains is required.

Objectives

A typical morphological feature of A. chrysogenum is the fragmentation of vegetative mycelium into arthrospores. These are uni- or binuclear cells, which develop during a prolonged cultivation under limited nutrient supply. Due to the known correlation of cephalosporin C production and arthrospore formation, we are interested in the identification of specific regulatory factors affecting both, cephalosporin C biosynthesis and morphological development. Here, we present a functional characterization of Axl2 in A. chrysogenum, a homologue of the yeast bud site selection protein Axl2p. In addition to that we are interested in the localization of Axl2 and other proteins of the bud site selection complex to verify their direct regulation of arthrospore formation.

Methods

Homologous recombination through split-marker approach; microscopic analysis

Conclusions

An axl2 deletion strain of A. chrysogenum shows an accelerated arthrospore formation in comparison to its recipient strain. Therefore it might act as putative regulator of septation processes of fungal hyphae and consequently affect the arthrospore formation. To further investigate this developmental process we were able to establish the visualization of filamentous actin with the live cell marker Lifeact.
Background
In nature, saprotrophic fungi have an essential ecological role in decomposition of organic matter. In soils, they break down material like leaves, wood, dead bodies, etc. Fungal colonization in natural environments requires advanced functional relationships, including various antagonisms and synergisms, between the different components of the biotope. Colonization can be described, analyzed at different scales (macroscopic and molecular) and modeled. In indoor environment, fungi find niches favorable to their development, but are most often undesirable. Indeed, micromycetes induce pathologies to the occupants (allergies, poisoning and infections) and can also cause irreversible damages to the materials they infest. Colonization of different man-made substrates (construction and decoration products) by mold is insufficiently described to develop colonization models: Is artificial substrate colonization done serendipitously or is it determined like in natural environment by a defined succession of fungi driven by antagonisms and synergisms? A better knowledge of the dynamic of fungal colonization would be helpful to imagine preventive strategy for health and artifacts preservation.

Objectives
In this context, our research in microbial ecology aims to describe and understand the fungal colonization mechanisms in enclosed spaces.

Methods
First, a macroscopic characterization of the fungal succession on two artificial substrates has been made by culture and molecular methods. Secondly, Volatile Organic Compounds (VOC’s) emitted by microfungi according to the growth substrates have been explored.

Conclusions
This new knowledge will offer alternatives to slow and limit fungal growth on building products to the use of traditional biocides.
IDENTIFICATION OF A SECOND LACTOSE PERMEASE GENE (LACPB) IN ASPERGILLUS NIDULANS

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Background

For most micro-organisms, lactose is a gratuitous carbon source that is slowly assimilated. In the filamentous fungus Aspergillus nidulans, lactose hydrolysis occurs intracellularly. Transport of lactose, which was previously shown to involve the lactose permease LacpA, is the limiting step in its catabolism in A. nidulans. Since lacpA deletion mutants are still able to grow on lactose, we assumed the existence of a second, physiologically relevant uptake system.

Objectives

Identification and characterization of the second lactose permease gene (lacpB, locus AN2814) in A. nidulans, which encodes a structural paralog of LacpA.

Methods

Single (lacpB) and double (lacpA, lacpB) lactose permease deletion mutants were created to test their phenotype (both growth and carbon source uptake) on solid and liquid media. Transcription profiles of lacpB and lacpA were also compared using Northern blot analyses.

Conclusions

Growth of lacpB-single and lacpA/lacpB double deletants was strongly reduced on lactose on solid media, while in submerged cultures, the lactose phenotype was even more pronounced. Expression of the two lacp genes differed markedly. In addition to lactose, lacpA transcription also responded to D-galactose and L-arabinose, while lacpB was found induced by cellobiose, sophorose as well as lactose. Neither of the genes appeared to be expressed on D-glucose, D-xylose or D-mannose. We conclude that lacpA and lacpB are jointly responsibe for wild-type lactose uptake by A. nidulans.
CONIDIAL ANASTOMOSIS TUBES AND HETEROKARYOSIS IN THE ASEXUAL PLANT PATHOGENIC ASCOMYCETE VERTICILLIUM DAHLIAE

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Background
Verticillium dahliae is a ubiquitous soilborne fungus that is known to reproduce strictly asexually. Therefore, different strains can only exchange and recombine genetic information through the parasexual cycle, starting with fusion (anastomosis) between hyphae of different genotypes to the formation of heterokaryons. Recently it was shown that parasexuality may alternatively begin in other fungi with fusion between conidial germlings via conidial anastomosis tubes (CATs).

Objectives
(i) To explore the possibility of self-/non-self fusion between V. dahliae germlings via CAT formation and study CATs' physiology, (ii) to study nuclear dynamics and heterokaryon formation through CATs, (iii) to identify signaling pathways that are required for CAT formation in V. dahliae.

Methods
(i) Live-cell imaging and analysis of fixed germling preparations with differential interference contrast (DIC) and fluorescence microscopy, (ii) tagging of fungal nuclei with fluorescent proteins for the study of nuclear migration, (iii) use of specific inhibitors of major signaling components and generation of corresponding gene deletion mutants via Agrobacterium tumefaciens-mediated transformation.

Conclusions
Typical CAT formation is frequent between V. dahliae germlings, although it is inhibited by the presence of nutrients and varies significantly between different strains. It is also dependent on pH, conidial density, and the history of the fungal cultures used. Cytoplasmic mixing and nuclear migration are permitted through CATs, allowing the formation of heterokaryons when conidia of different genotypes are combined. The formation of CATs in V. dahliae requires functional MAP kinase and ROS signaling components, similarly to normal development/morphogenesis and pathogenicity of the fungus.
Fungal development

CO2 REPRESSION OF FRUITING BODY FORMATION IN SCHIZOPHYLLUM COMMUNE IS MEDIATED VIA CAMP

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Background
The most important environmental cues impacting fructification in Schizophyllum commune are light and CO₂. Blue light is needed to initiate fructification, while high levels of CO₂ repress this process. The most ubiquitous mechanism of CO₂ sensing in prokaryotes and eukaryotes is based on cellular levels of cAMP. Regulation occurs through synthesis and degradation by adenylate cyclase (AC) and phosphodiesterase (PDE2), respectively. Most ACs localize to membranes and are activated by G proteins. However, there are also soluble ACs that are activated by bicarbonate or calcium ions. It has been proposed that fungal ACs are a hybrid between these two types and in most fungi there is only one gene for this protein.

Objectives
To study the mechanism for CO₂ sensing and its effect on fruiting body formation in S. commune.

Methods
CAMP was added to medium to assess its involvement in CO₂ sensing and fruiting in S. commune. Additional copies of pde2 were introduced by transformation and verified by qPCR. IBMX was used to repress PDE activity.

Conclusions
Addition of cAMP to the medium inhibited fructification but did not impact growth. Inhibition of fruiting body formation could be counteracted by overexpressing pde2. The pde2 overexpressor also fructified under repressing CO₂ levels where the wild type strain could not, reinforcing the existence of a link between CO₂ and cAMP levels. Addition of IBMX, an inhibitor of PDE2, to the medium abolished the effect of pde2 overexpression. Taken together, these results show that CO₂ levels are sensed via cAMP.
FUNCTIONS OF AUTOPHAGY-RELATED GENE SMATG12 IN FRUITING-BODY DEVELOPMENT OF SORDARIA MACROSPORA

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Background
In eukaryotes, autophagy is a highly conserved degradation process by which cytoplasm including organelles is non-selectively engulfed by double-membrane vesicles called autophagosomes. After fusion of the outer membrane with the vacuole/lysosome a vesicle surrounded by the inner membrane, the autophagic body, is degraded by hydrolytic enzymes and building blocks are recycled into the cytoplasm.

Objectives
In filamentous fungi, autophagy functions as a catabolic mechanism to overcome starvation conditions and to control diverse developmental processes under normal nutritional conditions. We recently showed that in the homothallic ascomycete Sordaria macrospora, autophagy related genes encoding components of the conjugation systems are required for fruiting body development or are essential for viability. Here, we cloned and characterized the S. macrospora (Sm)atg12 gene, encoding a ubiquitin-fold protein.

Methods
To examine the role of Smatg12 in detail, we replaced it with a hygromycin resistance cassette and generated a homokaryotic ΔSmatg12 deletion strain. GFP-labeled SmATG12 was used to localize the protein in vivo.

Conclusions
The ΔSmatg12 mutant displayed a decreased vegetative growth rate and was unable to form fruiting bodies. GFP-labeled SmATG12 was detected as punctured structures and co-localized with DsRED-SmATG8 at some places in the cytoplasm. We could show that lipidation of SmATG8 and proper autophagosome formation depends on SmATG12.
ENZYME ACTIVITY AND MALDI TOF/TOF MS ANALYSIS OF RESPONSES TO STRESS INDUCED BY ARSENIC AND ANTIMONY IN ACIDOMYCES ACIDOPHILUS FROM TIN-MINING SOIL

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Background
An Acidomyces acidophilus strain has been isolated from a tin mining soil in Cornwall, England that contained approximately 18970 mg kg\(^{-1}\) arsenic (As) and 228 mg kg\(^{-1}\) antimony (Sb) with the soil pH around 1. Culture experiments showed that the strain could tolerate up to 22500 mg kg\(^{-1}\) of As(V) and 100 mg kg\(^{-1}\) of Sb(V).

Objectives
This study examines the protein responses of A. acidophilus when exposed to As/Sb using MALDI TOF/TOF.

Methods
The strain was exposed for 24 h to 100 mg/L of sodium arsenate (Na\(_2\)HAsO\(_4\)) and potassium antimonate [KSb(OH)\(_6\)]. MALDI TOF/TOF MS analysis was carried out and tentatively identified that the following proteins were up-regulated in response to Na\(_2\)HAsO\(_4\) and KSb(OH)\(_6\): malate dehydrogenase, phospholipase B, phosphoglycerate kinase, glyceraldehyde-3-phosphate dehydrogenase and 3-isopropylmalate dehydrogenase. In addition, the activities of enzymes glutathione reductase, glutathione transferase, superoxide dismutase and catalase in the presence of As and/or Sb are still being investigated. Also, the isolated A. acidophilus and three A. acidophilus reference type strains (CBS 335.97, CCF4251 and CCF3679) were characterised by MALDI TOF/TOF and all four strains showed very distinctive spectra that distinguish them readily. It has also been found that A. acidophilus grown in different media exhibited different mass spectra, thus affecting the identification of this strain.

Conclusions
This provides further and in-depth opportunity to understand the extreme resistance to heavy metals by A. acidophilus and also identify the enzymes that are involved in the resistance mechanisms and thus contain biotechnological potential and future scientific uses such as bioremediation and clinical studies.
THE 7-TRANSMEMBRANE RECEPTOR GPR1 GOVERNS GROWTH, DEVELOPMENT AND MYCOPARASITISM-RELATED FUNCTIONS IN THE FUNGUS TRICHODERMA ATROVIRIDE AND INTERACTS WITH A SUR7 FAMILY MEMBRANE PROTEIN

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Background

Species of the fungal genus Trichoderma are among the most successful biofungicides in today’s agriculture although our understanding of the exact molecular mechanisms of their activity still is fragmentary. The biological control of plant diseases by Trichoderma includes direct antagonism of phytopathogenic fungi by parasitism. This mycoparasitic attack comprises sensing of the prey and chemotropic growth towards it followed by activation of “molecular weapons” such as cell wall-lytic enzymes, secondary metabolites, and infection structures. Consequently the receptors and signaling pathways involved in sensing and responding to the prey fungus are of special interest.

Objectives

The Trichoderma atroviride Gpr1 7-transmembrane receptor has been shown to be essential for regulating mycopathassism-relevant processes and it is implicated in the recognition of prey-derived signals. For further characterization of Gpr1, respective target genes being regulated by the Gpr1 receptor upon prey recognition have been identified by comparative genome-wide microarray-based transcriptomics of T. atroviride wild-type as well as gpr1 mutants.

Methods

Furthermore, direct interactors of Gpr1 were identified by the split-ubiquitin yeast-based two-hybrid system.

Conclusions

Functional characterization of one of the identified interactors, a plasma membrane protein belonging to the fungal-specific Sur7 family, by gene deletion revealed a role in the regulation of conidiation, polar growth, and the mycoparasitic attack of the prey fungus.
Background

The present study was conducted on fungal pathogen (Candida) causing ocular infections. The basis for effective treatment is rapid diagnosis of the disease and identification of its causative agent. Due to the limitations of classical methods for the detection of systemic fungal infections, it has become essential to develop a quick, sensitive and specific detection assay.

Objectives

A polymerase chain reaction (PCR) method was to be optimized that would be capable of detecting a wide range of medically important fungi from clinical specimens.

Methods

Eight Candida species sequences were downloaded from GenBank or EMBL databases. A set of unique primers were designed based on the conserved region in the given fungal species. To verify the specificity of the designed primers, the BLAST program at the NCBI website was employed to search the primers in short, near exact sequences. These primers were then analyzed by the AmplifX tool.

Conclusions

The results indicated that this study gives the strategy for the rapid detection of Candida species on the basis of genotype. A reproducible method for genotypic identification of Candida species will be very helpful for the doctors/practitioners to detect the specific species. In current medical facilities in Pakistan, the prediction of Candida sp. involved in ocular infection will be a valuable addition of information in the field of medicine. Pharmacists can make different antibiotics against different
diseases related to fungus *Candida* and can easily find out the specific cause of the related disease.
FEMS-0926
Fungal pathogenicity in humans

TRANSCRIPTIONAL PROFILE OF FUSARIUM OXYSPORUM AND FUSARIM SOLANI CHALLENGED WITH ANTIFUNGAL COMPOUNDS IN VITRO
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Background
Fusarium is a genus of filamentous fungi widely distributed and of high importance because several species of the genus are pathogens of plants and humans. In humans Fusarium is associated with superficial mycoses as onychomycosis and fungal keratitis in immunocompetent patients. However in immunocompromised patients, the infection can be disseminated and can cause the death of the patient due to the high resistance to antifungal treatments

Objectives
This project aimed to analyze the gene expression profile of both Fusarium oxysporum and Fusarium solani after the exposure to Posaconazole and Amphotericin B

Methods
The expression profile was performed by RNA-Seq of Fusarium spp and the genes with a significant expression were validated by qRT-PCR. We found an average of 20-30 of differentially expressed genes of the exposed strains relative to the controls

Conclusions
Some genes were associated with oxidative stress, ergosterol synthesis and cell wall structure, among others. The role of these genes in resistance against antifungal compounds is discussed. This study contributes to the general knowledge and understanding of the resistance mechanisms of Fusarium spp, which could improve the implementation of treatment strategies.
Fungal pathogenicity in humans

ANALYSIS OF GENE EXPRESSION PROFILES IN M. FURFUR USING NEXT GENERATION

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Background
The Malassezia genus is characterized by its lipophilic or lipid dependent species. These traits have been related with virulence, but until now the relation has not been clearly elucidated. M. furfur is a species presenting a high level of phenotypic variation. We previously described an atypical isolate of this species, which only assimilates tween 80. This finding suggested host adaptations mechanisms.

Objectives
We used microarray analysis to characterize the genetic expression that could give more clues about this metabolic advantage.

Methods
We analyzed gene expression differences between M. furfur CBS 1878 and the atypical M. furfur strain. In those experiments we used Dixon modified agar with and without tween 80. cDNA microarrays were designed and fabricated by Roche Nimblegen (NimbleGen Systems Inc., Madison, WI), using the Malassezia globosa genome. Normalization of the intensity values was performed using the VSN method. Differential expression analysis of the profiles was performed using pairwise comparisons and two different methods SAM and Ebayes algorithms. The enriched genes were analyzed by Gene Ontology (GO) terms.

Conclusions
The genes over expressed were associated with cyclophilins and thioredoxin, proteins involved in important cellular processes. We validated these results by qPCR. This analysis suggested that these candidate genes could be related with virulence.
ANTIFUNGAL SUSCEPTIBILITY TESTING OF YEASTS ISOLATED FROM THE PATIENTS WITH NOSOCOMIAL INFECTION IN SLOVAKIA.

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Background
Yeasts are common fungal opportunistic pathogens in humans playing a significant role in the morbidity and mortality of immunocompromised patients.

Objectives
The aim of the study was antifungal susceptibility testing of yeasts isolates to fluconazole, voriconazole, caspofungin, and anidulafungin.

Methods
From January 2013 through October 2014, 3200 isolates were obtained from hospitalized patients with nosocomial yeast infection. The distribution of the species obtained was Candida albicans (75.4%), C. glabrata (8.2%), C. tropicalis (4.8%), C. kefyr (3.4%), C. parapsilosis (2.5%), C. lusitaniae (1.9%), C. famata (1.4%), C. krusei (1.4%), C. guilliermondii (1%), C. sake (1%), T. asahii (1%) and T. mucoides (1%). MICs of antifungal agents were determined by Etest method.

Conclusions
All the isolates of C. albicans were susceptible to the fluconazole. Higher voriconazole MICs were observed in isolates exhibiting higher fluconazole MICs, frequently in C. krusei and C. glabrata. Isolates with echinocandins MICs suggesting reduced susceptibility were only sporadic cases.
VICISSITUDES IN ADHESION TO DENTURE ACRYLIC SURFACES OF ORAL CANDIDA DUBLINIENSIS ISOLATES FOLLOWING BRIEF EXPOSURE TO SUB-LICIDAL CONCENTRATIONS OF POLYENES, ECHINOCANDINS, AZOLES AND CHLORHEXIDINE GLUCONATE

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Background

Adherence of Candida to denture acrylic surfaces is implicated as the initial process in the pathogenesis of Candida-associated denture stomatitis. This condition can be treated with antimycotic agents belonging to polyene, echinocandin and azole groups of antimycotics. In addition chlorhexidine is also used in its management. In the niches of the mouth the intraoral concentration of these drugs fluctuates considerably due to the dynamics of the oral environment and Candida may undergo only to a brief exposure to therapeutic agents. It has been suggested that Candida dubliniensis may play an important role in the establishment and persistence of C. dubliniensis induced-denture stomatitis.

Objectives

Hence, the effect of brief exposure to sub-lethal concentrations of nystatin, amphotericin B, caspofungin, ketoconazole, fluconazole and chlorhexidine gluconate on adhesion of these isolates to denture acrylic surfaces was investigated.

Methods

After determining the minimum inhibitory concentration of the drugs, twenty oral isolates of C. dubliniensis were exposed to sub-lethal concentrations of these six drugs for one hour. Subsequently, the drugs were removed by dilution and the adhesion of these isolates to denture acrylic strips was assessed by an in vitro adhesion assay.

Conclusions
Compared to controls, exposure to nystatin, amphotericin B, caspofungin, ketoconazole, fluconazole and chlorhexidine gluconate suppressed the ability of *C. dubliniensis* isolates to adhere to denture acrylic surfaces by 74.68%, 74.27%, 73.33%, 57.31%, 44.57% and 56.53% (*P* < 0.001 for all drugs), respectively. Hence, brief exposure to sub-lethal concentrations of antimycotics seems to exert an antifungal effect by suppressing the adhesion to denture acrylic surfaces of *C. dubliniensis* oral isolates.
IDENTIFICATION OF EXTRACELLULAR ASPERGILLUS SPP COMPONENTS THAT INTERACT WITH IMMUNE RECEPTORS.

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**Background**

Aspergillus species are worldwide distributed and ubiquitous in the environment, among this genus relevant industrial and medical relevant species are present. For instance, *A. niger* is used in industrial production systems and has also been reported to cause infections. Extracellular compounds from Aspergillus spp can play various roles during human infections, for example by modulation of immune system response.

**Objectives**

Identify novel extracellular fungal components that interact with immune receptors.

**Methods**

Culture supernatants from various *A. niger* strains (WT and mutants strains) were screened via a Tricolor Assay enabling identification of components that interact with cellular immune receptors. Briefly, the culture supernatant was first co-incubated with a human immune cell mix (neutrophils, monocytes and lymphocytes). Subsequently, fluorescently labeled immune receptor antibodies were incubated with the pre-incubated supernatant-cells mix in a 96-well plate. Subsequently, the plate was centrifuged, washed and cells were fixed. Finally, antibody-antigen interactions were measured using FACS analysis.

**Conclusions**

We did not detect extracellular components in supernatants of *A. niger* wt (N402) that interact with immune receptor. However, a derivative of this *A. niger* wt strain, the mutant D15#26, did produce such components and which appear to be small since a 3 kDa filtrated supernatant was still active. This *A niger* mutant strain lacks extracellular proteases due to a PrtT mutation. However, it also contains additional undefined mutations due to UV mutagenesis and which for example prevents acidification of the medium. We are characterizing this mutant to identify the components and genes that are involved.
FEMS-2638
Fungal pathogenicity in humans

B39 - INVESTIGATION OF DERMATOLOGICAL SPECIMENS FOR SUPERFICIAL MYCOSES
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Background

The term ‘superficial mycoses’ refers to fungal infections usually confined to the outer layers of the skin, hair and nails. Superficial fungal infections are chronic and recurring conditions and are caused by dermatophytes as well as non-dermatophytes.

Infection by dermatophytes is cutaneous and generally restricted to the non-living cornified layers in patients who are immunocompetent. They are generally unable to penetrate tissues which are not fully keratinised. Reactions to infections can range from mild to severe depending upon the host’s immune response, the virulence of the infecting species, the site of infection and environmental factors. The dermatophytes are classified in three genera: *Epidermophyton*, *Microsporum* and *Trichophyton* species.

There are non-dermatophytes that can infect healthy skin, nails damaged by physical trauma, or pre-existing infection with a dermatophyte.

Objectives

Aimed at practising professionals in the field of microbiology, B39 is a diagnostic tool which describes and recommends the procedures that can be used to visualise and isolate dermatophytes, non-dermatophytes and other fungi from skin, nail and hair specimens.

B39 is developed by professionals in the field with specialist input and is intended as a general resource for laboratory staff, clinicians and healthcare commissioners. This document focuses on how detection and accurate identification can aid the management and control of superficial mycoses.

Methods

-

Conclusions
B39 provides guidance on the best minimum practice when investigating infection caused by dermatophytes, non-dermatophytes and other fungi. It provides information on rapid molecular methods and culture testing for these fungi and aims to help drive pathology modernisation by recommending new technologies.
Background
The dermatophyte Trichophyton rubrum is the most common causative agent of dermatophytosis in the world. Due the prominence of dermatophytosis and its socio-economic consequences, there is a need to develop of new antifungal agents. Most of the currently available drugs presents side effects and are ineffective against resistant strains of fungi. The plants are a significant source of biologically active natural products and many chemical structures are models for the synthesis of new drugs. The α-solanine is a glycoalkaloid extracted from potato sprouts (Solanaceae) and has several biological activities, including antimicrobial activity.

Objectives
The aim of this study was to determine the minimum inhibitory concentration (MIC) of α-solanine and to quantify the ergosterol content of T. rubrum exposed to natural compound α-solanine.

Methods
The susceptibility assay was performed using the microdilution test of α-solanine and control fluconazole in the range of to 1000-1.9 μg/mL toward the strain ATCC MYA-3108 of T. rubrum. Ergosterol content was extracted from mycelia exposed for 48 hours to MIC concentration of antifungals and was quantified by spectrophotometry.

Conclusions
The α-solanine compound showed a MIC of 3.12 μg/mL and fluconazole showed a MIC of 63 μg/mL toward the strain ATCC MYA-3108. The α- Solanine and fluconazole reduced 35% and 40.95% of ergosterol content, respectively. The α-solanine showed significant activity toward T. rubrum and reduced the level of ergosterol of cell membrane, which is a specific target of the fungal cell.
Background
Candidal vulvovaginitis is an infection of the vagina’s mucous membranes by *Candida* species. Up to 75% of women will have this infection at some point in their lives, and approximately 5% will have recurring episodes. It is the second most common cause of vaginal inflammation after bacterial vaginosis and usually caused by *C. albicans* but other species, including *C. glabrata*, *C. tropicalis*, *C. krusei* and *C. africana*.

Objectives
The aim of the present study was to molecular identification of *Candida* species isolated from patients with vulvovaginal candidiasis.

Methods
A total of 150 patients refer to Mazandaran University of Medical Sciences from 2013-2014. The vaginal swab specimens were cultured on Sabouraud Dextrose Agar (SDA) and identified based on conventional methods (Germ Tube Test, Chlamydospore production) and subsequently confirmed by PCR-HWP1 (produces 3 different DNA fragments: approximately 780 bp for *C. africana*, 941 bp for *C. albicans* and 250 bp for *C. glabrata*).

Conclusions
The current study shows that *C. albicans* as predominant species isolated from vulvovaginitis. It can be concluded that molecular diagnostic methods are reliable and would be useful for identification of medically important *Candida* species in clinical samples. Therefore considerable attention has been paid to prevention and treatment of microbial growth, which has resulted in improvement of patient management. Although recently molecular approaches have been developed for the identification and discrimination of *Candida* species, novel techniques for the diagnosis based on species level have been established in routine laboratories.
Background
Candida albicans is a frequent component of oral ecology found in up to 75% of humans; particularly among immunocompromised patients. Candida species can cause a multitude of disease ranging from mild oral disease to disseminated candidiasis.

Objectives
In this study, an attempt has been made to determine susceptibility pattern of four antifungal agents against the Candida species isolated from cancer patients with oropharyngeal candidiasis.

Methods
Isolation and identification of Candida spp was done based on standard procedures. Antifungal resistance pattern was carried out according to CLSI guideline. Identification of Candida species were confirm using the 18s Ribosomal RNA.

Conclusions
Leukemia and lymphoma were the most frequent cancer in the studied group, accounting for 17 (34%) and 12 (24%) respectively. A total of 29 Candida spp were isolated from 29 of cancer patients; of which 17 were C. albicans and 12 were C. non – albicans. All the Candida spp were confirmed using the 18s Ribosomal RNA. Among all the Candida spp, Candida non – albicans showed a high resistance pattern to amphotericin B (MIC 07 µg / ml) and ketokenazole (MIC=05 µg / ml). In conclusion, oropharyngeal Candidiasis is a serious infection among cancer patients. The isolated candida spp were resistant to common antifungal agents which may leads to longer hospital stay, more expensive/ toxic drugs and higher mortality. Therefore, interval surveillance is necessary in developing institutional guidelines.
Background

High-performance liquid chromatography (HPLC) is a reliable and robust methodology in medical diagnostics, including the detection of hemoglobin variants, vitamin D levels, and *Trichophyton rubrum* in skin and nail materials. Previous information reported that the latter was reliably identified using xanthomegnin activity via HPLC.

Objectives

In the present study, we aimed to discriminate between the two most common human pathogenic dermatophytic fungi, *T. rubrum* and *T. mentagrophytes* complexes, via the detection of xanthomegnin using HPLC methodology.

Methods

Hence, we investigated 32 reference dermatophyte strains: *Arthroderma* spp. (*n*=7), *T. rubrum* (*n*=18), and *T. mentagrophytes* complex (*n*=7) strains. In addition, we analyzed both clinical samples and strains from 13 samples identified as *T. rubrum* (*n*=9) and *T. interdigitale* (*n*=4) using ITS-sequencing. Twelve of 18 reference *T. rubrum* complex strains produced detectable amounts of xanthomegnin. Moreover, and more importantly, 2 *A. vanbreuseghemii* and 1 *T. interdigitale* reference strains also demonstrated xanthomegnin activity. Furthermore, from a total of 13 clinical samples, 11 clinical samples and 11 clinical strains were positive for xanthomegnin, including *T. interdigitale* strains.

Conclusions

Although xanthomegnin detection is a rapid and inexpensive method, we concluded that it is not a reliable method for discriminating between *T. rubrum* and *T. mentagrophytes* complexes given that only two-thirds of the reference *T. rubrum* complex strains and 3 strains from the *T. mentagrophytes* complex produced xanthomegnin activity. In addition, we believe that the results of the present study should be of interest to scientists and clinicians specifically in the field of medical mycology.
Background

*Candida glabrata* causes almost 20% of all fungal infections in humans. This yeast is distantly related to other medically important *Candida* species (such as the most widespread fungal pathogen *C. albicans*) and significantly differs from those species in the pathogenicity profile. One of essential steps in the microbial colonisation of human organism is the adherence of pathogen to host cells and the supporting extracellular matrix (ECM).

Objectives

*C. glabrata* cell surface is involved in initiating the contact between fungus and host during infection. The current study aims at identifying the major cell wall components that interact with human ECM proteins.

Methods

*C. glabrata* cell wall-associated proteins were isolated with β-1,3- or β-1,6-glucanases and allowed to interact with biotin-labelled fibronectin, vitronectin and laminin. The formed protein-protein complexes were fixed by chemical crosslinking with a heterobifunctional, photoactivatable reagent, sulfosuccinimidyl 2-[(4,4'-azipentanamido)ethyl]-1,3′-dithiopropionate. The stably coupled proteins were adsorbed on streptavidin-conjugated beads and identified by tandem mass spectrometry.

Conclusions

Binding of fibronectin, vitronectin and laminin to *C. glabrata* cell surface was confirmed and several fungal proteins involved in these interactions were identified. Some of them are common binders for all three investigated ECM proteins and predominantly belong to a group of loosely bound with the cell surface, cytoplasm-derived proteins. Examples of these ECM-binding proteins include enolase, alcohol dehydrogenase, glyceraldehyde-3-phosphate dehydrogenase, 6-phosphogluconate dehydrogenase and malate synthase.
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FEMS-2785
Fungal pathogenicity in humans

USE OF PCR-RFLP AND PCR-HWP1 FOR IDENTIFICATION OF CANDIA SPECIES ISOLATED FROM CYSTIC FIBROSIS PATIENTS.

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Background

Due to the predisposing conditions in patients with cystic fibrosis (CF) caused by defective mucociliary clearance facilitates of colonization and invasion with Candida species has dramatically increased. Molecular techniques utilizing amplification of target DNA provide quick and precise methods for diagnosis and identification of Candida species.

Objectives

The aim of current study was identification of the most medically common isolated Candida species from the air way of CF patients by PCR-RFLP and amplification of HWP1 gene.

Methods

A total of 42 CF patients presenting symptoms referred to pediatric respiratory diseases research center were screened for the presence of Candida spp. The isolates initially were phenotypically identified and confirmed by molecular approaches based on restriction fragment length polymorphism (PCR-RFLP) for discrimination of C. albicans of non albicans and amplification of HWP1 gene for discrimination of C. albicans from C. dubliniensis and C. africana was conducted.

Conclusions

Results show that C. albicans was the most frequently isolated species (83.8%) followed by non-albicans included C. parapsilosis (7.1%), C. glabrata (3.2%), and C. tropicalis (3.2%). The restriction patterns of each Candida species were perfectly specific. Since MspI is not able to discriminate between three morphological similar species, C. albicans, C. dubliniensis and C. africana, we used PCR amplification of hwp1 gene, which (7.1%) species from C. albicans identified as C. dubliniensis, however C. africana strains were not found. It can be concluded that molecular diagnostic methods are reliable and would be useful for identification of medically important Candida species in clinical samples.
NEWLY IDENTIFIED CAUSAL AGENTS OF FUNGAL KERATITIS

**Background**

Fungal keratitis is a serious suppurative, usually ulcerative corneal infection which may result in blindness or reduced vision.

**Objectives**

Our aim was to study the spectrum of fungi causing keratitis in South India based on the data collected at the Aravind Eye Hospital and Postgraduate Institute of Ophthalmology, Coimbatore, Tamilnadu between 2005 and 2014.

**Methods**

Fungal strains were isolated from corneal scrapings of keratitis patients. Morphological examination of the isolates and nucleotide sequence-based identification were performed.

**Conclusions**

The most frequently occurring genus was *Fusarium*. Representatives of five *Fusarium* species complexes were detected. Members of *Fusarium solani* species complex (FSSC) were the most frequent, followed by the species complexes *Fusarium dimerum* (FDSC), *Fusarium fujikuroi* (FFSC), *Fusarium oxysporum* (FOSC) and *Fusarium incarnatum-equiseti* (FIESC). *Fusarium napiforme* from FFSC and the sexual fungus *Neocosmospora vasinfecta* from FOSC were firstly detected from keratitis. *Aspergillus* proved to be the second most frequent genus with *Aspergillus*
flavus as the predominant species, followed by *Aspergillus fumigatus* and *Aspergillus terreus*. *Aspergillus tamarii, Aspergillus nomius* and *Aspergillus pseudotamarii* from section *Flavi, Aspergillus tubingensis, Aspergillus brasiliensis, Aspergillus neoniger* and *Aspergillus welwitschiae* from section *Nigri, Aspergillus sydowii* and *Aspergillus variecolor* from section *Nidulantes, Aspergillus amstelodami* from section *Aspergillus, Aspergillus melleus* from section *Circumdati* and *Aspergillus lentulus* from section *Fumigati* of the genus were firstly recognized as potential causal agents of fungal keratitis.

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Fungal pathogenicity in humans

FUSCIA IN HOSPITALIZED PATIENTS WITH PROLONGED FEVER WITH BACTEC METHOD
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Background
Fungemia is a very important infection in terms of incidence and mortality. It can be caused by several species of the genus Candida and should be diagnosed and treated quickly.

Objectives
This study was performed to assess prevalence of fungemia in patients with prolonged fever using two blood culture methods.

Methods
This descriptive study was carried on 330 blood samples obtained from patients with prolonged fever despite widespread antibiotic use in Beheshti hospital of Kashan using both BHI and BACTEC blood culture within 2010-2013.

Conclusions
Among 330 patients with prolonged fever, 16 cases of fungemia were detected which all of them were positive in BACTEC method and 12 were positive in BHI method. All of the isolated fungus were candida and the most frequent agent of candidemia was Candida Albicans. The predisposing underlying factors were malignancy, diabetes, using of widespread antibiotics and corticosteroids, trauma, and surgery. The rate of fungemia was 4.8% (16/330). We found Candida Albicans as the most common causative agent of candidemia in our samples.
FEMS-2630
Fungal pathogenicity in humans

CASE SERIES OF RHINOCEREBRAL MUCORMYCOSIS
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Background
Rhinocerebral mucormycosis (RCM) is a rare, fulminating opportunistic fungal infection caused by a fungus of order Mucorales

Objectives
The purpose of this paper is to draw attention to the clinical presentation of RCM and to emphasize need for high index of suspicion in diagnosis and treatment

Methods
This was a case series study conducted in Beheshti Hospital of Kashan, from 2001 to 2013. The charts of identified patients were reviewed for the following information: demographics, predisposing conditions, anatomical location of infection, treatment, length of hospital stay

Conclusions
Cases were between 21 to 88 year old. The most patient were female (sex ratio: 2). The clinical sign and symptoms of patients were hematemesis, epistaxis, melena, headache, pain, swelling and tenderness on the face and paranasal sinuses, malaise and weakness, fever, blurred vision, periorbital edema, black secretion and necrosis of nasal septum or hard palate, cranial nerve palsy. All the patients had history of diabet. CT scan of PNS revealed consolidation, increased thickness, necrosis and disruption of mucosal wall of sinuses. Amphotericin B was prescribed for all the patients. The length of hospital stay was more than 20 day. Blindness occurred in one patients. 3 patients died and the rest recovered

The diagnosis of rhinocerebral mucormycosis should be considered in the clinical setting of necrotic sinusitis in diabetic patients. Early diagnosis and treatment are crucial factors leading to a good outcome.
Background
The most common fungal infections in humans are candidiases caused by *Candida* yeast. A major, highly conserved yeast cell wall (CW) protein Bgl2p, which can also be secreted into the growth medium, plays a significant role in candidiasis. It was shown that the injected into a mouse's bloodstream *C. albicans* with *BGL2* deletion possessed decreased infectivity compared to the one without *BGL2* deletion (Sarthy et al., 1997). On the other hand, high level of antibodies against Bgl2p is prognostic factor, which defines the risk of mortality (Pitarch et al., 2006). Authors proposed to use Bgl2p from *C. albicans* for vaccine development. Earlier we demonstrated that Bgl2p from *S. cerevisiae* CW (Bgl2p homolog from *C. albicans*) could reveal amyloid properties (Kalebina et al., 2008). At the beginning of our work the process of Bgl2p amyloid formation remained unclear.

Objectives
It was suggested that Bgl2p as well as many other proteins of microorganisms appeared as a virulence factor due to its amyloid forming ability. In order to verify it we needed to study whether Bgl2p was able to form fibrils under conditions similar to those of a bloodstream. Primarily we were interested in investigating the influence of two factors on Bgl2p fibrillation: the pH value within a neutral range and the presence of regulatory molecules such as polyphosphates.

Methods
We investigated Bgl2p from non-pathogenic *S. cerevisiae* as a model using fluorescence confocal and electron microscopy, fluorescence spectroscopy, bioinformatics analysis.

Conclusions
Bgl2p fibril formation ability was strongly regulated by pH value (Bezsonov et al., 2013), polyphosphates, divalent cations and buffer components. The results obtained allowed us to conclude that Bgl2p might exist in fibrillar form under bloodstream-like conditions. The described Bgl2p amyloid formation ability is important for further fighting the candidiasis including vaccine development, using Bgl2p from non-pathogenic *S. cerevisiae*.
Fungal pathogenicity in humans

ANALYSIS OF THE PROFILE OF PARACOCCIDIOIDES SPP. IN RESPONSE TO THE DRUG BY TRANSCRIPTIONAL/PROTEOMIC APPROACHES AND IDENTIFICATION OF COMPOUNDS WITH ANTIFUNGAL POTENTIAL.

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Background

Paracoccidioides spp, a complex of several phylogenetic species, is the agent of paracoccidioidomycosis (PCM). PCM is endemic in Latin America, with 80% of cases reported in Brazil, where it is the eighth-leading cause of mortality among infectious and parasitic diseases, establishing it as a serious public health problem. The use of small molecules of natural origin as medicines or as prototypes of drugs is a well established theme in the drug development.

Objectives

Argentilactone, bioactive metabolite isolated from \textit{Hyptis ovalifolia} and inhibited the isocitrate lyase from \textit{Paracoccidioides}. Thiosemicarbazide is a small molecule terpene camphene derivative that inhibits the growth of \textit{Trichophyton mentagrophytes}. Indole alkaloids due their already described antifungal activity, we have included several carboline alkaloids isolated from \textit{Rubiaceae} species in our screening antifungal program aiming to discover compounds with potential to inhibit malate synthase from \textit{Paracoccidioides}.

Methods

In this study we performed the transcriptome, proteome analysis and virtual screening of \textit{Paracoccidioides spp.} yeast cells growth in the presence of compounds and/or itraconazole and amphotericin B.

Conclusions

A total of 1,058 genes were identified, of which 208 were up-regulated and 850 down-regulated in argentilactone response. Four indol alkaloids showed ability to reduce the \textit{PbMLSr} activity and inhibit the adhesion of the fungus and \textit{PbMLS} to extracellular matrix components. In our proteomic analyzes various classes of proteases were suppressed in the presence of thiosemicarbazide, were identified and sequenced 11
differentially expressed proteins in the condition with itraconazole, and 29 for the condition treated with amphotericin B.
ORAL MICROFLORA AND THEIR RELATION TO RISK FACTORS IN HIV+ PATIENTS WITH OROPHARYNGEAL CANDIDIASIS

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Background
Oropharyngeal candidiasis (OPC) is considered an opportunistic infection caused by a ubiquitous fungal organism that is routinely seen as normal oral flora.

Objectives
The purpose of this study was to determine the prevalence of oral microflora and association of oral candidiasis and multiple risk factors in HIV+ patients.

Methods

The present study included 100 HIV-infected patients participated in Imam Khomeini Hospital, Tehran, Iran for OPC and HIV. Samples were obtained from the oral cavity and direct microscopic examination, gram staining and culture on standard microbiological media were performed in all patients. CD4+ cell count/CD4+ percentage were also calculated.

Conclusions

A total of 460 bacterial colonies were obtained and Streptococcus mutans (15.4%) was the most frequently isolated species in the HIV+ patients. In addition, 254 yeasts (from 4 different genera) were isolated from the patient understudy. Candida species (94.4%) were the most frequently obtained genera, followed by Saccharomyces (2.4%), Kluyveromyces and Cryptococcus (1.6% for both) species. Candida albicans (37.2%) was the most common species isolated from HIV+ patients with OPC. The mean CD4+ cell counts were 154.5 cells/µL, with a range of 8 to 611 cells/µL. Thirty percent (30%) patients had a CD4+ cell count between 101 and 200 cells/µL (28.7% of total yeasts isolated). Our results showed that yeasts of the genus Candida were isolated at a comparable rate from the oral cavity of HIV+ patients and there was no significant difference of the variables CD4+ cell count and yeast counts.
Background
Intracellular proteinase A is an aspartic enzyme in *Candida albicans* that is expressed by *APR1* gene.

Objectives
The aims of this study were to evaluate and to compare *APR1* gene expression in *C. albicans* strains isolated from Malaysian and Iranian patients and mice infected by *C. albicans* obtained from Malaysian and Iranian patients.

Methods
The evaluation of *APR1* gene expression was performed using reverse transcriptase-polymerase chain reaction (RT-PCR) with two-pair primers and 18S rRNA as housekeeping gene. The expression of *APR1* gene and 18S rRNA was determined on agarose gel electrophoresis.

Conclusions
The results did not show any significant difference in *APR1* gene expression between *C. albicans* isolated from mice blood infected by *C. albicans* isolated from Malaysian and Iranian patients (*p*>0.05). The expression of *APR1* gene in *C. albicans* strains isolated from Malaysian patients was more than Iranian patients (*p*<0.05). In addition, the gene expression in *C. albicans* strains obtained from mice blood was significantly higher than *C. albicans* strains isolated from both Malaysian and Iranian patients (*p*<0.05). Considering to the important role of intracellular proteinase A in *C. albicans*, *APR1* gene as producer of this enzyme can act as an important gene in virulence of *C. albicans* in different environmental situations.
ANTIFUNGAL ACTIVITY OF ZATARIA MULTIFLORA, PLARGONIUM GRAVEOLENS AND CUMINUM CYMINUM ESSENTIAL OILS TOWARDS THREE SPECIES OF MALASSEZIA ISOLATED FROM PATIENTS WITH PITYRIASIS VERSICOLOR

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Background

Malassezia species have been recognized as members of the microbiological flora of human skin that cause skin diseases such as pityriasis versicolor (PV).

Objectives

To investigate the anti-Malassezia activities of Zataria multiflora, Plargonium graveolens and Cuminum cyminum essential oils (EOs) against different pathogenic Malassezia species isolated from patients with pityriasis versicolor (PV).

Methods

The EOs were obtained by water-distillation using a Clevenger-type system. Anti-Malassezia activity against Malassezia species was carried out using disk diffusion method in Dixon agar.

Conclusions

The main oil components were carvacrol (61.3%) and thymol (25.2%) for Z. multiflora, α- pinene (30%) and limonene (21%) for C. cyminum and citronelol (28.2%) and geraniol (22.1%) for P. graveolens. The three Malassezia species showed a similar susceptibility to the three plants tested, C. cyminum (mean value: 48.3 mm) being the most active, followed by Z. multiflora (mean value: 28.1 mm) and P. graveolens (mean value: 26.1 mm). This study indicated that Z. multiflora, P. graveolens and C. cyminum EOs have considerable anti-Malassezia activities, deserving further investigation for clinical applications for the treatment of PV.
DELETION LIBRARY REVEALS POTENTIAL GENES INVOLVED IN THE
VIRULENCE OF CANDIDA PARAPSILOSIS

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Background

Since the 1980s, fungi have emerged as causes of human disease. Some fungal species are opportunistic pathogens, that are only able to cause diseases in individuals with damaged physical barriers or/and altered immune status. The incidence of C. parapsilosis caused candidemia has dramatically increased over the past decade, therefore, the understanding of the pathomechanism of this species becomes urgent.

Objectives

Despite the growth of genome sequence information, a large number of fungal genes remain uncharacterized. In our previous work several fungal transcriptional factors have been identified using RNA-Seq data that were over expressed during host-pathogen interactions. Based on these data we aimed to generate a knock out library to be able to analyze these genes individually.

Methods

Fusion PCR was applied to generate gene specific deletion constructions to disrupt genes from the genome of a C. parapsilosis double auxotrophic strain. The growth abilities of the null mutant strains were tested in different conditions such as different medias and temperatures along with survival in the presence of various stressors.

Conclusions

We have found null mutants that show differences in appearance such as increased pseudohyphae formation, regressed growth on different temperatures and alkali-phobic phenotype. Comparison of the virulence of these deletion strains using infectious models is now in progress. In the future with the use of this method we are able to identify key regulatory factors that may play a role in the virulence of C. parapsilosis during host-pathogen interactions.
THE CFEM HEME-IRON ACQUISITION NETWORK IN PATHOGENIC FUNGI

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Background
Iron acquisition represents a challenge for all organisms due to the low solubility of iron under oxidizing conditions. This challenge is particularly acute for pathogenic microorganisms, due to the iron withholding mechanisms deployed by the host. Many microbial pathogens have therefore evolved mechanisms to extract iron from hemoglobin, the largest iron store in the host, by removing the heme cofactor from the globins and transferring it to the microbe’s cytoplasm. The fungal pathogen \textit{Candida albicans} uses a family of conserved extracellular proteins for that purpose, the CFEM proteins \textsuperscript{1}.

Objectives
The cell wall-anchored CFEM proteins Rbt5 and Pga7 are both able to extract heme from hemoglobin \textsuperscript{2}. The heme can then subsequently be transferred from one CFEM protein to the next \textsuperscript{2} until it reaches the plasma membrane, whereupon it is endocytosed by a mechanism involving the ESCRT pathway \textsuperscript{3}. A third, secreted CFEM protein, Csa2, is shown here to also participate in the heme-iron acquisition relay.

Methods
Biochemical and structural analysis

Conclusions
Biochemical and structural analysis of Csa2 reveals that CFEM proteins are unrelated, at either the sequence or structure level, to the known bacterial heme-scavenging proteins, suggesting that microbes have evolved several different independent solutions to the heme-iron acquisition problem.
Background

*Trichoderma* includes a great diversity of species, some of them with the ability to control the growth of fungal phytopathogens. Many of these strains produce secondary metabolites that are able to inhibit the growth of their fungal prey. However, pathogens can also produce metabolites that in some cases belong to the same family of those produced by *Trichoderma*.

Objectives

In the present work we study the interaction between Harzianum A (HA) produced by *T. arundinaceum* (Ta37) and botrydial (BOT) produced by *Botrytis cinerea*. HA exhibits antifungal activity, and BOT is a strong virulence factor in plant disease.

Methods

A procedure of confrontation between these two strains was implemented, followed by the analysis of gene expression by real-time PCR using mycelia isolated from the confrontation areas. To better characterize the regulatory responses produced by HA, the TaDTri5 mutant, isolated by disruption of *tri5* gene and blocked in the synthesis of HA, was also used.

Conclusions
It was observed that HA induced expression of genes involved in BOT biosynthesis when HA was added to *B. cinerea* growing alone. However, in confrontation experiments against TaDTri5 the expression of these genes was higher than in confrontation with Ta37, indicating that the mutant must produce some metabolite/s, responsible for such induction. A different metabolite production pattern was described for Ta37 and TaDTri5\(^{(a)}\), which is in agreement with the proteomic characterization of these two strains, suggesting that genetic disruption of *tri5* had a systemic effect of the fungal physiology not only restricted to the terpene pathway. 

(a) Doi: 10.1111/1462-2920.12514.
FEMS-1820
Fungal pathogenicity in plants

CHANGES OF THE ALTERNATIVE OXIDASE ACTIVITY OF BOTRYTIS CINEREA ISOLATES UNDER VARIOUS CONDITIONS
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Background

Botrytis cinerea (teleomorph Botryotinia fuckeliana [de Bary] Whetzel) is the causal agent of grey mould on a wide range of crops. Some of the pesticides used for chemical control inhibit the respiration electron transport chain. Resistance develops rapidly due to the high genetic variability of B. cinerea. In addition, a functional alternative oxidase (AOX) may also play a role in the survival of the fungus. Expression of AOX is often increased under biotic and abiotic stress, providing the organism adaptation capacity to different environmental conditions. It may also provide the respiratory electron transport chain an extra capacity when the cytochrome pathway is inhibited or saturated.

B. cinerea includes two cryptic species: group I (B. pseudocinerea) and group II (B. cinerea sensu stricto), which could differ in their pathogenicity, fungicide resistance, host variability and genetic diversity.

Objectives

In this study the differences between the alternative respiratory activity of B. cinerea belonging to group I and II were investigated. We also studied the effects of osmotic stress and the temperatures on the respiratory system.

Methods

Measurements were carried out and analyzed with polarographic oxygen electrode system. The cytochrome pathway was inhibited with KCN while SHAM was used for blocking the alternative respiration.

Conclusions

More intense alternative respiratory activity was observed in group II isolates under optimal growth conditions. The alternative pathway was more profoundly induced by osmotic and temperature stress compared to the cytochrome pathway, and this
correlated well with increased transcript formation of the aox1 gene encoding the AOX enzyme.
FEMS-1858
Fungal pathogenicity in plants

CHARACTERIZATION OF COMPONENTS PRODUCED BY AUREOBASIDIUM PULLULANS, ANTAGONISTIC AGAINST MANGO ANTHRACNOSE DERIVED BY COLLETOTRICHUM GLOEOSPORIOIDES
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Background

Mango anthracnose derived from Colletotrichum gloeosporioides reveal severe symptoms in terms of surface pitting, skin browning in the surface of mango fruits.

Objectives

In this study, We explore to screen the antagonistic yeasts against mango anthracnose and study the antagonistic mechanisms and characteristics of the components produced by the antagonistic yeasts.

Methods

Seven strains of C. gloeosporioides were isolated from 8 infected mangoes. On the pathogenic test, strain WU2L01-1 could infect mangoes and reveal pitting and skin browning as that of anthracnose, indicating the strain is a phytopathogen to mango.

Conclusions

Twelve of 578 yeasts exhibited antagonistic activity against C. gloeosporioides. These strains were classified into the member of the genus Aureobasidium, Cryptococcus and Jaminaea species, based on the sequences of the LSU D1/D domain of rRNA gene. The antagonistic components were characterized by analyzing the activity of degradative enzyme, siderophore and glycolipid. All the experiments demonstrated that these strains cannot produce high activity of siderophore and degradative enzymes, β-1.3–glucanase, chitinase, amylase, and ?. While, the extracts from culture broth derived from different yeast strains by methanol, ethyl acetate or dichloromethane were confirmed to have the antagonistic activity against C. gloeosporioides. The optimum medium was tested and obtained for producing antagonistic components. Till now, strains NN5L07, NN5L08, NN9L05,
GY10L05 and HM4L10 and could produce higher productivity of antagonistic components based on these results.
Background
Brazil is responsible for a third of world production of orange, as well as for 80% of international orange juice market. The European Union is the main market in this Brazilian agribusiness. But, the citrus crop is affected by a range of pathogens that decrease the production and the business viability. One of these diseases is Postbloom Fruit Drop (PFD) caused by one fungal species from the *Colletotrichum acutatum* species complex, which causes serious losses in citrus fruit production. Many works are being developed to study and control this phytopathogenic fungus.

Objectives
An alternative method to investigate the fungus infection process is the use of fluorescence protein.

Methods
For that, we applied the agrotransformation system to transform a *Colletotrichum* isolate from citrus symptomatic blossom, proven as pathogenic.

Conclusions
The methodology became efficient to insert genes in the fungus. The transformers have the ability to express the red fluorescent protein due the insertion of DsRed gene into the fungal genome. They were evaluated in fluorescence microscopy and is possible to identify spores, appressoria and fungus hyphae. Therewith, we efficiently applied the transformation methodology by *Agrobacterium tumefaciens* in the *Colletotrichum* citrus pathogen and these transformants can be used to infection process studies and evaluated for eventual pathogenic alterations as result of insertion mutagenesis.
FEMS-0744
Fungal pathogenicity in plants

ESTABLISHMENT OF GENETIC MANIPULATION SYSTEM FOR USTILAGO ESCULENTA
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Background: Ustilago esculenta is a fungal endophyte that triggers stem swelling of Zizania latifolia resulting in an edible aquatic vegetable. To date, characterization of the interaction between U. esculenta and Z. latifolia remains descriptive due to a lack of genetic manipulation and re-infection strategies.

Objective: In this study, a transformation system for U. esculenta was developed based on methods developed for U. maydis and evaluated.

Methods: To improve protoplast yield, regeneration rate and transformation efficiency, we systematically adapted several factors, including enzyme digesting conditions, osmotic pressure stabilizers, vectors and antibiotics. The adapted transformation protocol was confirmed by knocking out the UeICL (coding isocitrate lyase) gene and over-expressing the EGFP gene. Correct transformants were detected by PCR and fluorescence microscopy analysis.

Conclusions: The protocol was optimized that 100 mg fresh fungus yielded the maximum number of protoplasts after digested for 3 hours at 30°C in 2 ml SCS solution containing 15 mg/ml Lywllzyme with the osmotic pressure stabilizer of 0.4 M sucrose. The protoplasts obtained highest regeneration rate when growing on Reg-light-Agar medium containing 0.8 M sucrose. BsaI-mediated Golden Gate Reactions and counter-selection with hygromycin of 50 μg/ml or carboxin of 5 μg/ml were successfully used in constructing plasmids and identifying mutants. Similar to most filamentous fungi, flank length around 1kb with homologous ends was efficient for U. esculenta. Some sequences of U. maydis, like Potef and NLS (nuclear localization signal), worked well in U. esculenta. We successfully established a genetic manipulation protocol for U. esculenta through protoplast transformation.
Background
Anthracnose cassava incited by Colletotrichum is a prevalent disease in cassava producing regions.

Objectives
The aim of this study was to identify the Colletotrichum associated with cassava in Thailand.

Methods
Cassava anthracnose symptoms (circular, sunken lesions with spore masses produced in black acervuli) were observed on cassava leaves in Nakhon Sawan, Thailand. Four single-spore isolates were derived from fungal cultures isolated from 4 different infected cassava samples.

Conclusions
Colonies grown on HPDA changed from white to grey with an average colony diameter of 8.45x8.60 cm after 7 days. Their conidia were cylindrical and 4.75x 14.00 µm in size. To test the four isolates for pathogenicity, after 3 days, typical anthracnose symptoms had developed on the inoculated cassava leaves but not on that of the control treatment. The pathogen was re-isolated from the inoculated cassava but not from the control cassava. Based upon these morphological and cultural characteristics, the 4 isolates were initially identified as *C. gloeosporioides* f. sp. *aeschnomene*. To confirm identity of the four isolates, their internal transcribed spacer (ITS) regions of ribosomal DNA (rDNA), were amplified with ITS3 and ITS4 universal primers. The amplified DNA sequences of these isolates shared 99% homology with the ITS sequences of *C. gloeosporioides* f. sp.aeschnomene in the GenBank (JX131331, KC816043, JX139578 and JX258732). To our knowledge, this is the first report of *C. gloeosporioides* f. sp. *aeschnomene* causing anthracnose on cassava.
Background

“Penja Pepper” (Piper nigrum L.) is cultivated in the Penja region of Cameroon. It is famous for its exceptional aroma and taste. However, in Cameroon pepper is attacked by various diseases. Moreover, the support tree used for growing pepper vines in Cameroon, Spondias mombin, is also subject to disease attacks.

Objectives
In order to maintain the viability of Penja Pepper cultivation, these diseases must be identified and controlled. The aim of this study was to identify the principal pathogens of black pepper vines and its support tree.

Methods
Samples from pepper vines, S. mombin and the soil were collected from various fields in the Penja region. Using a combination of baiting and selective agar medium techniques several potential causal agents were isolated. The isolated microorganisms were identified based on morphology as well as ITS sequencing.

Conclusions
Eight oomycete isolates were obtained from pepper and 18 basidiomycete isolates from either pepper or S. mombin. The eight oomycete isolates of Pepper were tentatively identified as Phytopythium vexans and/or P. cucurbitacearum. The morphological identification of the basidiomycete isolates show that it is an Armillaria sp. Pathogenicity tests were carried out with 180 days old pepper plants. The plants were inoculated with discs of mycelium introduced in the soil and on the collar. This is the first time that Armillaria is found to be pathogenic to pepper as well as its support tree.
tree *Spondias mombin*. The importance of these finding for controlling these diseases to safeguard the sustainable production of the Penja Pepper of Cameroon is discussed
Background
Antracnose which is caused by *Colletotrichum* spp. is the major disease of tomato crops around the world. Antifungal substances have been reported as biocontrol agents against several phytopathogenic fungi. Previously, *Bacillus* sp. M10 was found to inhibit the growth of several fungal pathogens of tomato including *Colletotrichum capsici*.

Objectives
To purify and identify the antifungal protein from *Bacillus* sp. M10, and subsequently determine the effects of the antifungal protein to the fungus and its ability to prevent and/or protect tomato fruits from the post-harvest anthracnose disease caused by *C. capsici*.

Methods
Ammonium sulfate precipitation and DEAE anion exchange chromatography were performed to purify the antifungal protein. Excised band from SDS-PAGE was analyzed by MALDI-TOF MS. Light and scanning electron microscopy were used to observe the effects of the protein to the fungus. Finally, prevention and protection of postharvest anthracnose disease using the antifungal protein were evaluated.

Conclusions
After 40–60% ammonium sulfate precipitation and DEAE anion exchange chromatography, the antifungal protein was enriched 11-fold and was found to retain a specific activity of 189.57 AU/mg. SDS-PAGE revealed that the purified protein has a molecular weight of ~55.4 kDa. MALDI-TOF MS of the excised, enriched protein band revealed the closest similarity to the vegetative catalase (KatA) from *Bacillus amyloliquefaciens*. Microscopic observation showed that the protein induced abnormal hyphal elongation and conidial swelling and rupture. The protein caused a marked inhibition of anthracnose development on tomato fruits. Thus, this putative catalase may potentially be useful as a biocontrol agent against anthracnose disease.
FEMS-2415
Fungal pathogenicity in plants

AUXIN ROLE AND PRODUCTION IN CLAVICEPS PURPUREA
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Background

Biotrophic fungus *Claviceps purpurea* is known as a parasite of monocots, invading mainly rye florets (*Secale cereale*). The late infection is characteristic by the formation of dark purple sclerotia, which are the source of ergot alkaloids, substances used in pharmaceutical industry. During the infection, host plant does not show any symptoms of immune response and fungal hyphae grow unrecognized through the plant tissue. Plant defence responses are driven by phytohormone crosstalk and the fungus might utilize its own phytohormone biosynthesis for the host manipulation. Preliminary results showed that mycelium of *Claviceps purpurea* contains high amount of major auxin – indole-3-acetic acid (IAA).

Objectives

The aim of this work is to characterize an auxin biosynthesis in *Claviceps purpurea* and to determine an auxin role in *Claviceps* infection strategy

Methods

IAA concentration and its precursors and conjugates was analysed by TLC and HPLC coupled with the MS detection. Putative genes participating in different auxin biosynthesis pathways and translocation mechanism were deleted by the homologous recombination and transformants have been characterized upon the infection and in axenic cultures.

Conclusions

Genome data mining and measurements of auxin biosynthesis intermediates identified two possible pathways of IAA biosynthesis in *Claviceps purpurea* based on plant and bacterial models. Detailed characteristic of several deletion mutants and WT showed interested features indicating that auxin is secreted from fungal tissue upon the infection and actively translocated in fungi itself.
ROLE OF TETRAZOLIUM BROMIDE REDUCTION ASSAY IN DIAGNOSIS OF SUSPECTED CASES OF MULTIDRUG RESISTANT(MDR)PULMONARY TUBERCULOSIS

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Background

Early detection of multidrug resistant tuberculosis (MDR-TB) is of primary importance for both management & infection control. It is complicating TB control efforts in several low- & middle-income countries. Colorimetric method is one of the useful techniques for early detection of MDR-TB.

Objectives

To evaluate drug susceptibility testing by 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay and proportion method to diagnose suspected cases of MDR-TB.

Methods

Sputum from 37 previously treated smear positive patients with pulmonary tuberculosis having clinical suspicion of MDR-TB were included in the study. Culture from Lowenstein-Jensen(LJ) media was inoculated into 7H9 broth and after 7days the inoculum was tested for isoniazid(INH) & rifampicin(RIF) susceptibility testing by MTT assay. The formation of formazan product after addition of MTT (after 7days) was taken as positive for Mycobacterium tuberculosis.

Conclusions

In this study 18 cases were sensitive and 15 cases resistant to isoniazid by MTT assay. Twenty one cases were sensitive and 12 cases resistant by drug susceptibility testing of rifampicin by MTT assay. The sensitivity & specificity of MTT assay for INH was 90% & 100% and for RIF 90.90% & 90.90% respectively. The MTT assay is a simple, low cost susceptibility test that can be completed in approximately 7 days. Due to its high levels of agreement with proportion method, the MTT method has the potential to provide rapid detection of INH & RIF-resistant M. tuberculosis in clinical use.
Background
Drug resistance in tuberculosis (TB) is a growing global problem.

Objectives
The aim of this study was to determine the resistance rates of Mycobacterium tuberculosis complex (MTC) isolates to major antituberculosis drugs (ATBDs) (isoniazid, rifampicin, ethambutol, streptomycin) within six years period in a tertiary referral hospital.

<table>
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<th>Clinical Sample Counts</th>
<th>2009</th>
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<th>2011</th>
<th>2012</th>
<th>2013</th>
<th>2014</th>
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<tr>
<td>Isolate Count (%)</td>
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<td>50 (4)</td>
<td>25 (1.6)</td>
<td>33 (1.6)</td>
<td>56 (3.1)</td>
<td>34 (1.5)</td>
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<td>Resistant Isolates (%)</td>
<td>26 (31.7)</td>
<td>22 (44)</td>
<td>5 (20)</td>
<td>11 (33.3)</td>
<td>13 (23.2)</td>
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<td>Single INH Resistance (%)</td>
<td>11 (13.4)</td>
<td>7 (14)</td>
<td>1 (4)</td>
<td>5 (15.2)</td>
<td>4 (7.1)</td>
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<td>Single RIF Resistance (%)</td>
<td>1 (1.2)</td>
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<td>Single EMB Resistance (%)</td>
<td>3 (3.7)</td>
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<td>Single SM Resistance (%)</td>
<td>0 (0)</td>
<td>1 (2)</td>
<td>1 (4)</td>
<td>0 (0)</td>
<td>2 (3.6)</td>
<td>2 (5.9)</td>
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<tr>
<td>INH+RIF (%)</td>
<td>2 (2.4)</td>
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<td>0 (0)</td>
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<tr>
<td>INH+RIF+SM (%)</td>
<td>0 (0)</td>
<td>1 (2)</td>
<td>1 (4)</td>
<td>3 (9.1)</td>
<td>5 (8.9)</td>
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<tr>
<td>INH+RIF+EMB (%)</td>
<td>5 (6.1)</td>
<td>2 (4)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>INH+RIF+SM+EMB (%)</td>
<td>3 (3.7)</td>
<td>0 (0)</td>
<td>2 (8)</td>
<td>2 (6.1)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Total MDR (%)</td>
<td>10 (12.2)</td>
<td>4 (8)</td>
<td>3 (12)</td>
<td>5 (15.2)</td>
<td>5 (8.9)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Total PDR (%)</td>
<td>1 (1.2)</td>
<td>2 (4)</td>
<td>0 (0)</td>
<td>1 (3)</td>
<td>1 (2)</td>
<td>1 (2.9)</td>
</tr>
</tbody>
</table>


Methods
Clinical specimens were cultured and the susceptibilities of isolates to the major ATBDs were tested using the BACTEC MGIT 960 system. In a period of six years, 280 MTC strains are isolated from 10121 clinical samples. A decreasing trend was found on the MTC isolation rates from clinical samples with years (7.5% in 2009 to 1.5% in 2014). Of the 197 (70.4%) isolates were found sensitive to all four tested ATBDs and 83 (29.6%) of the isolates were found resistant to at least one ATBD.
Rate of single drug resistance for isoniazid was 10.4% (n=29), for ethambutol 3.9% (n=11), for streptomycin 2.1% (n=6) and for rifampicin 1.4% (n=4). Resistance rates of isoniazid, which was found as the most resistant drug among others, were decreasing (13.4% to 2.9%) while only rifampicin resistance was increasing (1.2% to 5.9%) by years. Rate of at least isoniazid and rifampicin resistant -multidrug resistant (MDR)- isolates was 9.6% (n=27). Number of MDR isolates was decreasing by years.

Conclusions
Even though TB rates are decreasing by years, persistence of high isoniazid resistance rates and increasing resistance of rifampicin requires more strict applications of precautions and limitation of rifampicin use both for treatment and prophylactic purposes in other diseases than TB.
EXPLORING THE EVOLUTIONARY SUCCESS OF MYCOBACTERIUM TUBERCULOSIS BY INVESTIGATION OF RARE CLINICAL ISOLATES WITH SMOOTH COLONY MORPHOLOGY

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Background

*Mycobacterium tuberculosis*, one of the most successful human pathogens, and other members of the *M. tuberculosis* complex (MTBC) show a mainly clonal population structure with 99.9% of genetic similarity. In contrast to these strains with predominantly rough colony morphology, rare clinical isolates with smooth morphology (*M. canettii*) show a greater genetic divergence yet are less virulent and less persistent.

Objectives

In order to better understand the mechanism underlying the smooth/rough variation and its implication in virulence, we are currently investigating smooth and rough morphotypes of *M. canettii* strains in more detail.

Methods

Stable smooth and rough morphotypes of different *M. canettii* strains were isolated after several passages on solid medium and were subjected to a whole genome sequence analysis by Illumina technology. Single-end short reads were aligned against the reference genome by using SHRiMP. Lipid analysis was done by chloroform/methanol/water extraction followed by thin layer chromatography.

Conclusions

Comparative genome analysis of the smooth and the rough morphotype revealed a difference in the *pks5* gene locus, which codes for a polyketide synthase that was found to be involved in lipooligosaccharide (LOS) synthesis in the fish pathogen *M. marinum*. More detailed analysis of this locus suggested the importance of *pks5* and a gene of the *pap* family (polyketide synthase associated protein) to be involved in
the phenomenon. Thus, the *pk5* locus seems to be involved in the smooth/rough variation in *M. canettii* through production of lipooligosaccharides, which could also play a role in the reduced virulence phenotype.
IDENTIFICATION OF NONTUBERCULOUS MYCOBACTERIA IN POTABLE AND RECYCLED WATER IN MEXICO CITY UNIVERSITY

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Background

Nontuberculous mycobacteria (NTM) are ubiquitous microorganisms that can act as human pathogens in both, immunocompromised and immunocompetent individuals. Because until now it has not been described human-to-human transmission, the main source appears to be the exposure to water containing the microorganisms. All over the world an increasing number of NTM species with different profiles causing human disease represent a public health threat. However, a fast and reliable method to identify NTM in the water distribution system is lacking. The water distribution system and the recycled water for irrigation of Ciudad Universitaria campus of Universidad Nacional Autónoma de México (UNAM) were screened for NTM. We compared hsp65 gene and 16S rRNA-23S rRNA (ITS) sequence in its ability to distinguish between NTM closely related species.

Objectives

The aim of this work was to compare the ITS and hsp65 gene PCR-RFLP to identify NTM species isolated from recycled water and human use water in distribution systems of UNAM

Methods

56 recycled water samples and 163 potable water samples NTM’s were grown in Lowenstein-Jensen medium and Middlebrook 7H10 medium. DNA from the isolated strains were identified by ITS and hsp65 gene PCR-RFLP specific sequences. Amplifications were digested by HaeIII and CfoI for ITS and HaeIII and BstEII for hsp65 gene. The electrophoresis gels of digested products were analysed with the Roth and Telenti algorithm.

Conclusions

Markers ITS and hsp65 gene PCR-RFLP identified twelve species. Moreover the use of ITS complemented the hsp65 restriction profiles. The sequences used are a good molecular tool for NTM species and subspecies identification.
THE ANALYSIS OF PEDIGREE AMONG MYCOBACTERIUM AVIUM SPP. PARATUBERCULOSIS INFECTED DAIRY HERD
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Background
Johne’s disease, chronic granulomatous enteric disease in dairy cattle, is caused by Mycobacterium avium spp. paratuberculosis (MAP) infection. The prevalence of the pathogen in US dairy farm is an approximately 68% and the pathogen induced $200 to $250 million economic loss in US dairy industry due to decreasing milk production and increasing cow replacement costs.

Objectives
Analysis of MAP positive pedigree for the evaluation of vertical MAP infection comparison with horizontal infection among dairy herd last 10 years.

Methods
The serum MAP antibody detection enzyme-linked immunosorbent assay (ELISA) was applied for the identification of MAP infection among dairy herd and the test was performed biannually. The MAP positive or negative dam, offspring, sibling and offspring of sibling were recorded for 10 years (2005-2015).

Conclusions
The MAP positive or negative pedigree was constructed based on the ELISA test record and the association of each MAP positive individuals among dairy herd was not related with each other. The result from this study could suggest that the horizontal infection would be more considered MAP transmission rather than vertical infection among dairy herds, and it could provide a guideline of future MAP intervention program among dairy herds.
MOLECULAR DETECTION AND CHARACTERIZATION OF RESISTANT GENES IN MYCOBACTERIUM TUBERCULOSIS COMPLEX FROM DNA ISOLATED FROM TUBERCULOSIS PATIENTS IN THE EASTERN CAPE PROVINCE SOUTH AFRICA

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Background
Tuberculosis (TB) in both animals and humans is caused by Mycobacterium tuberculosis complex (MTBC) primarily transmitted by inhalation of aerosolized droplets containing the organism. Multi-drug resistance (MDR) and extensive drug resistance (XDR) are evolutionary features of Mycobacterium tuberculosis to subvert the antibiotic regimes in place.

Objectives
The heavy burden of TB in South Africa motivated the investigation of MTBC prevalence among TB patients, amplification and sequencing of DNA amplicons known to confer resistance to TB drugs in Port Elizabeth.

Methods
Three thousand eight hundred and ten (3810) sputum specimens were processed and DNA isolated from the Eastern Cape Province, South Africa. DNA was amplified using the Seeplex® MTB Nested ACE detection assay. The agar-dilution proportion method was used to perform drug-sensitivity testing using 7H10 Middlebrook medium. Target genes known to confer resistance to first and second-line drugs were amplified and the amplicons sequenced.

Conclusions
One hundred and ninety (5%) DNA samples tested positive for MTBC and from the resistant profiles of the 190 positive samples, we noted that multidrug-resistant TB was identified in 189 (99.5%) with 190 (100%) patients infected with MTB resistant to isoniazid and 189 (99.5%) having MTB resistant to rifampicin. Other percentages of drug resistance observed including 40% pre-XDR and 60% of XDR. This study provides valuable data on the different kinds of mutations occurring at various target loci in resistant MTBC strains isolated from samples obtained from the Eastern Cape Province and reveal a high incidence of MDR amongst the positive samples from
Eastern Cape Province, South Africa.
MYCOBACTERIAL TUBERCULOSIS CELLWALL ASSOCIATED PROTEIN PROMOTES BACILLARY SURVIVAL BY INHIBITING OXIDATIVE STRESS AND AUTOPHAGY PATHWAYS IN MACROPHAGES AND ZEBRAFISH.

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Background

*Mycobacterium tuberculosis* (*Mt*) cell wall associated glycoproteins are involved in host pathogen interactions and pathogenesis.

Objectives

In our study, we identified mannosylated glycoproteins from purified *Mt* cell wall. Our objective was to functionally elucidate the role of this protein in host immune responses and in pathogenesis.

Methods

*Mycobacterium tuberculosis* gene is cloned and expressed in *Mycobacterium smegmatis*. Orthologue of *Mt* gene in *Mycobacterium marinum* was also deleted. Invasion assay was performed in HCT-116 and intracellular survival assay, ROS, autophagic and MAPK pathways were analyzed in infected macrophages.

Conclusions

The PRT activity measurement in *Mycobacterium smegmatis* expressing recombinant gene confirmed the mass spectrometry data. Using *Mycobacterium marinum*-zebrafish and the surrogate *MsmPRT* infection models, we proved that PRT is indeed involved in mycobacterial virulence. Histological and infection assays showed that *M. marinum* mutant was strongly attenuated in adult zebrafish and also survived less in macrophages; whereas infection with wild-type and the complemented PRT *M. marinum* strains showed prominent pathological features such as severe emaciation, skin lesions, hemorrhaging, greater weight loss and more zebrafish death. This intracellular survival reveals inhibition of autophagy, reactive oxygen species and reduced activities of superoxide dismutase and catalase enzymes in infected macrophages. Infection with *MsmPRT* also activated the MAPK pathway, NF-κB and inflammatory cytokines. In summary, we show that a novel mycobacterial mannosylated PRT acts both as a virulence and an immunomodulatory factor, suggesting that it may constitute a novel target for antimycobacterial drugs.
FEMS-1779
Mycobacteria

“TB-SPRINT ULTRA” : DEVELOPMENT OF A MICROBEAD-BASED METHOD FOR DETECTION OF MUTATIONS ASSOCIATED WITH SECOND-LINE DRUGS AND ETHAMBUTOL RESISTANCE IN MYCOBACTERIUM TUBERCULOSIS ISOLATES

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Background
New cost-effective, portable and personalized molecular methods to predict Mycobacterium tuberculosis drug-resistance are needed

Objectives
To up-grade “TB-SPRINT”, a previously developed assay that targets simultaneously CRISPR polymorphism and first-line drug resistance, to detect resistance to second-line drugs.

Methods
An assay targeting gyrA codon 94, rrs nucleotides 1401, 1402 and 1484, eis promoter -10, -12 and -14, and embB codon 306 was built on 16 reference DNA. The previous 59-Plex complexity was thus increased to a 75-Plex. To validate this assay a first set of 111 DNA samples from different geographical origin (Lithuania and Colombia) were blindly analyzed. Thirty-eight samples with pyrosequencing data allowed first estimates of performances.

Conclusions
Sensitivity, specificity and overall concordance rates of the first version of TB-SPRINT ultra in comparison with pyrosequencing were: 66.7% (6/9), 96.2% (25/26) and 88.6% (31/35), respectively, for FLQ; 96.2% (25/26), 100% (10/10) and 97.2% (35/36), respectively, for KAN; 90.9% (10/11), 100% (25/25) and 97.2% (35/36), respectively.
respectively, for AMK; 100% (2/2), 100% (35/35) and 100% (37/37), respectively, for CAP; and 100% (12/12), 100% (5/5) and 100% (17/17), respectively, for EMB. Altogether, 'TB-SPRINT ultra 1.0" was able to detect most prevalent mutations associated with second-line drug resistance with very good specificities on pure and concentrated samples. We are in the process of increasing sensitivity to broaden the applicability of this test. This work shows that tuberculosis resistance to first- and second-line drugs will soon be available as an 'all-in-one tube" assay. This assay will later be tested on biological samples directly.
INFECTIONOUSNESS RATE IN PULMONARY TUBERCULOSIS PATIENTS WITH SMEAR POSITIVE OF SPUTUM IN KASHAN, IRAN

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Introduction
Identifying time predictors of sputum smear conversion in patients with pulmonary tuberculosis (TB) could be used for planning and the counseling of TB patients. This study was conducted to assess the duration in which sputum smear was turned negative and end of infectiousness of smear positive in pulmonary tuberculosis patients in Kashan.

Materials & Methods
This was a cohort study was conducted on 50 smear positive Tb patients in Kashan from the year 2012 to 2013. Sputum smear was taken from pulmonary TB patients in Moslem TB Control Center. Sputum smear was done at the end of first, 2nd, 3rd, 4th and 8th week of treatment. A questionnaire including demographic information and clinical symptoms in addition to the result of HB and ESR were collected. Data analyzed was performed by SPSS 16.0.

Results
The result of analysis indicated that 23(46%) of the AFB smear positive tuberculosis were male and 27(54%) were female. The mean age of the patients was 54.7 ± 22.7 years, while 21(42%) were under the age 49 and 29(58%) were over the age 49 years. The result also showed that 20 (40%) patients were Iranian and 30(60%) was from Afghanistan. The most frequently observed cases (90%) were from urban areas and 68% of the cases were outpatient. The highest frequency of clinical symptoms were fever 36 (72%) and dyspnea 21(42%), respectively. The negative result of sputum smear at the end of 2th week was observed in 8(16%), 3th week in 20 (40%), 4th week in 23(46%) and 8th week in 31(62%) patients, respectively. There was positive smear in 19(38%) patients. There was no significant association between the age, sex, nationality and positive result of smear at the second month of treatment. The result of ANOVA revealed that there was a significant effect of time on the infectiousness.

Conclusion
Considering the high rate of smear positive AFB at the end of second month of treatment, assessment of Dots quality and the incident of drug resistance among mycobacterium tuberculosis in Kashan is necessary and important.
MULTIPLE COLD ABSCESSES DUE TO MYCOBACTERIUM TUBERCULOSIS

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Background
The musculoskeletal system involvement of tuberculosis (TB) other than spinal TB is a rare and less-known entity. Furthermore symptoms of musculoskeletal involvement of TB infection are non-specific, relatively slow and indolent

Objectives
We report a 55-year-old patient suffering multiple cold abscesses in adjacent soft tissues; emphasising significant clinical and imaging findings

Methods

Case presentation:

A 55 year old man from Kashan-Iran presented with a 4×4 cm mass without discharge and pain in left buttok from 9 months ago treatment with cephalexin was not successful and other masses appeared in anterior of left leg, left clavicula and under right 12 th rib and right scapula. other complains were 30 kg weight loss, anorexia, anemia and occasional night fever. he was chashectic and pale and afebrile.

there was no tendeness and discharge in all masses. in sonography, collection having debri and gas extending from bone to soft tissue was reported. Mass of elbow was aspirated and acid fast bacilli was seen in smear of aspirated sample. anti TB treatment was started and after 6 month he was well and all masses were disappeared

Conclusions

TB must be considered in differential diagnosis of multiple cold abscesses even in a immunocompetent patient and treatment can be effective even in advanced stage.
MOLECULAR TYPING OF MULTIDRUG-RESISTANT AND DRUG SUSCEPTIBLE MYCOBACTERIUM TUBERCULOSIS IN ASIAN COUNTRIES
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Background

Emergence and spread of multidrug-resistant Mycobacterium tuberculosis (MDR-TB) is a serious problem. The treatment needs significantly longer period than in drug susceptible TB and the outcome tends to be disappointing.

Objectives

To understand the mechanism of the development of MDR-TB, we should have information about the genetic background of each M. tuberculosis strain. In current study, we collected TB isolates in three Asian countries, Nepal, Bangladesh and Japan, and compared the genetic characteristics between MDR-TB and drug susceptible TB groups.

Methods

TB isolates were collected in each country as follows and the spoligotype was determined, Nepal: 154 MDR-TB and 95 susceptible, Bangladesh: 182 MDR-TB and 50 susceptible, Japan: 88 MDR-TB and 29 susceptible. In MDR-TB, drug resistance determining regions of rpoB, katG and inhA were also analyzed.

Conclusions

In current study, we found some of the spoligotype lineages increased its percentage among MDR-TB, e.g. Beijing type and T1 (SIT53), which seemed to suggest their clonal expansion. Those isolates usually possessed a so-called low-fitness-cost mutation, katG G944C, and they might have additional advantageous characteristics.
Further investigation seems to be needed to elucidate the mechanism of the emergence of transmissible MDR-TB to prevent the spread of the lethal pathogen.
Background

The reemergence of tuberculosis worldwide has forced efforts to control infections caused by mycobacteria, developing tools to assist in the diagnosis of early, quickly and efficiently, with the aim of reducing morbidity related to Mycobacteria.

Objectives

The aim of this work was to detect and differentiate between an infection caused by Mycobacterium tuberculosis complex (MTBC) and atypical mycobacteria (NTM) in patients with clinically suspected pulmonary tuberculosis.

Methods

Multiplex PCR for detection of CMTB and NTM was assessed; 120 sputum samples were collect from patients with suspected TB, which were divided for PCR analysis and culture.

Conclusions
In 39.2% of the samples, mycobacterial DNA was detected; by culture, only 12.5% were positive. These results highlight the importance of the implementation of molecular techniques in the detection of mycobacterial infection with clinical correspondence. In the samples we detect MTBC (38%), *M. avium* (4.2%), *M. intracelullare* (4.2%); *M. indicus* (2.1%), and the remaining 51.5% identification was not possible since only the 16S rRNA and rpoB genes were detected, representing a high percentage of infection NTM, most of these samples were not possible to cultured. In previous years it has been detect by culturing *M. fortuitum* infection, the next step is to design specific primers for the precise identification of the NTM that represent a high percentage of infection in the population of Oaxaca. **Acknowledgements:** This work is found by Fondo Mixto CONACYT-Gobierno del Estado de Oaxaca, FOMIX 193298
RAPID DIAGNOSIS OF MYCOBACTERIUM TUBERCULOSIS WITH TRUENAT MTB: A NEAR CARE APPROACH

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Background
Control of the global Tuberculosis (TB) burden is hindered by the lack of a simple and effective diagnostic test that can be utilized in resource-limited settings. There has been considerable interest in the miniaturization of the PCR platform as this would confer advantages such as reduction in cost of instruments and tests, faster turnaround times and enhancement in the availability and accessibility of PCR tests in resource-poor geographies. With the combined advantages of affordability, simplicity in operations, diagnostic sensitivity and portability, micro-PCR devices are strong candidates for widescale use among the peripheral laboratories of India

Objectives
Objectives are as follows:
Methods
We evaluated the performance of Truenat MTB, a chip-based nucleic acid amplification test in the detection of Mycobacterium tuberculosis (MTB) in clinical sputum specimens from 226 patients with suspected pulmonary tuberculosis (TB). The test involved sputum processing using Trueprep-MAG™ (nanoparticle-based protocol run on a battery-operated device) and real-time PCR performed on the Truelab Uno™ analyzer (handheld, battery-operated thermal cycler). Specimens were also examined for presence of MTB using smear microscopy, liquid culture and an in-house nested PCR protocol. Results were assessed in comparison to a composite reference standard (CRS) consisting of smear and culture results, clinical treatment and follow-up, and radiology findings.

Conclusions
This preliminary study shows that the Truenat MTB test allows detection of TB in approximately one hour and can be utilized in near-care settings to provide quick and accurate diagnosis.
**Background**
Isoniazid (INH) is known to have a bactericidal action against actively replicating *M. tuberculosis* (*Mtb*). However, it has little effects against slowly replicating bacilli. *Mtb* reverts from nongrowing bacilli in M1 macrophages to growing bacilli in M2 macrophages and can be susceptible to INH. Therefore, the modification of host immune system is crucial for more effective elimination of persisters in non-replicating state.

**Objectives**
Therefore, the modification of host immune system is crucial for more effective elimination of persisters in non-replicating state.

**Methods**
Mouse splenocytes were cultured for 5 days and non-adherent cells were removed. The fermented soybean paste isolated Bacillus lentus (BL) was applied to cells for M1 and *Lactobacillus brevis* (LB) isolated from Kimchi, the fermented vegetable, for M2 with 48 hours of infection period. LPS for M1 and IL-4 for M2 were used as a positive control and were run in parallel with every assay. Activated cells were infected with *M. tuberculosis* H37Rv and INH was added 3 days after infection. After 4 days of incubation, number of intracellular *Mtb* was determined at 14-16 days of infection.

**Conclusions**
BL had been shown to induce M1. However, LB induced high levels of IL-10 and arginase-1 secretion which is characteristics of M2 macrophage polarization. LB was shown to induce M2 allowing the replication of mycobacteria that increased the efficacy of INH. It could be concluded that the combined use of anti-tuberculosis chemotherapy and immunotherapeutic microbiology might be more effective for eradication of persisters *Mtb* in macrophages.
Background

*Mycobacterium tuberculosis* (MTB) is able to survive under physical chemical insults, such as nitrosative and acid stress, within the macrophage host niche. By gene expression studies under stress conditions in the model organism *Mycobacterium smegmatis* mc²155 we show the down regulation of the MSMEG_3765 gene, whose deduced amino acid sequence shares 74% identity with *Mycobacterium tuberculosis* H37Rv Rv1685c gene product. Both genes are defined as members of TetR transcriptional regulators, often serving as repressors.

Objectives

Role of the MSMEG_3765 gene in stress response. Isolation and characterization of a *M.smegmatis* mutant strain, carrying a null mutation in MSMEG_3765.

Methods

Transcriptional analysis of the MSMEG_3765 locus was performed by RT-PCR; the MSMEG_3765 gene expression was evaluated by Real Time PCR after acid-nitrosative multi-stress.

The mutant strain was constructed by a two step homologous recombination process. The upstream and downstream regions of MSMEG_3765 (1-Kb flanking region) were cloned into a pNIL series vector together with marker genes obtained from the pGOAL series, leading to a suicide delivery system. The pFP1 recombinant vector was used to transform *M.smegmatis* competent cells, and the *M.smegmatis Δ3765* mutant strain was isolated.

Conclusions

The MSMEG_3765 was shown to be the last of a four genes operon. Real Time PCR analysis revealed that MSMEG_3765 is four times less expressed under acid-nitrosative multi-stress than standard growth conditions. To elucidate the role of MSMEG_3765 gene under stress conditions, stress survival and proteome analyses of mutant strain are in progress.
INCIDENCE OF PARATUBERCULOSIS IN WILD RUMINANTS IN BULGARIA - FIRST VETERINARY STUDY WITH IMPLEMENTED COMPLEX DIAGNOSTIC TECHNIQUES

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Background: Paratuberculosis is one of the most important diseases in cattle and wild animals and with great importance to the international trade market. That is why the application of combination of rapid methods will allow developing a methodology for control of paratuberculosis in Bulgaria, to conduct epidemiological analysis and to assess the risk to animals and humans.

Objectives: Application of pathomorphological, microbiological and molecular methods for detection and identification of Mycobacterium avium subsp. paratuberculosis (MAP) in tissue samples.

Methods: Study samples included 50 diagnostic materials (mesenteric lymph nodes, small intestine) from deers and mouflons in Bulgaria, suspected for paratuberculosis in the period of 2009-2013. Bacterial cultivation was carried out by using Middlebrook 7H9, Herrolds and Löwenstein media. For molecular methods was used MAP Vet kit (Sacace Biotechnologies, Italy).

Results and Conclusions: Typical for MAP pathomorphological lesions were observed in 4 mouflons and 3 deers. MAP isolates grow very well in the Middlebrook 7H9 and Herrold media but did not grow on the Löwenstein-Jensen medium even after 6 months cultivation. The presence of MAP was confirmed by PCR methods. In conclusion, we present here the first veterinary data for diagnostics of paratuberculosis in wild animals in Bulgaria. Development of this scientific field will deepen the knowledge of the properties and early detection of MAP, its significance for infectious pathology, prevention and control and will add new information to the global database for distribution of paratuberculosis in Bulgaria.
Mycobacteria

THE PEPTIDOGLYCAN REMODELLING ENZYME RPFB OF MYCOBACTERIUM TUBERCULOSIS IS TRANSCRIPTIONALLY REGULATED BY MTRA AND SIGMA FACTOR B.

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Background

*Mycobacterium tuberculosis*, the causative pathogen of tuberculosis, is a global burden to human health. With the emergence of multi drug resistance, extensive drug resistance and co-infection with HIV, tuberculosis kills 1.7 million people every year. *Mycobacterium tuberculosis* lies dormant within its host over a long time, emerging from dormancy only when conditions are favorable. Both dormancy and reactivation represent processes which are likely to be good therapeutic targets.

Objectives

The resuscitation promoting factors are hydrolytic enzymes, which are required for resuscitation of dormant mycobacteria. Resuscitation promoting factor B (RpfB), a peptidoglycan remodelling enzyme is required for reactivation of mycobacterium from chronic infection *in vivo*, underscoring the need to understand its transcriptional regulation.

Methods

Transcriptional regulation of *rpfb* has been analyzed using promoter GFP-fusions and in vitro transcription assays. Transcription start point was determined by 5’-RACE. In vitro transcription and EMSA confirmed the recognition of the *rpfb* promoter by sigma factor, SigB and transcription factor, MtrA respectively. Streptavidin pull down assay with a biotinylated DNA fragment and chromatin immunoprecipitation confirmed association of MtrA with the *rpfb* promoter region. Binding site of MtrA on the *rpfb* promoter was identified by DNasel footprinting assay.
Conclusions

SigB and MtrA are the two regulators of \textit{rpfB}. MtrAB coordinates cell wall homeostasis. A likely role in cell wall metabolism is strengthened by this report that MtrA regulates expression of the peptidoglycan hydrolase, RpfB. The insights into regulation of \textit{rpfB} are therefore relevant in our quest to uncover new bacterial targets for drug development.
ROLE OF THE N-TERMINUS OF THE MYCOBACTERIAL MEMBRANE PROTEIN MCTB IN THE LOCALIZATION, STRUCTURE AND RESISTANCE TO COPPER.

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³Department of Chemistry, Tsinghua University, Beijing, China
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Background
We previously described the protein Rv1698 of Mycobacterium tuberculosis as involved in an unknown mechanism of resistance to toxic copper ions: mutants of M. tuberculosis and M. smegmatis lacking functional expression of the gene accumulated copper while wild-type cells did not. Rv1698 is a cell envelope, channel-forming protein and we hypothesized it could be part of a copper efflux machinery. We named it MctB (Mycobacterial copper transport protein B).

Objectives
To gain insight on the contribution of MctB to copper resistance, we aimed to characterize its structure.

Methods
We solved the structure of recombinant MctB lacking its putative signal peptide by X-ray crystallography. We determined the composition of the mature protein by mass spectrometry analysis. By expressing translational fusions of mctB to various reporter genes within phenotypic assays, we investigated the role of the N-terminal hydrophobic α-helix on the localization and function of the protein. We analyzed the oligomeric state of MctB by size-exclusion chromatography and electron microscopy (EM).

Conclusions
In contrast with the established biophysical properties, the crystal structure of recombinant MctB did not show characteristics of a classical channel protein, namely the presence of a channel or contiguous hydrophobic surfaces. We showed that the N-terminus – thought to be a classical signal peptide – (i) is still present in the mature form of MctB, (ii) is required for the translocation and activity of MctB outside the cytosol and (iii) plays a critical role in its oligomerization. Finally, we propose an oligomeric model of MctB based on EM data that helps explain MctB properties.
MOLECULAR CHARACTERIZATION OF MYCOBACTERIUM ORYGIS ISOLATES FROM WILD ANIMALS OF NEPAL

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Background
Mycobacterium tuberculosis complex (MTC) species cause tuberculosis (TB) in humans and animals. Recently, we isolated acid fast bacilli from a spotted deer (Axis axis) and a blue bull (Boselaphus tragocamelus) in Nepal. They showed atypical colony characteristics.

Objectives
Genotypic characterization of bacterial isolates helps to understand the nature of bacteria. In this study, we aimed to characterize the bacilli isolated from the spotted deer and the blue bull in detail.

Methods
TB suspected lesions from the lung and lymph node in the deer and the lung in the blue bull were collected during postmortem and cultured on Löwenstein-Jensen media. Spoligotyping, multi-locus variable number of tandem repeats analysis (MLVA), region of difference analysis (cpn32, RD9 and RD12) and SNP typing were performed to identify species and to get molecular epidemiological insights.

Conclusions
Isolates from both animals showed a spoligotype SIT587 in SITVIT WEB database as of January 2015. RD analysis results (cpn32: +, RD9 - and RD12-) and SNP data in gyrB, mmpL6, Tbd1, PPE55 and Rv2042c identified these isolates to be M. orygis. In addition, MLVA of these isolates showed a similar pattern with many other reported M. orygis isolates. M. orygis have been isolated from many animals and humans of South Asian countries. Our finding may indicate M. orygis to be endemic in the subcontinent. Additionally, the identification of a new member of MTC in Nepal may suggest the complexity of tuberculosis in the country. Further investigation with increased sample size and various host animals including human will help to understand the ecology and epidemiology of M. orygis.
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Mycobacteria

MIXED INFECTION IN TUBERCULOSIS: THE CRUCIAL PHENOMENA IN OUTCOME OF TREATMENT
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Background

*Mycobacterium tuberculosis* (MTB) is among the most successful human pathogenic bacteria worldwide and is responsible for extensive morbidity and mortality, with approximately 2 million deaths each year thought to be due to primary infection, reactivation of latent infection, or re-infection with a new strain. Molecular fingerprinting has differentiated the MTB strains. Most of these studies have shown that each patient is infected with one strain. However, by using more sensitive PCR based technique such as MIRU-VNTR, it is possible to determine co infection with multiple strains of MTB.

Objectives

The aim of the current study was to determine the prevalence of mixed infection and clonal complexity in Iranian patients by MIRU-VNTR technique. Additionally, we aimed to determine whether mixed infection results in differences in clinical manifestation or response to treatment.

Methods

The PCR was designed to amplify a standard set of 24 MIRU-VNTR loci from genomic DNA extracted using primers described by Supply et al. PCR products were electrophoretically separated on 1.8% agarose gels. In addition, the presence of double alleles was confirmed by two independent rounds of PCRs

Conclusions

The MIRU-VNTR method was able to identify mixed infection in at least 25% (17/67) of the cases. Mixed infections occur at high prevalence among Iranian TB patients. The presence of multiple strain infections had high impact on the clinical manifestation and response to anti tuberculosis treatment for individual patients.
PRESENCE OF MUTATIONS IN GENES ASSOCIATED WITH ANTIBIOTIC RESISTANCE IN MYCOBACTERIUM TUBERCULOSIS STRAINS FROM MICHOACÁN, MEXICO

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Background

The six-month treatment against tuberculosis involves rifampin, isoniazid, pyrazinamide, ethambutol and streptomycin. Associations between mutations in specific genes (rpoB to rifampicin; katG and promoter mabA to isoniazid; pncA to pyrazinamide; embB to ethambutol; rrs and rpsL to streptomycin) and resistance to antibiotics have been reported. The frequency of these mutations varies in each country and region within a country, so in order to design diagnostics tools, the relationship between mutations and first line resistance patterns in each country must be known.

Objectives

Identify mutations associated to antibiotic resistance in strains of M. tuberculosis from patients in the state of Michoacán, Mexico.

Methods

Thirty-three strains of Mycobacterium tuberculosis were analyzed. The susceptibility patterns were determined using BACTEC-MGIT. Mutations were detected by PCR amplification, sequencing, and comparison with the TB Drug Resistance Mutation Database.

Conclusions

21.2% of the studied strains were resistant to rifampin. The genes katG and rpoB showed the highest mutation percentage (30.3% and 27.2%, respectively). 54.5% of the strains presented at least one mutation in any of the analyzed genes. Our analysis of resistant strains found both new and previously reported mutations such
as the katG Ser315Thr. The four analyzed strains isolated from patients with meningeal tuberculosis showed extensive drug susceptibility of first line antibiotics. Only two of the latter strains had mutations in the katG gene: strain 049 with synonymous Ala411Ala mutation having no influence on antibiotics sensitivity, and 040 with Arg463Leu mutation reported as producing resistance in some cases. 62.5% of the strains resistant to drugs showed some resistance mutation.
VARIATIONS IN THE SEQUENCES OF VIRULENCE GENES HARBORED BY MYCOBACTERIUM TUBERCULOSIS STRAINS FROM MICHOACAN

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Background
The Mycobacterium tuberculosis genome contains two large families of genes called pe and ppe, some of which encode proteins involve in virulence and the evasion of the immune response. These genes are highly polymorphic and these polymorphisms have been associated with the pathogenicity of strains that are presented.

Objectives
To find relationships between polymorphisms in pe/ppe gene families and certain genotypes.

Methods
59 M. tuberculosis strains obtained of tuberculosis patients from Michoacán, México, were analyzed. The species, lineage, family and genotype of the strains were previously determined. The pe_pgrs18 and pe_pgrs26 genes were amplified and sequenced. Polymorphisms were found by alignment with ClustalX and MEGA6 software against homologous sequences retrieved from GenBank.

Conclusions
The pe_pgrs18 gene sequences from strains belonging to T, LAM and some no previously reported (NPR) families from the Euro American lineage, EAI strains from Indo Oceanic lineage and Beijing strains from East Asia lineage, harbor SNPs in positions 54-266. Sequences from the same gene, but from Harlem, X and different NPR strains from Euro American lineage, NPR strains from Indo Oceanic lineage and Ghana strains from West Africa lineage, showed SNPs and insertions in positions 295-510. The pe_pgrs26 gene harbor SNPs, insertions and deletions; X and T strains shared the same SNPs, while EAI strains showed their own set of shared SNPs.

Some detected polymorphisms are consistent with previous reports, while others are not. The gene pe_pgrs18 was highly polymorphic. There is a relationship between genotypes and certain polymorphisms in pe and ppe families.
GENE EXPRESSION PROFILE OF BIOMARKER CANDIDATES IN MYCOBACTERIUM AVIUM SUBSP. PARATUBERCULOSIS INFECTED CATTLE

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Background

Johne’s disease (JD) is a chronic disease of ruminants caused by *Mycobacterium avium* subsp. *paratuberculosis* (MAP). JD has very long incubation periods, resulting in difficulties in diagnosing the disease, especially regarding the fecal shedders of MAP without any clinical signs. Current MAP diagnosis techniques lack sensitivity in the detection of early stage of MAP infection. Therefore, attempts were made to findout biomarker to detect early responses to MAP infection.

Objectives

To develop enhanced diagnosis of early stage of MAP infection, biomarkers were evaluated by analysis of immune-related genes in MAP-infected cattle.

Methods

Total RNA was extracted from whole blood of 32 cattle characterized by ELISA and IS900/IS-MAP PCR. Expression of 24 genes was analyzed by real-time PCR to validate the biomarker candidates. The expression level was determined by the $2^{-\Delta\Delta Ct}$ method using β-actin, as a reference. The expression level was compared to the control group to determine the expression-fold change of each gene.

Conclusions

Based on the present study, four biomarker candidates showing significant changes were proposed as potential biomarkers of JD. CCL4 and CXCR3 expression were significantly decreased in group 1. On the contrary, CXCL9 expression was significantly decreased in group 2. KLRB1 expression was significantly increased in group 2. These biomarkers might be used for development of enhanced diagnostic tool of early stage of MAP infection in further study. This work was carried out by the grant from RDA, Korea(Project No. PJ008970012015)
EVALUATION OF MYCOBACTERIUM AVIUM SUBSP. PARATUBERCULOSIS ANTIGENS TO DETECT SUBCLINICAL STAGE OF INFECTION IN CATTLE
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Background

Johne’s disease, caused by Mycobacterium avium subsp. paratuberculosis (MAP) is one of the most widespread and economically important disease in cattle. Current diagnostic methods are based on detection of MAP antibodies in milk or serum samples or isolation of causative agent. However, these diagnostic techniques are often not applicable in subclinical infection because of relatively low sensitivity. Therefore, finding new antigen candidates highly react with host immune system has been required.

Objectives

To detect subclinical stage of infection effectively, several antigen candidates were selected from previous reports. Selected antigen candidates were analyzed about their efficiency.

Methods

Complete gene sequences of antigen candidates were cloned and expressed by cold shock expression system with pCold™ I vectors and One shot® chemically competent E. coli Top10 cells. Recombinant proteins were tagged by 6-histidine and purified by His SpinTrap™. Serological samples were prepared from cows with different stages of the disease divided into four groups according to stage of infection. Reactivity with these four groups of sera and each antigen was evaluated by western blot.

Conclusions

Total of 9 antigen candidates were selected (MAP0862, MAP3817c, MAP2077c, MAP0860c, MAP3954, MAP3155c, MAP1204, MAP1087, MAP2963c) which have MAP specific and/or have high immune responsibility to infected animals. Using four groups of serological samples, it could be evaluated whether the antigen possibly
detect subclinical infection or not. Possible antigen cocktail will be used for
development of sensitive ELISA. This work was carried out with Project No.
PJ008970012015.
COMPLEX INFECTION AS A POTENTIAL MECHANISM FOR HETERO-
RESISTANCE AND RIFAMPIN RESISTANCE IN MYCOBACTERIUM TUBERCULOSIS

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Background
Rifampin is a key component of the first-line therapy against Mycobacterium tuberculosis. Rifampin resistance was reported to be caused by mutation in rpoB gene, while 5% of the resistant strains showed no mutation. Hetero-resistance is considered as the preliminary stage to the full resistance in M. tuberculosis.

Objectives
This study investigated the possible mechanisms conferring hetero-resistance and rifampin resistance without rpoB gene mutations.

Methods
Two hundred and twelve clinical isolates were subjected to rpoB gene sequencing to examine the occurrences of rpoB mutations and hetero-resistance. The standard 24-locus variable-number tandem repeat (VNTR) method was used to identify complex infection, which consisted of mixed infection (the presence of strains with different VNTR patterns at ≥ 2 loci in one sample) and clonal heterogeneity (different VNTR patterns at a single locus).

Conclusions
We found that 3.8% of isolates (n = 8) had rifampin resistance of no mutation in rpoB gene, in three among which dual peaks from the sequencing results were revealed and therefore identified as rifampin hetero-resistant. The main peak was wild type and the secondary one was mutated (Cys-526, Leu-531 and Tyr-526). All three hetero-resistant isolates were caused by complex infection (18, 8.5%), which included mixed infection (7, 3.3%) and clonal heterogeneity (11, 5.2%). Rifampin resistance with no mutation in rpoB gene was significantly associated with complex infection ($\chi^2$, 31.21, $P < 0.01$), especially mixed infection ($\chi^2$, 56.78, $P < 0.01$). Accordingly, it is necessary to examine complex infection, a possible mechanism for hetero-resistance and rifampin resistance without rpoB gene mutations.
Background

Herbal drugs have been used since ancient times as remedies for the treatment of a variety of ailments. *Premna* is a member of family Verbenaceae. Some of *Premna* species are known to exhibit cytotoxic, antimicrobial, antioxidant, antipyretic, hypoglycemic and cardiotonic activities.

Objectives

In this study *P. resinosa* methanolic extract as well as its constituents were evaluated for antimicrobial and cytotoxic activity.

Methods

*Premna resinosa* powdered plant was extracted with methanol and constituents were isolated by different chromatographic techniques and identified by spectroscopic methods. The extracts and compounds were tested for antimicrobial *in vitro* against *Staphylococcus aureus*, *Bacillus subtilis*, *Acinetobacter baumannii*, *E. coli*, *Salmonella typhimurium*, *Shigella flexneri* and *Enterococcus faecalis*. In addition, they were tested against human medulloblastoma (Daoy), human hepatocellular carcinoma (HepG2) and human melanoma (SK-Mel-28).

Conclusions

Seven constituents were identified from *P. resinosa*. They are Quercetin (1), 3-Methoxyquercetin (2), Kaempferol (3), 3-Methoxykaempferol (4), Myricetin 3,7,3'-trimethyl ether (5), Lupeol (6), Stigmasterol (7). The dichloromethane extract exhibited the highest antimicrobial effect, meanwhile, Noticeable cytotoxic activity was shown by the crude methanolic extract against Daoy cell line. The antimicrobial activity of the isolated compounds showed higher sensitivity of Gram-positive bacterial strains than the Gram-negative.
NEW INSIGHTS IN THE BUTYRATE PHOTOASSIMILATION IN RHODOSPIRILLUM RUBRUM S1H
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Background

Purple non-sulfur bacteria display a great metabolic versatility and are able to assimilate volatile fatty acids (VFAs) through photoheterotrophic metabolism. One of these bacteria, Rhodospirillum rubrum S1H, was subsequently selected by the European Space Agency for its ability to metabolize VFAs and to remove them in its bioregenerative life support system.

Objectives

Here we aim to offer a better view of the still misunderstood photoheterotrophic metabolism of butyrate.

Methods

A global understanding of butyrate photoassimilation in purple non-sulfur bacteria was obtained through a multidisciplinary approach including quantitative SWATH-MS proteomic analysis, activity assays, bacterial growth analysis and VFAs consumption monitoring.

Conclusions

Butyrate is partially converted into acetyl-coA. Because R. rubrum is an isocitrate lyase lacking organism, an alternative anaplerotic pathway to glyoxylate cycle should be used to replenish the TCA cycle with intermediates for biosynthesis. We suggest here that butyrate could be converted into propionyl-CoA and succinate through ethylmalonyl-CoA pathway and valine degradation pathway.

We also observed that butyrate assimilation occurs only if carbonate is supplied. In accordance with the observed upregulation of RuBisCO, CO2 fixation is most probably used as redox balancing reaction necessary to sustain butyrate assimilation.

Finally, we showed an inhibition of the butyrate assimilation by acetate when they are provided together at an equivalent net carbon concentration. This effect could be due
to the observed downregulation of key enzymes of the butyrate assimilation pathway when acetate is used as carbon source.

Altogether our data allows better understanding of metabolic pathway involved in butyrate photoassimilation in R. rubrum.
THE UNIQUE SHEATHS OF METHANOGENIC ARCHAEA ARE FUNCTIONAL AMYLOID STRUCTURES
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Background
Amyloids are highly organized protein or peptide polymers characterized by a cross-β-sheet structure. These protein structures were for many years solely considered an abomination associated with incorrect protein folding, leading to devastating neurodegenerative diseases, such as Alzheimer's disease. However, within the last decade it became clear that the amyloid protein fold has many beneficial roles in nature. In bacteria they are used as adhesive fimbriae and for the coating of spores. Fungi use them to modulate the hydrophobicity of aerial hyphae and spores and in the form of prions as a means of phenotype regulation. In mammals, amyloids are used as a scaffold for the skin pigmentation and as a storage form for hormones. However, functional amyloids have never been shown within the Archaea domain of life.

Objectives
Members of the archaeal genera *Methanospirillum* and *Methanosaeta* grow as filamentous chains of rod-shaped cells within tubular paracrystalline sheaths. These sheaths share many properties with the functional amyloids. They are proteinaceous and display high stability toward chemical and thermal denaturation and display a cross-β-sheet structure. Accordingly, we hypothesized that the archaeal sheaths could represent a novel type of functional amyloid.

Methods
We employed amyloid-specific conformation antibodies and biophysical techniques to confirm the amyloid nature of the sheaths.

Conclusions
Our work represents the first report of a functional amyloid from the archaeal domain of life. The amyloid nature explains the extreme resistance of the sheath, the elastic properties that allow penetration of diffusible substrates, and how the sheaths are able to split and elongate outside the cell.
EXTRACELLULAR STRUCTURES OF PELAGIC FORMOSA
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Background

*Flavobacteria* contribute to the degradation of complex organic matter, in particular proteins and polysaccharides. During the decay of a spring phytoplankton bloom in the German Bight of the North Sea near Helgoland in 2009 a successive occurrence of different *Flavobacteria* clades represented up to 60% of the bacterial community (Teeling, H. *et al.*, 2012). Strains of the genus *Formosa* in the marine clade of *Flavobacteriaceae* were recently isolated from this site (Hahnke, R.L. *et al.*, 2014). Large extracellular structures were detected (Hahnke, R.L. *et al.*, 2014), however their function is still not understood.

Objectives

We started to study such appendages to gain new insights into their biological role within the pelagic bacterioplankton.

Methods

Light microscopy, transmission electron microscopy (TEM) and scanning electron microscopy (SEM) showed at cells of *Formosa* clade A segmented chain-like appendages with a width between 40 to 100 nm. A first membrane preparation by cells disintegrated by pressure change and by differential centrifugation revealed TonB-dependent receptors, an OmpA protein, one SusD-like protein and two hypothetical proteins in the outer membrane enriched fraction. Similarities in the protein pattern were observed for the other pelagic strain, *Formosa* clade B. In contrast, *Formosa agariphila* isolated from green algae revealed a different protein pattern.

Conclusions

In this contribution, we will report the outer membrane/appendage proteome as prepared by different protocols. In addition, the appearance of appendages as
response to the physiological situation of the cells may provide first insight into the ecological importance of these appendages.
INVESTIGATION OF THE SIGNAL LEADING TO SURFACE LOCALIZATION OF LIPOPROTEINS IN CAPNOCYTOPHAGA CANIMORSUS

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Background

The gram negative Bacteroidetes are prominent members of the human intestinal flora, where they are responsible for the breakdown of all major classes of host and dietary glycans. This ability is encoded by so-called Polysaccharide Utilization Loci (PUL), of which the starch utilization system (Sus) of B. thetaiotaomicron is the archetype. The Sus is a cell envelope-associated multiprotein complex characterized by the coordinated action of several proteins and lipoproteins involved in starch binding, degradation and internalization into the periplasm. Bacteroidetes genome sequencing projects showed that PUL are a hallmark of the phylum. PUL-encoded lipoproteins are surface exposed and little is known on how these proteins are recognized and targeted to the cell envelope.

Objectives

We aimed at understanding how the bacterial cell distinguishes between periplasmic and surface lipoproteins.

Methods

We used as model organism Capnocytophaga canimorsus, an oral commensal of dogs and cats, which can cause severe human infections. An in silico approach allowed the identification of a putative signal region at the N-terminus of the lipoproteins and we therefore performed a mutagenic screen on this region on the PUL-encoded surface mucinase Ccan_17430. The mucinase surface localization of the N-terminal mutants was detected by monitoring mucin degradation.

Conclusions

We found a region needed for protein stability, preventing proteolytic cleavage and/or miss-localization of the protein. Further work is needed to characterize the signal and to determine whether the protein degradation observed for several mutants is due to protein instability or is a consequence of miss-localization.
THE STACK: A NEW BACTERIAL STRUCTURE ANALYZED IN THE ANTARCTIC BACTERIUM PSEUDOMONAS DECEPTIONENSIS M1T BY TEM CRYO-TEM AND TOMOGRAPHY

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Background

Improvements in TEM and Cryo-TEM techniques and the use of Cryo-electron tomography (CET) of plunge-frozen whole bacteria and vitreous sections (CETOVIS), allow visualization of biological samples close to their native state, enabling us to refine our knowledge of already known bacterial structures and to discover new ones.

Objectives

Application of these techniques to the Antarctic cold-adapted bacterium Pseudomonas deceptionensis M1T to demonstrated the existence of a previously undescribed cytoplasmic structure.

Methods

We used cryo-immobilization of specimens by rapid cooling followed by freeze substitution (FS) and sectioning, freeze fracture (FF) and observation of replica, cryoelectron microscopy of vitreous sections (CEMOVIS) and CET of plunge-frozen whole bacteria and vitreous sections to analyze this new structure.

Conclusions

P. deceptionensis M1T and other bacteria contain a new structure we termed "Stack". Each "Stack" consisted of a pile of oval disc-like subunits, each disc being surrounded by a lipid bilayer membrane. They are found at different locations within the cell cytoplasm, in variable number, separately or grouped together, very close to the plasma membrane (PM) and oriented at different angles (from 35º to 90º) to the PM. Stacks did not appear to be invaginations of the PM. Regardless of their position, stacks were mostly observed very close to DNA fibers. This new structure deserves further study to refine its configuration and to establish whether its function could be related to chromosome dynamics.
STRUCTURAL AND FUNCTIONAL INSIGHT IN THE HELICOBACTER PYLORI BLOOD GROUP ANTIGEN BINDING ADHESIN BABA

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Background

*Helicobacter pylori* is one of the most successful human pathogens, chronically infecting up to half the world population and thereby causing chronic gastritis, duodenal ulcers and even gastric cancer. The blood group antigen binding adhesin (BabA) of *H. pylori* mediates attachment to stomach glycoreceptors containing ABO and Lewis b blood group antigens.

Objectives

Our aim was to gain structural insight in the interaction between BabA and ABO/Lewis b blood group antigens.

Methods

BabA belongs to a paralogous family of *Helicobacter* outer membrane proteins (Hops) and we demonstrate BabA to feature a novel topology with an OmpA-like 8-stranded non-contiguous β-barrel that is disrupted by an extracellular domain that mediates the adhesive properties to BabA. The crystal and small angle X-ray scattering structure for the BabA extracellular domain reveals a two-domain architecture composed of an α-helical head domain and a coiled-coil stem domain making the connection to the β-barrel domain. The X-ray structure of the BabA ectodomain bound to its natural glycan receptor Lewis b is presented. Receptor binding is located in a small subdomain grafted into the conserved head domain and critically depends on a disulphide-bound loop in the binding site. The ligand-bound structures of two BabA isomorphs provide a molecular rationalization of the observed blood group antigen polymorphisms in *H. pylori* clinical isolates.

Conclusions

Our results represent the first structural insight in how the important human pathogen *H. pylori* recognizes and interacts with its human host.
FUNCTIONAL STUDIES ON THE HMG-COA REDUCTASE GENES OF MUCOR CIRCINELLOIDES

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Background

Terpenoids have several biological functions. Among others, members of this family of chemical compounds are cell membrane components (e.g. ergosterol in fungi), pigments (e.g. carotenoids) or functional groups of certain proteins (e.g. RAS proteins). HMG-CoA reductase is a membrane anchored protein catalysing the central step of the acetate-mevalonate pathway of the terpenoid biosynthesis. Genome of the carotenoid producing zygomycete fungus Mucor circinelloides contains three HMG-CoA reductase genes (i.e. hmgR1, hmgR2 and hmgR3). Earlier, we analysed the expression of these genes under different growth conditions by qPCR studies (1).

Objectives

In this study, function of these genes and subcellular localization of the encoded proteins were analysed. Functional studies were carried out by overexpression and silencing the three genes.

Methods

For silencing, the antisense RNA technique was used. Carotene and ergosterol content and morphology of the overexpressed and silenced transformants were examined. To analyse the subcellular localization of the enzymes, they were fused with the green fluorescent protein and different dyes were used to analyse their co-localization to different compartments of the cell.

Conclusions

Our results suggest that HMGR2 and HMGR3 co-localize to the endoplasmic reticulum. HMGR2 plays role in the general isoprenoid biosynthesis, while HMGR3 is especially active under anaerobic growth condition and participates in the morphogenesis and the germination of the spores.

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References

Background

Outer-inner membrane vesicles (O-IMVs) were recently described as a new type of membrane vesicle (MV) secreted by the Antarctic bacterium *Shewanella vesiculosa* M7T. Their formation is characterized by the protrusion of both outer and plasma membranes, which pulls cytoplasmic components into the vesicles [1].

Objectives

The aim of the current study was to determine whether these new O-IMVs are also secreted by pathogenic Gram-negative bacteria whose MVs have been repeatedly reported to contain cytoplasmic components and DNA.

Methods

For this purpose, three pathogenic bacteria: *Pseudomonas aeruginosa* PAO1, *Acinetobacter baumannii* AB41 and *Neisseria gonorrhoeae* DSM15130 and their derived MVs were analyzed by Transmission Electron Microscopy (TEM) and Cryo-TEM. The presence of the cytoplasmic constituents such as DNA and ATP was also evaluated. The encapsulation of DNA inside the O-IMVs was confirmed by gold DNA immunolabeling. A proteomic analysis of *N. gonorrhoeae*-derived membrane vesicles was performed to identify proteins from the cytoplasm and plasma membrane.

Conclusions

TEM and Cryo-TEM techniques enabled us to demonstrate that Gram-negative pathogenic bacteria produce the recently described new type of vesicles, O-IMVs, together with conventional outer membrane vesicles (OMVs). The existence of O-IMV extends the hitherto uniform definition of MVs in Gram-negative bacteria and explains the presence of components in MVs such as DNA, cytoplasmic and inner membrane proteins, as well as ATP [2].

References

2. **C. Pérez-Cruz et al.** 2014. Outer-Inner Membrane Vesicles Naturally Secreted by Gram-negative Pathogenic Bacteria. Plos one (Accepted).
Background

The ability to sense and respond to external stimuli is a key feature of all living organisms. Motility is essential to achieve this goal. Archaea are able to swim using a rotating surface structure, the archaellum. It functions analogous to the bacterial flagellum, but its components are unrelated. Instead the archaellum subunits are homologous to components of type IV pili. In bacteria, transfer of external stimuli to the flagellum is mediated by the chemotaxis system. The response regulator CheY plays herein a central role. Euryarchaea display photoaxis, aerotaxis and chemotaxis. Euryarchaea have acquired chemotaxis components from bacteria via horizontal gene transfer. This raises the question how the bacterial chemotaxis machinery has been evolutionary adapted in order to interact with subunits of the archaeal motility structure.

Objectives

We aim to elucidate the functional and structural details interaction of the archaeal CheY with the archaellum motor complex to alter the direction of swimming.

Methods

We have applied crystallography to reveal the structure of the euryarchaeal CheY in 1.6 Å resolution. In addition we have determined its phosphorylation state, as well as that of CheA. CheA is an upstream chemotaxis signaling component, which in bacteria is responsible for phosphorylation of CheY.

Conclusions

The crystal structure of the archaeal CheY shows high similarity to the bacterial CheY. However, the interface of flagellum interaction is very basic in the bacterial CheY, while this same site shows a negative charge in the archaeal CheY. This suggests that the archaeal CheY is structurally adapted for interaction with the archaellum.
IDENTIFICATION OF NOVEL BINDING PARTNERS OF CAMPYLOBACTER JEJUNI FLAGELLINS

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Background
Flagella are propeller-like structures embedded in the bacterial envelope, which allow bacteria to swim in directions dictated by diverse sensory signals. In many pathogenic bacterial species, flagellar motility is required to reach the ultimate infection niche and to establish infection. The human pathogen *Campylobacter jejuni* is a highly motile organism that possess a flagellum on each pole. As many as 20,000 subunits of flagellin are assembled into a single filament via highly regulated process. Most flagellar proteins are bound to specific chaperones before export by the flagella assembly machinery. *C.jejuni* encodes for three different flagellins, however flagellar chaperones have not been characterized yet.

Objectives
Flagellar chaperones are well-studied in Gammaproteobacteria but not Epsilonproteobacteria including the human pathogens: *Campylobacter jejuni* and *Helicobacter pylori*. The goal of this study is to characterize the flagellar chaperones in *C.jejuni*, the main cause of bacterial food-borne infections.

Methods
Potential flagellar chaperones of *C.jejuni* were cloned and expressed in *E. coli*. Interaction of the recombinant putative chaperones with purified *C.jejuni* FlaA flagellin was tested via *in vitro* and *in vivo* pull-down experiments. The effect of targeted mutagenesis of the chaperones on flagella production was tested by a motility soft agar assay and (electron)microscopy methods. Western-blotting and qRT-PCR were used to detect differences in the FlaA, FlaB or FlaC flagellin expression levels.

Conclusions
Targeted mutagenesis of putative flagellar chaperones influenced bacterial motility, flagellin levels and flagella length. The observed phenotype can be explained by direct interaction of chaperones with FlaA flagellin as demonstrated by pull-down experiments.
LEA PROTEINS ARE INVOLVED ON DESICCATION RESISTANCE AND OTHER ABIOTIC STRESSES IN AZOTOBACTER VINELANDII

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Background

In plants, the expression of Late Embryogenesis Abundant Proteins (LEA) correlates closely with the acquisition of tolerance against drought, frezzing and salinity stresses. In *A. vinelandii* we found two genes with homology sharing identity to putative LEA proteins from Arabidopsis thaliana, the sequence of the genes contains a part of the LEA eukaryotic motif, but differs from the distinctive motive found in other bacteria, so this could be an example for the possibility of horizontal gene transfer between domains of life.

Objectives

The aim of this work is to analyze if LEA proteins are involved on the resistance to desiccation, heat, frezzing and osmotic stress in both cysts and vegetative cells of *A. vinelandii*. 

Fig. 1 Tolerance to osmotic stress caused by a) NaCl and b) Sorbitol in *A. vinelandii* cells
Fig. 4 Tolerance to high temperatures in a) vegetative cells, b) cysts of *A. vinelandii*.

Fig. 5 Tolerance to freezing in *A. vinelandii* vegetative cells.

% Desiccation Resistant Cysts
<table>
<thead>
<tr>
<th>Strain</th>
<th>WT</th>
<th>LEA1</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>9.52%</td>
<td>1.40%</td>
</tr>
<tr>
<td>2</td>
<td>16.23%</td>
<td>0.0%</td>
</tr>
<tr>
<td>3</td>
<td>7.25%</td>
<td>2.30%</td>
</tr>
<tr>
<td>Mean</td>
<td>11.00%</td>
<td>1.23%</td>
</tr>
</tbody>
</table>

**Methods**

We constructed mutants for these genes, LEA1 and LEA2 and a double mutant LEA12. Cysts and vegetative cells were treated to desiccation for 3 months; osmotic stress was caused by different concentrations of NaCl and Sorbitol. We tested the tolerance of these mutants to high temperatures (60°C for 15 minutes) and freezing (-20°C for three months) for both cysts and vegetative cells.

**Conclusions**

LEA proteins are necessary to confer tolerance to desiccation on cyst cells after 3 months treatment and are necessary to survive at high osmotic agents concentrations in vegetative cells too. LEA proteins can also protect cyst and vegetative cells against high and low temperatures since LEA mutants were unable to grow at 50°C or -20°C.
Background
Curli are proteinacious fibers that constitute the major component of the extracellular matrix in pellicle biofilms in many Gram-negative bacteria.

Objectives
Numerous fundamental questions about the curli biogenesis pathway remain unsolved. Curli formation requires a dedicated protein secretion machinery comprised of the outer membrane lipoprotein CsgG responsible for the transport of curli subunits across the outer membrane and two soluble accessory proteins. Here we propose a novel diffusion-based and entropy-driven transport mechanism that is as a working model for the CsgG translocator.

Methods
Recently the X-ray structure of this protein was solved within our group, revealing a nonameric complex embedded in the outer membrane by a 36-stranded β-barrel. Based on this structure and preliminary single-channel results we observe that the CsgG pore on its own is an ungated, non-selective protein secretion channel. As earlier described in literature we confirm that the addition of certain amounts of the CsgE chaperon protein blocks the CsgG pore.

Conclusions
Together these findings support the working model we propose. Future structural and single-channel analysis give even more insight in the process and will probe how subunit secretion is coupled with fiber assembly.
GAC IS ESSENTIAL FOR GROUP A STREPTOCOCCUS AND DEFINES A NEW CLASS OF MONOMERIC DTDP-4-DEHYDRORHAMNOSE REDUCTASES

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Background

Group A Streptococcus (GAS) is a preeminent Gram-positive human pathogen. GAS expresses a characteristic cell wall polysaccharide, known as Group A Carbohydrate (GAC), consisting of rhamnose and N-acetylglucosamine (GlcNAc), which is critical for full virulence. The final step of the four-step dTDP-L-rhamnose biosynthesis process is catalyzed by dTDP-4-dehydrorhamnose reductase (RmlD) in an NAD(P)H-dependent manner. Previous structure elucidation of RmlD from Salmonella enterica demonstrated that Mg²⁺-dependent dimerization is essential for enzymatic activity. The first gene in the GAC gene cluster, gacA, is annotated as an RmlD enzyme.

Objectives

We aimed to identify the structure, function and biological importance of the gacA gene product.

Methods

We generated recombinant GacA protein for biochemical and structural analysis. Also, a saturated transposon library of GAS was screened by whole genome sequencing (Tn-seq) and subjected to Bayesian analysis to identify essential genes. Finally, gacA was expressed in a S. mutans rmlD knockout to analyse whether this would restore its attenuated phenotype.

Conclusions

GacA catalyses the biosynthesis of dTDP-rhamnose in a novel Mg²⁺-independent monomeric manner. The 1.1 Å high-resolution structure of GacA confirms conservation of key active site residues, but GacA lacks the dimerization motif characteristic for RmlD in Gram-negative bacteria. GacA may therefore represent a new class of monomeric RmlD enzymes. We identified gacA as an essential gene and confirmed enzymatic function by restoring the growth and morphology of an S. mutans rmlD mutant. Our report provides a framework for future screenings to identify novel inhibitors that interfere with dTDP-rhamnose biosynthesis in human pathogens.
CYDIV, A CRITICAL PROTEIN IN THE CELL DIVISION IN ANABAENA SP. PCC7120.

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Background

Bacterial cell division is a fundamental and complex process that requires coordinated mechanisms for elongation and septal formation in the middle of the cell. Most researches in cyanobacterial division have been focused on identifying proteins that interact specifically with the FtsZ protein, also revealing that the division machinery presents differences in composition and regulation compared to the division of gram-negative and gram-positive model microorganism. Studies carried out in our laboratory have determined a gene common to all filamentous cyanobacteria (cyDiv), which encodes a protein with one transmembrane and coiled-coil domains. The phenotype of the mutant strain in this gene presents serious defects in cell division.

Objectives

To determine the participation of CyDiv protein in the cell division process of the filamentous cyanobacteria Anabaena sp. PCC7120.

Methods

We analyzed CyDiv localization by immunofluorescence generating specific polyclonal antibodies against the protein from a synthetic peptide. Furthermore, using bioinformatic analyses, we predicted the possible role of this protein.

Conclusions

We showed that CyDiv is localized in the cell poles and in the middle of the cell depending on the division phase, co-localizing with the Z-ring. These results, together with previous phontypical observations of the cyDiv mutant, suggest that CyDiv is essential for cell division in filamentous cyanobacteria. In silico analyses determined that this protein has a similar topology to the cellular division proteins FtsB and DivIC, from E.coli and B. subtilis respectively.

Acknowledgments: Fondecyt Grant #1131037
Background

The Gram-negative bacterium Acinetobacter sp. Tol 5 shows autoagglutinating nature and noteworthy adhesiveness independent of biofilm formation through a trimeric autotransporter adhesin (TAA), namely AtaA. TAA is one of the major outer membrane proteins widely distributing in Gram-negative bacterial species. Many studies have shown that TAAs from pathogens adhere to ECM proteins, such as collagen, fibronectin and laminin. In contrast, we previously reported that AtaA mediates bacterial cell adhesion to various abiotic surfaces nonspecifically. However, we have investigated the adhesive property mediated by AtaA only in a cellular level.

Objectives

Here, we report the characteristics of AtaA as a molecule. We attempted to purify the AtaA molecule and characterize its biophysical and adhesive properties.

Methods

The structural stabilities in thermal treatment and pH change were analyzed by transmission electron microscopy (TEM) and circular dichroism (CD). The adhesion property of the purified AtaA molecule was analyzed by quartz crystal microbalance (QCM).

Conclusions

The TEM images of the purified AtaA showed slightly curved fiber structure of 225 nm in length with two globular domains at the amino-terminal and near the carboxyl terminal. The adhesive function of the purified AtaA remained after pH treatment. In contrast, that of the purified AtaA decreased with disruption of its structure after thermal treatment more than 80°C. These results suggest that AtaA molecule has relatively high resistance against strict conditions and its fiber structure plays crucial role in its adhesiveness to abiotic surfaces.
Background
Halophytes are highly salt tolerant plants able to survive and flourish in coastal salt marshes and salt steppes. Lebrija marshes are situated at SW-Spain (36º 64' 29'' N - 6º 12' 92'' O), covering an area of 13000 has. Several halophytes are able to grow in this zone. Among them, we have selected *Arthrocnemum macrostachyum*, a plant with proven desalination capacity, as source for isolating halotolerant rhizosphere microorganisms.

Objectives
Isolate halotolerant rhizosphere microorganisms.

Methods
A total of two hundred bacteria have been isolated from plants grown in two different zones of the Lebrija-marshes at two different seasons (autumn and summer). These strains have been tested for their capacity to grow under high salinity conditions and six strains have been selected.

Rhizospheric soil samples were suspended and homogenised in sterile mineral buffer. Serial dilutions were plated on Nutrient Agar supplemented with 300 mM NaCl and cycloheximide (100 mg/l), adjusted at pH 7.4, at 28 °C for 5 days.

Conclusions
Three of the six strains, named Cs16b, Cs25 and Hvs 18 were unable to grow without salt in the culture media (halophyte strains) being the last one (Hvs18) able to support up to 20% NaCl. The other strains (Hv14b, Hv16 and C14) were halotolerant ones. All of them have been sequenced and characterized. Cs16b, Cs25 and Hv16 have been described as novel species within the genera *Microbulbifer*, *Labrenzia* and *Kokuria*, respectively.

This work was supported by INIA Project RTA-2012-00006 cofunded by FEDER.
Background
Diarrhea is influenced on specific pathogenic microbes. Wood creosote is an anti-diarrhea medicine to have been used for about 100 years in the northeastern Asia.

Objectives
We investigated the change of intestinal microbiota in rats to be fed with wood creosote.

Methods
Sixty rats (7 weeks old) were divided to 5 groups; control, antibiotic group, 0.2%, 0.4% and 0.8% wood creosote groups. The wood creosote and the antibiotic were supplied with feed for 4 weeks. To analyze intestinal microorganisms, the pyrosequencing was performed with 16S rRNA genes of microorganisms from intestinal contents.

The control group distributed 6 phyla including 229 species of microorganisms, the antibiotic group 7 phyla including 245 species, 0.2% wood creosote group 7 phyla including 284 species, 0.4% wood creosote group 7 phyla including 276 species and 0.8% wood creosote group 7 phyla including 324 species, respectively. *Clostridium* to belong to *Clostridia* that include many species of pathogenic microorganisms was 33.4, 44.3, 37.0, 35.5 and 11.6 per cents in the control group, the antibiotic group, 0.2, 0.4 and 0.8% wood creosote groups, respectively. The wood creosote dose-dependently decreased *Clostridium* in the large intestine.

Conclusions
We think that the supply of the wood creosote increases the diversity of the intestinal microorganisms and inhibits the propagation of specific pathogenic microorganisms.
FEMS-2378
Microbial communities

DEEPWATER HORIZON SPILL’S IMPACT ON SHIPWRECK MICROBIOMES AND SURROUNDING ENVIRONMENT

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Background
Shipwrecks serve as artificial reefs in the deep ocean. Due to their random distribution and inherent diversity compared to the surrounding environment, shipwrecks are ideal ecosystems to study pollution impacts, microbial distributions and marine corrosion.

Objectives
To comparatively assess the Deepwater Horizon spill’s impacts on shipwreck microbiomes and the synergistic effects on these communities and surrounding environment. Field activities will generate material and environmental samples required to address questions related to contaminant effects resulting from the spill and the interaction of spill related contaminants on the microbiota. Microbial action at each site will be interrogated to determine if exposure to hydrocarbons and dispersant-based chemicals have impacted the artificial reefs.

Methods
Porewater from sediment samples were analyzed by ion chromatography and inductively coupled plasma optical emission spectrometry to quantify major elements. Concentration and stable isotope ratios for sediments were determined using an elemental analyzer interfaced to an Isotope-Ratio Mass Spectrometer (IRMS). The molecular analysis of sediment from wreck sites was performed on the Ion Torrent Personal Genome (PGM) machine to provide a comprehensive assessment of microbiome composition. An analytical pipeline to extract taxonomic information from PGM sequences was completed to provide metabolic functional information from DNA sequences.

Conclusions
Six shipwrecks in the northern Gulf of Mexico were investigated. Various laboratory tasks have been completed since the second field research expedition funded by this project. Taxonomic classification of shipwreck microbiomes and metabolic information extracted from sequence data yielded new understanding of microbial processes associated with shipwrecks in the deep biosphere.
PREDICTING THE DYNAMICS AND BIODEGRADATION POTENTIAL OF RIVER MICROBIAL COMMUNITIES IN ANTHROPIZED WATERSHEDS: A WET LAB APPROACH.

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Background
River waters represent less than 0.0002 % of global water resources but they are essential for human life. Yet they are particularly exposed to anthropogenic stresses and risks of degradation. In a context of diminishing access to clean resources, lotic microbial communities play a key role in water self-purification. However they remain largely uncharacterized, and the link between lotic bacterial diversity and bacterial community stability and performance has not been studied.

Objectives
The goal of our project is to determine which factors drive the response of lotic microbial communities to organic pollution and how they tolerate, resist, exploit this pollution and/or go back to original status (resilience) downstream sewage outfalls. More specifically, the role of (i) native community structure (both phylogenetic and functional), (ii) wastewater bacteria released with organic matter and (iii) type of disturbance (regular/occasional, low/high loads) are studied using bacterial strains isolated from the Zenne river, Belgium.

Methods
Random artificial assemblages of these strains are used to test the effect of microbial community structure (richness, evenness, architecture, metabolic diversity…) on community performance and recovery from stress.

Conclusions
We hypothesize that such simplified, “wet lab” approaches can help to develop predictive tools of the dynamics and activity of bacterial communities in urban watersheds and in fine contribute to a better protection and restoration of river ecosystem services.
Background
Mixed microbial cultures have been proposed as an efficient way to deconstruct plant waste. The characterization of specific microbial consortia could be the starting point for novel biotechnological applications related to the efficient conversion of lignocellulose to cello-oligosaccharides, plastics and/or biofuels.

Objectives
Here, the diversity, composition and functional profiles of two novel microbial consortia are reported, on the basis of replicated aerobic wheat straw enrichment cultures.

Methods
Soil as microbial source and three sequential-batches were evaluated by amplicon-pyrosequencing (bacterial 16S rRNA and fungal ITS1) and whole metagenome sequencing.

Conclusions
Both systems appeared to indeed incite a reshaping of the bacterial communities, with reductions in richness and increases in prevalence of particular members of the Enterobacteriales, Pseudomonadales, Flavobacterales and Sphingobacteriales. Among the fungal players with high biotechnological relevance, co-existing with bacteria ones, we detected member of the genera Coniochaeta, Acremonium and Trichosporon. Metagenomic analyses revealed an overrepresentation of diverse carbohydrate transporters, plant polysaccharides sensing proteins and several genes encoding enzymes of the glycosyl hydrolase families GH2, GH43, GH92 and GH95 in the two consortia, as compared to the soil inoculum. These analyses are the starting point for synthetic expression systems and could enhance our understanding of microbial lignocellulose bioconversion in engineered or natural soil environments.
FEMS-0801
Microbial communities

MYCORRHIZAS AND THE ECOLOGY OF FOREST SUCCESSION
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Background
Identifying the drivers of plant community succession is a central theme in ecology. Research has shown that plant-soil feedbacks drive community assembly and has inspired many theories of species diversity. An important unsettled issue is whether mycorrhizas mediate plant-soil feedbacks to determine tree community succession.

Objectives
We show that ectomycorrhizas (but not arbuscular mycorrhizas) define interactions among trees and therefore have the ability to trigger and eventually to arrest tree community succession.

Methods
Using reciprocal invasion design with ectomycorrhizal mycorrhizal (EcM) and arbuscular mycorrhizal (AM) tree species, we tested for the potential for shared compatible mycorrhizal fungi to modulate interactions between invading seedlings (‘daughters’) understorey and resident trees (mycorrhizal ‘mothers’) in replicate mesocosms.

Conclusions
Characterizing mycorrhizal fungal communities using pyrosequencing, we find that EcM seedlings favored EcM fungi over AM fungi, but AM seedlings exhibited no mycorrhizal preference. The results from growth survey and physiochemical data confirm (i) invasibility of EcM fungi (and hence EcM seedlings) to AM-dominated communities, (ii) detrimental effects of EcM fungal diversity on AM daughters and mothers, and (iii) the ability of EcM fungal network in reducing the relative growth of EcM daughters to EcM mothers. The results demonstrate that invasion of EcM fungi facilitates establishment of EcM trees on the presence of AM trees, and eventually arrests the relative growth of EcM daughters to EcM mothers, a relationship potentially regulating tree community succession. The observed shifts from positive to negative plant-soil feedbacks involving EcM may explain why the climax temperate forests are predominantly ectomyorrhizal.
Microbial communities

MICROBIAL INTERDEPENDENCIES IN AN ENRICHED AEROBIC MIXED METHANOTROPHIC COMMUNITY

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Background
Aerobic methanotrophic bacteria are capable of both energy conservation and growth with methane as carbon source. Next to this unique feature they can be primary producers, supporting microbial foodwebs, particularly in aquatic environments. Recently it has been suggested that disparate microbial populations, including the methanotrophs are responsible for methane cycling rather than individual methanotrophs. The interactions between methanotrophs and heterotrophic partners have also been shown to be highly specific and beneficial to the rate of methane oxidation. Methane is an important greenhouse gas and methanotrophic communities could be used for both mitigation and biotechnological applications, warranting investigations into their composition.

Objectives
The aim of this work was to gain insight in the primary and secondary consumers from an active laboratory methanotrophic enrichment culture.

Methods
A methanotrophic mixed culture from various sources was incubated with both $^{13}$C labeled methane and $^{12}$C methane. At several points in time the 16S rRNA genes of the community were amplicon sequenced from both the light and heavy fractions to identify the first partners to incorporate labeled methane (methanotrophs and primary consumers) and later partners (secondary consumers). Interesting timepoints were selected based upon denaturing gradient gel electrophoresis profiles at 24 and 60 hours of incubation.

Conclusions
The main methanotrophic community driver was Methylophilus sp. Unique primary partners in the heavy fraction at 24h were members of the genera Acidivorax, Curvibacter, Pseudoxanthomonas, Methylophilus and Rhodobacter. At 60h the genera Sphingobacter, Kaistia, Sphingomonas, Flavobacterium, and Methylobacillus represented the heavy fraction. Isolation of these partners from the community could help elucidate their influence on methane cycling.
IMPROVEMENT OF SEQUENCE-BASED BACTERIA IDENTIFICATION IN POLYBACTERIAL SPECIMENS BY RIPSEQ MIXED SOFTWARE

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Background

Sequencing-based molecular methods have become a substantial tool in microbiology but analysis of polybacterial samples still remains a methodological challenge. RipSeq Mixed represents software-based technique, when up to 3 mixed chromatograms can be separated in silico, however this possibility can be inadequate for some types of microbiological material.

Objectives

We present the use of Denaturing Gradient Gel Electrophoresis (DGGE) followed by sequencing in combination with analysis of polymicrobial sequences by RipSeq Mixed software. Combination of these methods was used for detection of bacteria in urine and ureteral stent sonicates. Bacteria were detected also by culture-dependent techniques.

Methods

A total of 45 specimens of 15 patients were collected. The V3-V5 region of the 16S rRNA gene was amplified and multiple length homogeneous amplicons were separated by DGGE. Bands were excised and reamplified PCR products were sequenced. RipSeq Mixed software was used in case of mixed chromatograms.

Conclusions

There were 40 PCR-positive specimens (80 % of all specimens), 32 specimens were able to analyze only with PCR-DGGE. Ten bands (20 % of all bands) exhibited mixed nature, so that 8 specimens were identified by the combination of PCR-DGGE and RipSeq Mixed. The combination of established PCR-DGGE method and RipSeq Mixed analysis can eliminate failures of both approaches. More than 3 bacteria in a specimen can be detected and mixed chromatograms can be separated.

SOIL METHANOTROPHIC COMMUNITIES IN AN ALPINE WETLAND OF THE TIBETAN PLATEAU

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Background
Methane (CH₄) plays an important role in global warming and has been responsible for approximately 20% of the Earth’s warming since the pre-industrial period. However, most CH₄ is consumed by methanotrophs before its emission into the atmosphere.

Objectives
To understand the methane oxidizers in wetlands, the largest source habitat of atmospheric CH₄.

Methods
We used quantitative qPCR and functional gene-targeted 454 pyrosequencing to elucidate the abundance and diversity of aerobic and anaerobic (N-DAMO) methane oxidizers in the Zoige Wetland, which is located on the Tibetan Plateau and is the largest alpine wetland in the world.

Conclusions
Our results demonstrated that type I methanotrophs were the primary aerobic methanotrophs, and *Methyllobacter* comprised approximately 50% of the community. Type II methanotrophs, represented by the genus *Methylocystis*, were also detected, and the presence of N-DAMO bacteria in the Zoige Wetland was confirmed by the quantification and pyrosequencing analyses of N-DAMO 16S rRNA and *pmoA* genes. Furthermore, aerobic *pmoA* gene ranged from 10⁷ to 10⁸ copies·g⁻¹ of fresh soil, and the 16S rRNA gene of the NC10 phylum ranged between 10⁴ and 10⁵ copies·g⁻¹ of fresh soil. The results suggest that the aerobic and anaerobic methane-oxidizing communities together affect CH₄ cycle at the Zoige Wetland.
TRACKING RELATIONS AMONG BACTERIAL AND PROTOZOAN COMMUNITIES IN WASTEWATER TREATMENT PLANTS

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Background
Understanding the interactions between prokaryotic and eukaryotic populations in complex environments can be challenging. Particularly, there is a recognized difficulty in establishing how the interactions between the bacterial and the protozoan populations can affect the performance of wastewater treatment plants (WWTP).

Objectives
In order to determine the relationships between microbial communities (protozoa, metazoa and filamentous bacteria) and abiotic parameters (physical-chemical and operational), thirty seven WWTP with activated sludge process were studied in Portugal, during two years, in a trimestral sampling strategy.

Methods
Samples were collected to enable a molecular characterization of the microbiota. In order to ensure the diversity of microorganisms, 100 samples were selected based on their geographic localization and time-spaced to track relationships within bacteria and protozoa. A 16S rRNA gene PCR-DGGE approach was carried out for bacterial community fingerprinting. The resulting profiles were analysed together with the results obtained from the survey of the protozoa populations. Finally, the study of the correlations between the physical-chemical and operational parameters and the microbial communities was also investigated.

Conclusions
The study of these interactions constitutes a strategy for the understanding of how the different metabolic groups of prokaryotic and protozoa are affected by the relations between this populations, allowing a better prediction of the overall community dynamics. This will allow setting the conditions that better contribute to improve the wastewater treatment while increasing the knowledge of overall communities networking in complex environments.
INTRA-AMNIOTIC AND VAGINAL MICROBIOMES AT THE END OF PREGNANCY
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Background
The microbiota of the genital tract of pregnant women remains to be elucidated. Recent studies of the bacterial populations in different groups of pregnant women have yielded considerable amount of information on their bacterial communities. However, few comparisons have been performed between the vaginal and intra-amniotic microbiomes of individual women, and of women of different racial backgrounds.

Objectives
To investigate the compositions of the vaginal and intra-amniotic bacterial populations in late pregnancy of women from different racial backgrounds, and to compare the compositions of their genital microbiomes.

Methods
Vaginal and intra-amniotic swabs were obtained from 90 women at delivery by Caesarean section. Participants were administered a questionnaire including demographic and pregnancy data. The identity of the taxa present was determined by ultrafast sequencing of the V3V5 regions of the 16S rDNA gene of samples. Species diversity and richness, and Shannon’s diversity analyses were performed on the sequence data of each woman. SIMPER analysis showed the contribution of each species to the vaginal and intra-amniotic microbiomes. PERMANOVA and PERDISP analyses showed that both genital microbiomes depend on racial background.

Conclusions
At variance with commonly held assumptions, the analyses identified in the intra-amniotic cavity of the women in the study rich and diverse bacterial populations that were significantly different to the vaginal populations. The results suggested that the intra-amniotic microbial profile remains quite stable independently of the variability of the vaginal microbial profile. The outcomes of this study have potentially important implications for the management of urogenital infections during pregnancy.
Background
Interactions among members of polymicrobial infections can result in altered pathogen behaviours such as enhanced virulence, biofilm formation or antibiotic tolerance, which may influence the disease phenotype and clinical outcome of the infection. *Pseudomonas aeruginosa* and *Staphylococcus aureus* are important opportunistic human pathogens and are both part of the polymicrobial infection communities in human hosts. Cell-cell communication can play a major role in the interaction behaviors between these bacteria, which subsequently can affect the physiology of the individual strains. However, the extent to which evolutionary processes may remodel interspecies interactions and bacterial physiology during the course of infection and therapy is currently not understood.

Objectives
Therefore, the aim of this study was to analyze *in vitro* interactions between *S. aureus* and a human host adapted *P. aeruginosa* strain from a dominant lineage, DK2, that have evolved through decades of growth in chronically infected patients.

Methods
By using a combination of *in vitro* agar assays, molecular genetics and MALDI-TOF imaging mass spectrometry we demonstrate that DK2 – in contrast to other *P. aeruginosa* strains - show a commensal-like interaction with *S. aureus* during *in vitro* co-culturing, where the growth activity of *P. aeruginosa* was increased in the presence of *S. aureus*. The commensal interaction was observed both under aerobic and anaerobic conditions as well as during growth on artificial CF sputum medium, however was abolished with *S. aureus* strains mutated in the *agr* quorum sensing system.

Conclusions
Our results suggest that metabolic transformations during interaction with *S. aureus* enhance the growth activity of *P. aeruginosa*. 
OCCURRENCE AND ACTIVITY OF METHANOGENIC ARCHAEOA IN AGRICULTURAL BIOGAS PLANTS

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Background

Anaerobic digestion of organic substrates is a complex process which is accomplished by the cooperation of different microorganisms. The last part of the biogas process, the methanogenesis, is mostly the bottle neck of the whole process and is still not completely understood. Recently, molecular analyses of the microbial populations shed light on the microbiological ‘black box’, but on DNA-level, the whole population consisting of dead and inactive cells is analyzed.

Objectives

In this study, biogas plants were analyzed to investigate which fractions of the methanogenic community are transcriptionally active under different process conditions.

Methods

Different lab- and full-scale biogas reactors with different substrate mixes and process conditions were analyzed for the community composition of methanogens on DNA- and mRNA-level. After DNA- and RNA-extraction, (RT-)PCR-cloning targeting the mcrA/mrtA gene coding for methyl-coenzyme M-reductase was conducted. Sequences were checked for chimeras, aligned and a phylogenetic tree was constructed.

Conclusions

Hydrogenotrophic methanogens dominated in the investigated agricultural biogas plants, and acetoclasts were only present at high retention times. In well running systems only small differences were seen between the whole (DNA) and active (cDNA) methanogenic communities, and certain Methanosarcinaceae and Methanobacteriaceae dominated. During process disturbances, the active community mostly differed considerably from the whole community, and certain
*Methanobacteriaceae* took over the main transcriptional activity, not necessarily with productivity loss. A small part of the present methanogens obviously can take over the main part of the activity. The community should thus be analyzed on cDNA level in order to analyze the active part of the community reliably.
PUBLIC HEALTH SURVEILLANCE OF NOSOCOMIAL INFECTIONS IN SOUTH-EAST, NIGERIA

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Background
Hospital-acquired infections (HAIs) are constituting major threat to hospitalization world-over. They occur in several different patient groups, especially patients on ventilators in intensive care unit where the rate of the pneumonia may be as high as 3\% per day.

Objectives
This study was undertaken to unveil the burden of HAI in South-East, Nigeria.

Methods
Structured questionnaires totaling 660 were administered personally to healthcare workers consisting of 96 doctors, 170 nurses, 24 pharmacists and 40 medical laboratory scientists from randomly selected government hospitals and 90 doctors, 180 nurses, 30 pharmacists and 30 medical laboratory scientists from randomly selected private hospitals. Study was carried out between April and July 2013. The study shows that the prevalence of nosocomial infection in the hospitals surveyed is in the following descending order: urinary tract infection (34.9\%), Staphylococcus aureus (31.0\%), Gastroenteritis (27.1\%), Hospital acquired pneumonia (20.3\%), Candida albicans/Aspergillus spp (10.8\%), Pseudomonas aeruginosa (10.5\%), Tuberculosis (9.0\%), Clostridium difficile (3.9\%), ventilator associated pneumonia (3.6\%), Methicillin and vancomycin resistant bacteria (1.5\%). The study also showed that the frequency of occurrence of HAP is higher in government than private hospitals. This can be attributed to the higher population of patients leading to overcrowding. The modes of transmission of hospital acquired pneumonia was observed to follow this order airborne > contact with blood and body fluids > contaminated instruments > contaminated hands > needle sticks.

Conclusions
The prevalence of HAI in South-east, Nigeria is high. Efforts are needed to reduce the incidence, improve the quality of life of patients and healthcare workers and
improve healthcare delivery.
Characterizing and monitoring the dynamics of microbial communities has become an important research field in microbial ecology. Yet conventional microbial (e.g. heterotrophic plate counts) and molecular (e.g. Denaturing Gradient Gel Electrophoresis) workflows are time consuming and laborious.

Objectives
Here, we applied a novel statistical pipeline on flow cytometry data in order to perform rapid high-resolution and objective monitoring of the microbial community dynamics in an oligotrophic engineered aquatic system. The obtained results were then related to the imposed operational conditions (pH, temperature, conductivity).

Methods
A full-scale engineered, oligotrophic cooling water system operating on a nuclear test reactor was used as model system. This system operated under 3 week cycles and was monitored throughout two full cycles by flow cytometry and on-line physicochemical sensors. Aside from generating phenotypic community fingerprints, flow cytometric analysis also allowed the assessment of total cell counts, nucleic acid content and cell viability. Additionally, the exact microbial community throughout the cycles was determined with 16S rRNA amplicon sequencing.

Conclusions
Community fingerprinting indicated a highly dynamic community shift during the operation of the cooling water system. Five distinct phenotypic communities related to start-up, operational periods and inactivity of the reactor were identified. Conductivity and temperature were the predominant physicochemical factors but only partially explained the observed community shifts. The cooling water microbial community further showed an oscillating pattern in cell concentrations and nucleic acid content on both weekly and daily scale. The output of this phenotypic “fingerprint” approach was validated using high-throughput 16S rRNA amplicon sequencing.
COMPARATIVE ANALYSIS OF BACTERIAL DIVERSITY IN HEAVY METAL POLLUTED SOILS ESTIMATED BY MOLECULAR AND CULTIVATION-BASED TECHNIQUES

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Background

Pb-Zn smelter - KCM is the largest enterprise in the Balkan Peninsula producing lead, zinc and precious metals.

Objectives

To assess bacterial diversity in heavy metal polluted soils via molecular and cultivation-based techniques.

Methods

16S rDNA approach, as well as cultivation-based techniques were applied.

Conclusions

Molecular analysis showed that microbial community in the studied samples consisted in different extent of Proteobacteria (a, b, g and d), Actinobacteria, Holophaga/Acidobacteria, AD1, Cytophaga/Flavobacterium/Bacteroides, Cytophaga, BD and novel bacteria. In the highly polluted soil sample - KCM 3 g-proteobacteria were not detected. Cultivation-based analysis showed presence of heterotrophic aerobs, heterorphic anaerobs, sporeforming bacteria, denitrifying bacteria, amonifying bacteria, nitrifying bacteria, Fe(II)- oxidizing bacteria, Mn(II) oxidizing bacteria, Fe(III) redicing bacteria, Mn(IV) reducing bacteria, colourless sulphur bacteria, cellulose degradating bacteria, oligocarbophiles, metal-leaching bacteria Acidithiobacillus thiooxidans, Acidithiobacillus ferrooxidans, Acidithiobacillus denitrificans,
Acidithiobacillus thioparus except fungi and actinomycetes. Both analyses revealed that the microbial communities in the polluted area were strongly affected.
INFLUENCE OF VERY EARLY EXPOSURE TO CEFUROXIME VIA THE MOTHER ON GUT MICROBIOTA COMPOSITION IN TERM BABIES BORN BY CAESAREAN SECTION

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Background
Cefuroxime is an antibiotic effective against a wide range of Gram negative and Gram positive bacteria. It is standard procedure to give the mother a single prophylactic dose (1500 mg) of cefuroxime in relation to caesarean section (CS). If the antibiotic is given before the umbilical cord is cut the antibiotic will pass from the mother to the infant. However, the effect of this very early exposure to cefuroxime on infant gut microbiota (GM) colonization is still not known.

Objectives
To investigate if very early exposure to cefuroxime in CS born infants influences GM colonization.

Methods
42 pregnant women (BMI<30) scheduled for elective CS were recruited at Odense University Hospital, Denmark. Two mothers were expecting twins meaning that 44 infants were included in the study. The mothers were randomly assigned to either receiving cefuroxime before skin incision or immediately after the umbilical cord were cut. Faecal samples were collected from all infants 10 days after birth and GM composition determined by MiSeq-based tag-encoded 16S rRNA gene targeted high throughput amplicon sequencing.

Conclusions
The GM of both groups were dominated by the genera *Bifidobacterium*, *Clostridium* (relative abundance of both genera ~5-10%), *Veillonella* (~25-30%) and family Enterobacteriaceae (~25-30%). Testing with analysis of similarity (ANOSIM) on both unweighted (p=0.872, R=-0.0401) and weighted (p=0.140, R=0.0317) uniFrac distance matrices analysis revealed no major differences between the 2 groups. In conclusion, early and short exposure to cefuroxime does not have major effects on infant GM colonization.
QUANTITATIVE CHANGES OF CLOSTRIDIUM SP. IN HUMAN INTESTINAL TRACT OF DIFFERENT AGE GROUPS

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Background

Characterization of changes in gut microbiota during life-time is the first step in elucidating its role in health and disease. Bacteria belonging to the Clostridium sp. both commensal and pathogenic are highly abundant in the human gastrointestinal tract. However, during life the abundance of different predominant Clostridium species of healthy persons is not well studied.

The aim of the study was to assess the bacterial counts in Clostridium groups of intestinal microbiota of healthy humans within different life cycles.

Objectives

The study group comprised healthy infants (17 male/8 female; 11.9±0.6 month), children (13 male/12 female; 5.3±0.2 year-old), adults (4 male/21 female; 48.2±6.6 year-old) and elderly (9 male/14 female; 72.9±5.0 year-old).

Methods

DNA from fecal samples was isolated by using QIAamp DNA stool mini kit (Qiagen). The counts (plasmid copy number/g feces) were detected by real-time PCR with specific primers.

Conclusions

Four dominant Clostridium groups (C. leptum, C. perfringens, C. coccoides and C. difficile) were detected. Infants were more colonized with C. perfringens group bacteria in comparison to children, adults and elderly (p<0.001, for all respectively). Children were more colonised with C. coccoides group bacteria in comparison to elderly (p=0.005). The counts of C. leptum group bacteria were highest in adults than in elderly and infants (p<0.001, both respectively). C. difficile prevailed in elderly and infants than in adults (10/23 (43.4%), 7/25 (28%) vs. 0/25 (0%), p<0.001, p=0.005; respectively).
Our study confirms that the quantitative composition of human intestinal *Clostridium* groups changes during life differentially. This data may be used for comparable analysis of the gut microbial composition between healthy persons and persons with particular disorders in different age groups.
MICROBIOLOGICAL COMMUNITIES OF THE WATER OF GELLÉRT BATH (BUDAPEST, HUNGARY) STUDIED BY CLASSICAL AND SPECIAL CULTIVATION TECHNIQUES

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Background

Thermal baths are unique water environments combining a wide variety of natural and anthropogenic ecological factors. There is limited information on the microbiology of thermal baths in their complexity, tracking community shifts from the thermal wells to the pools.

Objectives

In the present study the “normal” microbial community of the well and the pool water in Gellért bath was studied in detail.

Methods

To isolate bacteria 10% R2A and minimal synthetic media (with „bath water”) with agar-agar and Gelrite were used after prolonged incubation time, moreover polyethylene blocks covered with media were also applied. Strains were identified by sequencing their 16S rRNA gene after grouping them by ARDRA.

Conclusions

On each applied media and from each sample the dominance of a-Proteobacteria was characteristic though their diversity differed among samples. The largest groups in the pool water belonged to the Tistrella and Chelatococcus genera. The most dominant member in the well water was a new taxon, its similarity to Hartmannibacter diastrophicus as closest relative was 93.62%. From each sample novel species could be isolated belonging to genera Nocardioides, Rhizobium and Brevundimonas. Deinococcus and low G+C Gram positives appeared only in the pool water. In order to cultivate novel taxa the best method was the combined use of minimal synthetic media, special cultivating techniques and increased incubation time.
CO-OCCURRENCE OF ATOPOBIUM VAGINAE AND GARDNERELLA VAGINALIS IN MALE URETHRA INDEPENDENTLY OF SEXUAL EXPOSURE

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Background
Bacterial vaginosis (BV) is the most common genital infection among women of reproductive age. It is characterized by a transition in vaginal microflora from Lactobacilli to anaerobic bacteria.

Objectives
Though BV has been strongly linked to sexual behaviour, an endless controversy over the sexual transmission of BV remains elusive. Hence, we aimed to investigate prevalence and co-occurrence of BV-associated bacteria in urethras of healthy adult men in accordance with sexual exposure by using species-specific PCR.

Methods
First-catch urine specimens, representative of urethral swabs, from 114 healthy male volunteers were collected between January-June 2014, in Konya/Turkey. DNA was extracted using QIAamp mini kit (Qiagen, Germany) from pellets of 15 mL of urine. Lactobacillus crispatus, Lactobacillus jensenii, Lactobacillus gasseri, Lactobacillus iners, Gardnerella vaginalis, Atopobium vaginae, Peptoniphilus spp., BVAB1, BVAB2, BVAB3, Megasphaera type I, and Prevotella spp. were investigated by species-specific PCR assay.

Conclusions
Of 114 participants, 49 were sexually-unexperienced, and 65 were sexually-experienced, and all were circumcised. The most common species were Peptoniphilus spp. (53.5 %), L. iners (50.8 %), G. vaginalis (50.8 %), and A. vaginae (20.1 %). There was no significant relation between any BV-associated bacterium and sexual exposure. The only significant difference was in the co-occurrence of A. vaginae and G. vaginalis (p=0.025) in the male urethra independently of sexual exposure. This result was compatible with similar reports studied on females. Since both A. vaginae and G. vaginalis are dedicated as key microorganisms in BV pathogenesis, more comprehensive studies on male counterparts are warranted.
VALIDATION OF THE INTELLICAP® SYSTEM AS A TOOL TO STUDY CHANGES IN THE SMALL INTESTINAL MICROBIOTA IN A DIETARY INTERVENTION STUDY

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Background

The gut microbiota is involved in regulation of host metabolic and immune pathways, and is an important target for dietary interventions. Human microbiota studies are mainly based on the analysis of fecal samples. The microbiota in the small intestine differs substantially from the fecal microbial community, but is at least as relevant for human health. Nevertheless, this ecosystem has barely been studied due to the invasive sampling procedures. The IntelliCap® system is an electronic medical device for the site-specific delivery of drugs in the gastrointestinal tract. The IntelliCap® system has been adapted to aspirate liquid from its environment, and may thereby offer a minimally-invasive tool for sampling from the human small intestinal tract.

Objectives

To explore whether the IntelliCap® system can be used as a tool to study the human small intestinal microbiota composition.

Methods

A randomized cross-over controlled dietary intervention trial in 10 healthy male volunteers will be performed in January 2015. The study is approved by the Ethics Committee and subjects signed informed consent. Two diets are used: a three-day high-protein versus high-carbohydrate diet, each preceded by a washout diet. These diets aim to target microbiota with different fermentation requirements. The IntelliCap® capsule is administered after both intervention periods to collect a distal small-bowel sample. Fecal samples are collected from the same individuals. Microbiota composition is analyzed by sequencing using Illumina technology and dedicated bioinformatics.
Conclusions

The IntelliCap® system may be applied to generate important new insights into the role of the human small intestinal microbiota in health and disease.
SPECIES DISTRIBUTION OF NON TUBERCULOSIS MYCOBACTERIA ISOLATES FROM NORTH OF IRAN

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Background

Nontuberculous mycobacteria (NTM) differ from tuberculous mycobacteria as most of them are ubiquitous and saprophytic. A few are considered to have the potential to infect humans. The immunological status of a person determines the advance of disease. On the other hand among environmental factors, water plays the main roles as a resource of contamination in transferring this group of microorganisms to human.

Objectives

The aim of this study is to identify and isolate NTM in the different region of Iran surface water.

Methods

62 water samples were collected from different regions of Northern Iran. Cetyl pyridinium chloride (cpc) 0.01% was used to decontaminate the water samples. After enrichment using filtration, all samples were incubated on Lowenstein-Jensen (LJ) medium at temperatures of 30 and 37 °C. Classical culture, biochemical and enzymatic methods are described for the identification of mycobacteria. hsp65 gene has more variability in its sequences, compared to the some more conserved genes in NTM, for identification of mycobacteria to species level.

Conclusions

There were different hsp65 gene PCR-RFLP profiles produced by digestion with BstEII and HaeIII. Dominant isolates were M. fortuitum. Other isolates of Mycobacteria were differentiated using hsp65 genes PCR-RFLP. This study indicates that notable percentage of water sources are contaminated with mycobacterium species. The exposure of immuno-compromised individuals to these sources can have crucial consequences. Interestingly the prevalence of mycobacterium species differs in diverse geographical areas.
BACKGROUND

In natural bacterial communities, the number of species existing at a low density levels is large and harbors the most of uncharted genetic information in nature. However, simple metagenomic sequencing methods can reveal those information for actively growing species unless a special method such as deep-sequencing or copy-number normalization is employed.

OBJECTIVES

In this study, we aimed to develop a method increasing relative abundance of rare-species genomes while reducing that of predominant species. We hypothesized that treating community samples by a germicidal agent normalize copy number of genomes in metagenome samples.

METHODS

We treated soil and insect samples by alkalinizing chemicals including quicklime (CaO), NaOH, and KOH, which kills microbial cells in general. For each sample treated with different concentrations of alkalinizing chemicals, bacterial community was analyzed by t-RFLP and pyrosequencing of 16S rRNA gene, and by enumeration and identification of culturable bacteria.

CONCLUSIONS

There were slight decrease of species richness in t-RFLP profile with stronger treatment (0-13% w/w CaO). However, overall diversity estimation (Shannon-Weaver diversity index) showed a steep increase at 3-5% CaO treatment. Chao1 estimates of richness for pyro-reads increased from 639±121 to 2,933±103 by optimal CaO treatment for soil samples. Rarefaction analysis on pyro-reads implied that richness of the untreated sample reached saturation while other richness values did not reach saturation. Shannon-Weaver diversity for pyro-reads increased from 1.8 to 6.7 by the optimal treatment. It could be concluded that the suggested treatments can increase
sensitivity of detecting rare species so that richness estimates increase more than five folds.
LONG-TERM SURVIVAL OF ENTEROCOCCUS FAECIUM OUTSIDE THE HUMAN HOST

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Background

Enterococcus faecium is a Gram-positive commensal of the gut of human and animals. However, in the last few decades it has acquired resistance to several antibiotics and has become an important cause of nosocomial infections. E. faecium can survive outside the human host for an extended period of time, which increases its ability to spread throughout the hospital and cause outbreaks.

Objectives

Our goal is to uncover the mechanisms used by E. faecium to survive outside the human host.

Methods

Using prolonged batch culture in brain heart infusion broth (BHI) or incubation in nutrient-free phosphate buffered saline (PBS) we determined the capacity of two clinical E. faecium strains (E745 and Aus0004) to survive at 20°C and 37°C.

Conclusions

We show that temperature has a critical role in the long-term survival of E. faecium. At 37°C viable counts in PBS drop by 90% after 2 – 3 days while at 20°C the viable counts remain stable for 5 days. During prolonged culture in BHI, viable counts decrease by 90% after 7 days, but at 20°C viable counts remain stable for 10 days. Our finding suggest that E. faecium has specific mechanisms to survive outside the human host for a prolonged period of time. Currently, high-throughput functional genomic experiments are being performed to identify the mechanisms which contribute to survival of E. faecium in nutrient-poor conditions at ambient temperature.
DEVELOPMENT OF A SIMPLE ANIMAL MODEL TO STUDY CHRONIC PSEUDOMONAS AERUGINOSA INFECTIONS

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Background

Chronic infectious disease is increasingly recognized as a major public health threat. The development of novel, more effective antimicrobial therapies is of utmost importance. Antimicrobial drug discovery suffers from high attrition rates in later stages of drug development due to a lack of \textit{in vivo} effectiveness of candidate drugs. Validation in an \textit{in vivo} infection model early in the process is likely to remedy this problem. However, animal models that enable the \textit{in vivo} study of chronic bacterial infections are typically complex and generally involve higher animals such as mice.

Objectives

We sought to develop a simple animal model that would allow the study of a prolonged bacterial infection by a clinically relevant pathogen.

Methods

We selected the opportunistic pathogen \textit{Pseudomonas aeruginosa} and the crustacean host \textit{Daphnia magna}. \textit{D. magna} juveniles were infected with varying doses of a virulent strain of \textit{P. aeruginosa}. While the highest infectious doses resulted in acute toxicity and complete mortality within 2 days after exposure, lower doses did not cause increased mortality compared to uninfected control populations up to 2 weeks after exposure. Infection with a \textit{P. aeruginosa} strain expressing green and red fluorescent proteins allowed visualization of infection sites using fluorescence microscopy. Labeled bacteria were still readily identified 2 weeks after exposure, revealing a stable chronic infection.

Conclusions

Together, our results strongly suggest that the simple \textit{P. aeruginosa-D. magna} model is a suitable tool for studying chronic bacterial infections \textit{in vivo} and support its further development for use in validation of candidate antimicrobials.
NEW INSIGHTS INTO PERSISTERS METABOLISM

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Background

Persistence is defined by a phenotypic switch occurring at low frequency (≈10⁻⁶) and generating a subpopulation tolerant to antibiotics. After removal of the antibiotic, persistent bacteria are able to regrow. Persistence is therefore thought to be responsible for chronic infections and could play a key role in the survival of biofilms to antimicrobials. Despite these observations, molecular mechanisms underlying persistence remain unclear.

Objectives

The aim of this study is to shed light on the molecular mechanisms by which phenotypic variability is generated in bacteria, allowing them notably to switch from an antibiotic susceptible state to a persistent state.

Methods

In a first approach, the main experimental parameters affecting E. coli persistence in our experimental conditions (aeration rate, overnight culture duration, growth medium, carbon source, ...) were identified. A standard and reproducible method for measuring persistence was then set up and validated. Using this method, we are currently investigating persisters genetics and metabolism. Approaches combining the use of deletion mutants, fluorescent transcriptional reporters and biosensors are used to determine which metabolic pathways are activated/shut down in persisters.

Conclusions

Our data confirmed that persisters consist of highly heterogeneous populations. In addition, different classes of antibiotics appear to generate different persistent populations, characterized by the activation of distinct metabolic pathways and stress responses. This indicates that molecular mechanisms underlying persistence are not unique and a variety of mechanisms are involved. Population (FACS analyses) and
single-cell (time-lapse microscopy) behaviours are currently analysed to get further insights into persisters mechanisms diversity.
DURING CHRONIC INFECTIONS SALMONELLA ACQUIRES MUTATIONS IN REGULATORS OF METABOLISM AND VIRULENCE GENE EXPRESSION

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Background

Salmonella enterica is responsible for diverse infections in mammals, including chronic infections. For Salmonella enterica serovar Typhi, the causative agent of typhoid fever, up to 6% of patients, will become life-long carriers.

Objectives

Despite the importance of chronic infections, the molecular mechanisms of chronic host association are poorly understood. The objectives of this study are to elucidate the mechanisms employed by Salmonella for establishment of chronic infections.

Methods

Mice were infected with low doses of Salmonella, and the development of chronic infections was followed over the course of the infection. After six weeks of infections mice were sacrificed and the livers and spleens were analyzed for the presence of Salmonella. Dominant clones of livers, spleens and fecal samples were whole genome sequenced to identify adaptive mutations.

Conclusions

Most clones had acquired one or two non-synonymous point mutations (single nucleotide polymorphism, SNP). Most SNP’s were associated with metabolism and virulence regulation. For example, a dominant clone from a fecal sample had a SNP in kdgR, a repressor of genes of the Entner-Doudoroff pathway, indicating metabolic adaptation in this clone. Re-infection experiments revealed that the kdgR-SNP mutant exhibited rapid and superior colonization of the mouse intestines resulting in a super-shedder phenotype within 1-2 weeks post-infection. Furthermore, the KdgR-SNP regulator protein could not be induced leading to constitutive repression of its target genes, supporting metabolic remodeling in the kdgR-SNP mutant. In conclusion, during a short chronic infection, Salmonella adapts to its host by modifying the activity of central regulators of metabolism.
FEEDBACK BETWEEN MICROBIAL INTERACTIONS AND EVOLUTION IN CHRONIC CF INFECTIONS

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Background

Although many infections are assumed to be monoclonal it is becoming clear that interactions between bacterial pathogens and other microbes at the infection site may alter pathogen behaviour. We have limited understanding of the evolution of microbial interactions and the extent to which interactions constrain or promote evolution of the interacting partners during the course of infection.

Objectives

Chronic airway infections in cystic fibrosis (CF) patients provide opportunities to study microbial evolution and its relation to microbial interactions. The CF airways are nearly always infected with a number of microorganisms, including *Pseudomonas aeruginosa*. CF infections are associated with genetic adaptation of *P. aeruginosa*, and to understand the adaptive mechanisms, we have focused on two transmissible *P. aeruginosa* lineages – DK1 and DK2 - that share a similar natural history: They both entered the Copenhagen CF Clinic more than 40 years ago, adapted in one or a few patients and subsequently spread to a large number of patients. Our aim is to uncover the genetic changes that determine the apparent dominance of these clones, and to investigate in which way microbial interactions shape the evolutionary pathways.

Methods

The systematic collection of bacterial isolates from lung expectorates of CF patients at the Copenhagen CF Clinic since 1973 has made it possible to investigate within-host evolution of both lineages in relation to both inter- and intraspecies interactions.

Conclusions

We show that the microbial CF community context impact evolution of the individual lineages, and that evolved changes within individual species can modify the behavior of the infection community.
THE ROLE OF CHLAMYDOPHILA PNEUMONIAE IN THE ETIOPATHOGENESIS OF SCHIZOPHRENIA: RELATIONSHIP BETWEEN SCHIZOPHRENIA BRAIN-DERIVED NEUROTROPHIC FACTOR (BDNF) AND NEUROTROPHIN-3 (NT-3) A WORLDWIDE RETROSPECTIVE STUDY

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Background

It's known that, in the occurrence of schizophrenia, genetic predisposition, neurodevelopmental disorders and environmental factors play role. It was suggested that synthesis of brain-derived neurotrophic factor (BDNF) and neurotrophin3 (NT3) were upregulated from monocyte/macrophages infected with Chlamydophila pneumoniae, considered among environmental factors.

Objectives

In this study we aimed to show at the worldwide level, for the first time, the relation of C.pneumoniae infection, BDNF and NT3 levels.

Methods

In this crosssectional retrospective study 50 patients with schizophrenia and 35 healthy controls(HC) were included. The C.pneumoniae DNA was investigated by RT-PCR from PBMC and IgA, IgG, IgM were investigated by immunofluoresans in both
group's serum. BDNF and NT3 levels were determined by ELISA. Chi square student's t and Mann Whitney U tests were used for statistical analyses.

Conclusions

A significant difference was found between PG and HC cases for chronic C. pneumoniae infection seropositivity (p<0.05). The mean serum BDNF and NT-3 levels were found to be significantly lower for PG members than for HC members (p<0.001). We couldn't detect any C. pneumoniae DNA in PBMC of 85 participants. In conclusion, although in HG, compared to HC, the presence of chronic C. pneumoniae infection was found remarkable high, BDNF and NT3 levels, were found significantly low in PG compared to HC, which is the first international study in the basis of C. pneumoniae, schizophrenia relation, and the detection of BDNF and NT3 levels, a conviction wasn't reached and we suggest that new studies, especially based on cohort and performed with large series including these tree parameters supported by molecular methods are needed.
MOLECULAR MECHANISM ANALYSIS OF PERSISTERS FORMATION WITH A CELL DIVISION MARKER

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Background
Persisters are multidrug tolerant cells due to their suppression of cell division. When bacterial population is exposed to antibiotics, persisters could survive and cause for the recurrence of infections. Although molecular mechanisms of bacterial persistence to antibiotics have been studied, a crucial gene had not been identified. Since the mechanisms of persisters formation are redundant, analysis targeting a specific single gene is not appropriate for persister researches.

Objectives
The objective of our research is to develop a marker for persisters that enable to isolate persisters and to analyze an exhaustive gene expression.

Methods
To distinguish non-dividing cells from dividing cells, fluorescent protein YFP and CFP were fused at the both N and C terminals of FtsZ as a cell division marker. The fusion proteins emit CFP fluorescence in non-dividing cells, moreover they emit fluorescence resonance energy transfer (FRET) signal at Z-rings in dividing cells. This mechanism enables to visualize non-dividing cell and dividing cell. Non-dividing persisters were isolated using a cell sorter and their transcriptome was explored using a microarray.

Conclusions
The fusion FtsZ emitted FRET fluorescence at the Z-ring. Sorted cells in low FRET signal fraction exhibited about 50 times higher survival rate after exposure to antibiotics (ofloxacin) than cells in high FRET signal fraction. Hence, our method attained an objective. Transcriptome analysis suggested following three hypotheses; i) persisters take up energy by anaerobic respiration, ii) membrane protein is sensing changes of the environment, iii) clinical thread is not caused only by persistence, but also by secretion system.
IN VITRO EVOLUTION OF PERSISTENCE BY CYCLIC TREATMENT WITH AMINOGLYCOSIDES IN PATHOGENIC ESCHERICHIA COLI AND SALMONELLA

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Background

Bacteria can evade antibiotic eradication by forming transiently antibiotic-tolerant phenotypic variants, called persisters. Persistence is linked to the recurrent and chronic nature of many bacterial infections, but mechanisms underlying this phenomenon have mainly been investigated in non-pathogenic Escherichia coli lab strains.

Objectives

Infections by many Gram-negative pathogens are becoming increasingly difficult to treat due to resistance towards multiple antibiotics. Since aminoglycosides often retain activity against these strains, there is revived interest in their clinical use. We investigated the evolution of persister levels in uropathogenic E. coli and Salmonella under periodic aminoglycoside treatment.

Methods

Previously, a method for the experimental evolution of persistence by cyclic antibiotic treatment was optimized in our lab. Here, we apply this method to investigate the evolution of persister levels in uropathogenic E. coli UTI89 and S. enterica Typhimurium SL1344.

Conclusions

Once daily, high dose aminoglycoside application rapidly selects for mutants with extremely high persister levels, while resistance levels (minimum inhibitory concentrations) remain unchanged. Likewise, mutants show increased persister levels in biofilms. Moreover, evolved uropathogenic E. coli mutants display cross-tolerance to fluoroquinolones, although high persister levels of S. enterica Typhimurium mutants seem antibiotic-specific. The dosing scheme of our in vitro experiments is roughly comparable to current clinical practice, underscoring the relevance of our findings. Future clinical evaluations of antibiotic dosing regimens
should include an assessment of persistence, as this could be an overlooked factor in failure of antibiotic therapy.
A SURVEY OF NASAL CARRIAGE RATE OF STAPHYLOCOCCUS AUREUS IN DIALYSIS PATIENTS

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Background

Staphylococcus aureus is the most common cause of cutaneous pyogenic infection. Some groups of individuals seem to be particularly prone to colonization with S.aureus. For example, patients undergoing chronic hemodialysis and users of illicit intravenous drugs

Objectives

The purpose of this study was determination of nasal carriage rate of S.aureus in dialysis patients

Methods

This survey was carried out in a descriptive analytic cross-sectional research in 2012. 71 patients of chronic renal failure were selected who had undergoing dialysis in dialysis ward of kashan. After obtaining consent form, Specimens with swabs were performed from anterior nares nose of the patients. The questioners having demographic and related risk factors were filled through interviewing. Anitimicrobial resistance S aureus of positive cases were determined by disk diffusion

Conclusions

The patients were 41 females (53.2%) and 36 males (46.8%). The mean of age in patients was 48.43 years and the mean of duration of dialysis was about 23.54 months. 12 cases were nasal carriers of S.aureus (16.9%). There was no significant statistical correlation between sex, age, job, underlying disease and duration of dialysis and nasal carriage. The most resistance of S.aureus was seen to meticillin and the most sensitivity was to rifampin. Resistance to cefazolin, cotrimoxazole 33.3%, vancomycin 50% and cloxacillin 75%. There was no S aureus was fully sensitive to vancomycin.

Considering to 16.9% frequency of S aureus carriage among dialysis patients and 50% resistance to vancomycin, periodic examination and proper treatment due to antibiogram results is recommended.
Background

*Haemophilus influenzae* is a commensal bacteria but can cause severe infections including meningitis, sepsis, bacteraemic pneumoniae, otitis media, sinusitis, and chronic bronchitis

**Objectives**

The present study was conducted to investigate the reservoir role of tonsils and adenoid tissue for colonization with *H. influenzae* bacteria in patients with chronic tonsillitis and adenoiditis and genotyping analysis of the strains isolated from both sites of the same patient.

**Methods**

A total of 200 specimens (100 tonsils and 100 adenoid tissue) were taken from 100 chronic tonsillitis and adenoiditis patients who undergoing adenotonsilectomy. All samples were bacteriologically cultured and Pulsed field Gel electrophoresis (PFGE) method was applied for all *H. influenzae* isolates which revealed a high growth in tonsil and adenoid tissue.

**Conclusions**

**Results.** It was found that 82 out of 100 (82%) tonsils and 79 out of 100 (79%) adenoid tissue cultures a total of 161 samples were exhibited 274 bacteria from different bacterial species. *H. influenzae* strains were formed % 32 (87/274) of the total bacterial growth. A twenty eight isolates from each of tonsils and adenoid tissue of 28 patients (56 isolates) were analyzed by PFGE method which revealed that 52 out of 56 (92.85%) *H. influenzae* strains of 26 patients were similar whereas four strains of two patients were different.

**Conclusion.** The tonsils and adenoid tissue are playing a reservoir role not only for colonization with *H. influenzae* and other different bacterial species but each site is also a reservoir for others and vice versa. For our opinion *H. influenzae* strain could be a key for prevention of chronic tonsillitis and adenoiditis.
MYCOBACTERIAL PERSISTENCE: USING MICROFLUIDICS AND SINGLE-CELL MICROSCOPY TO INVESTIGATE HOW MYCOBACTERIA SURVIVE ANTIBIOTIC EXPOSURE

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Background

Up to a third of the world’s population is estimated to be infected with Mycobacterium tuberculosis, the causative agent of tuberculosis (TB). A significant ability of this pathogen is its capacity to adapt to and survive stress. This may in part be the reason why TB treatment requires such a lengthy time – at least 6 months of antibiotics. Both of these observations may result from a propensity for mycobacteria to form antibiotic tolerant persisters.

Objectives

We investigate how growth rate affects mycobacterial survival when exposed to antibiotics, with an initial assumption that growth arrest is a major mechanism promoting bacterial survival under these conditions.

Methods

A microfluidic platform was used that allows observation of single-cells of M. smegmatis by widefield microscopy, with constant flow of defined medium, and rapid switching between different conditions. The drug rifampicin was used to dissect persister formation due to its bactericidal properties, as well as its clinical relevance in TB treatment. Nutrient limitation was used to determine how this affects persister formation.

Conclusions

Growth rate of individual cells prior to rifampicin exposure appears to have no impact on their ultimate survival, however growth arrest is clearly observed during antibiotic exposure in all cells. Nitrogen limitation and starvation appear to significantly decrease survival, whilst carbon limitation appears to have little effect. This work, though preliminary, may enhance our understanding of how persisters form in mycobacteria, and help in the design of novel intervention strategies to combat tuberculosis.
FREQUENCY OF ANTIBIOTIC APPLICATION DICTATES RAPID EVOLUTIONARY ADAPTATION OF BACTERIAL PERSISTENCE

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Background

Microorganisms employ different strategies to cope with changing environments. Typically, a dedicated sensor perceives a specific environmental trigger and signals an adaptive response. However, if changes are rare and/or abrupt, maintenance and time lag of sensory systems become too costly. Consequently, bacteria evolved the capacity of switching stochastically between phenotypes to hedge their bets. Such a bet-hedging strategy based on a non-genetic switch between a sensitive growth state and an antibiotic-tolerant, non-growing persister stage, protects bacteria against eradication by antibiotics. Persistence causes antibiotic therapy failure and is therefore responsible for the chronic and sometimes incurable nature of infections.

Objectives

Little is known about the evolutionary forces that determine persister levels. We therefore aim to delineate the evolutionary dynamics of persistence. Furthermore, we want to explore the theoretically suggested and intuitively appealing correlation between the level of persistence and the frequency of antibiotic application.

Methods

We used experimental evolution in \textit{Escherichia coli} to demonstrate rapid, reproducible and reversible changes in tolerance levels that optimize the population’s fitness in function of the treatment frequency. Daily antibiotic attacks select for single-nucleotide changes that confer high persister levels (up to 100\%) and shorten persister lag phase. Persister levels decrease again in the absence of selective pressure. When time between antibiotic strikes increases, intermediate persister levels evolve that correlate with treatment frequency.

Conclusions
We show that persistence is an evolvable phenotype that quickly adapts to environmental dynamics, which may have important implications for developing optimal treatment strategies of chronic bacterial infections.
FEMS-1539
Microbial persistence and chronic infections

GENOMIC AND TRANSCRIPTOMIC CHARACTERIZATION OF BURKHOLDERIA PSEUDOMALLEI WITHIN-HOST EVOLUTION IN FIVE CHRONICALLY INFECTED AUSTRALIAN CYSTIC FIBROSIS PATIENTS
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Background
Cystic Fibrosis (CF) is a genetic disorder that results in impaired mucociliary clearance of inhaled microorganisms. Over time, the airways of CF patients become colonised with different microorganisms, resulting in infections that cannot be eradicated. An uncommon pathogen of CF patients, but of rising significance, is Burkholderia pseudomallei (Bp). Bp is an opportunistic environmental pathogen that causes melioidosis, a potentially fatal disease that is endemic in most tropical regions, particularly northern Australia and Southeast Asia. Melioidosis is being increasingly described in CF patients returning from or living in melioidosis-endemic regions. Presentations of melioidosis in CF patients range from acute infection with rapid deterioration, to a chronic infection with slowly progressive deterioration in lung function.

Objectives
Unfortunately, the molecular basis for this chronic presentation remains entirely unknown.

Methods
To address this knowledge gap, we performed whole-genome sequencing on paired longitudinal Bp isolates from five Australian CF patients. To complement the genetic analysis, we have also performed stranded, deep sequencing of the transcriptome using Illumina RNA-seq. To best mimic the bacterial expression profiles in the CF lung, isolates were grown in an artificial sputum medium and RNA was extracted at late-log phase.

Conclusions
We will be reporting on the genomic and transcriptomic changes detected in the CF isolates, which have been shaped by within-host evolution. Similarly to common CF pathogens such as P. aeruginosa, we have observed a rise in multi-antibiotic resistance and changes to virulence factor expressions and metabolism.
ANTIBIOTICS STRONGLY AFFECT COST AND BENEFIT OF COMPETENCE IN BACILLUS SUBTILIS
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Background
Bacillus subtilis differentiates stochastically into the state of competence for transformation. Transformation is thought to speed up adaptive evolution in the presence of antibiotic stress.

Objectives
Here we address the question if the state of competence confers a strong benefit during episodic antibiotic treatment and acts as a persister state.

Methods
Using head-to-head competition experiments between strains with different rates of differentiation into the state of competence, we found that the cost of competence increases with the probability of differentiation. Competence was highly beneficial when competitors were treated episodically with antibiotics that inhibit cell wall synthesis, translation, transcription, or proton motive force. This benefit was independent of transformation. Time-kill kinetics in conjunction with single cell time lapse microscopy showed that the state of competence is a persister state. Addition of antibiotics did not increase the fraction of competent cells, indicating that competent B. subtilis are type II persisters. Finally, we tested whether the generation of phenotypic heterogeneity in terms of competence served as a bet-hedging strategy when B. subtilis was treated episodically with antibiotics. In competition experiments between strains with different probability of differentiating into the state of competence we found that populations that generate phenotypic heterogeneity minimize the cost of competence under benign conditions while increasing the probability of survival when challenged with antibiotics.

Conclusions
We conclude that B. subtilis uses a bet-hedging strategy for increasing its chance of survival when challenged with antibiotics by stochastically differentiating into the state of competence which acts as a persister state.
Background

Formation of small colony variants (SCV) of *Staphylococcus aureus* is a strategy (not only for *Staphylococcus aureus*) to overcome antibiotic treatment and to escape the immune system. This phenotype can persist for long time periods in patients with low pathogenic activity. One problematic feature of SCV is the ability to switch easily back to a more virulent phenotype.

Objectives

The aim of the study was to establish a test to quantify the ability of SCV’s to switch back to a non SCV phenotype and to test several clinic isolates.

Methods

Five *Staphylococcus aureus* SCVs, isolated from five different patients with cystic fibrosis were tested. All SCV’s were not able to grow in Mueller Hinton (MH)-broth. From each isolate 500 single colonies grown on Lysogeny Broth (LB)-Agar for 24 h showing SCV colony type were inoculated in 200µL MH broth for 24h. Growth in MH was confirmed by plating on MH agar plates and identification of *S. aureus* via MALDI-TOF analysis.

Conclusions

Four of the five SCV isolates showed stable phenotype with only 0-0.8% phenotype changes. In contrast one SCV changed the phenotype in 25.2%, which means at least one bacterial cells restored the ability to grow on/in MH media.

This test method is based only on one SCV characteristic and does not document if there is a total reoccurrence of the original wild type phenotype. Nevertheless it is an easy assay, and it will be a possibility to investigate the influence of different conditions on the stability of SCV’s.
BACKGROUND
In November, the University of the Basque Country (UPV/EHU) organized the 14th edition of the Science Week (ZientziaAstea). This annual event, aimed at audiences of all ages, tries to bring scientific concepts, methodologies and innovations to a non-specialist public by means of children's workshops, scientific shows, conferences and other activities. The microbiology has been present in this event from the first edition.

OBJECTIVES
In ZientziaAstea, we aim to bring the public diverse aspects of microbiology, to clarify certain misconceptions and to show its importance in our daily lives.

METHODS
Every year, our Department’s staff prepares materials and activities and attends the questions of school children and general public in a stand. Initially, the proposals consisted mainly in microbial cultures, photographs, diagrams and microscopic preparations. Year-to-year, proposals have become more complex, including models, videos, and posters with a common thread (hydrothermal vents, food, interactions) or hot topics (ebola). In the latest edition, we have incorporated the workshop Microorganisms working to refute the bad image of microorganisms and to highlight their benefits and their ecological and industrial importance.

CONCLUSIONS
We have observed a good response for assistance and acceptance by the public in these activities, but we have also detected the need to renew the issues and approach. As consequence, for the 15th edition we are preparing the workshop Small microbiology for small scientists that, through games and activities, will try to introduce children the microbial world (the microbial diversity in a big drop of water or in our body, the bacterial growth, etc…).
FEMS-2625
Microbiology education

WORKING ON ACCESSIBILITY OF MICROBIOLOGY EDUCATION FOR DEAF COMMUNITY: DEVELOPMENT AND USE OF A NEW SCIENTIFIC GLOSSARY IN SIGN LANGUAGE
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Background

University should be an open and accessible environment to all people, where knowledge and opinion can be exchanged. Deaf community often finds scientific language as a hard barrier blocking their education. Knowing that most of Sign Languages are usually extremely colloquial, the development of an international scientific Sign language seems to be as useful as necessary.

Objectives

This work aims to develop a handy and rigorous Sign Language that allows to all Deaf community stablish an independent university education and research avoiding the problems associated with continued use of interpreters.

Methods

We have created a sum of 320 signs for terms commonly used in science and related with Microbiology. These terms have been attributed to a sign, maintaining a logic in the use of Sign Language and a scientific adequacy. All the signs have been validated and used both by qualified interpreters and by deaf students from Complutense University of Madrid. All the terms have been included in a glossary and diffused firstly by using the Complutense University “Virtual Campus” and then on the web page of Spanish Society for Microbiology.

Conclusions

The main objective of this project has allowed us to develop teaching resources adapted to deaf students. The glossary has been applied to design dynamic accessible slides including explanations in Sign Language and illustrative multimedia tools making easier the educational process to deaf students in both, theoretical and practical lessons on Microbiology.
ANTIBIOTIC RESISTANCE: DEFINING THE SCIENTIST’S RESPONSIBILITY.
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Background

Antibiotics transformed global health and the fields of scientific discovery. The use of antibiotic tools is ubiquitous but the rise of antibiotic resistance is threatening modern medicine. Scientists are asking manufacturers to stop selling antibiotics as growth promoters; new initiatives to stimulate antibiotic research; and a better educated public and clinicians to champion responsible use of antibiotics.

Objectives
The next generation of biologists and microbiologists are taught the mechanisms of antibiotic resistance in formal academic settings. They learn about the ‘urgent need’ for new antibiotics, from a teaching and scientific research community urging responsible use of antibiotics. Yet students attend laboratory-based practical classes relying heavily on antibiotics. Should the scientific community examine how learning opportunities encompassing antibiotics and antibiotic resistance are incorporated across university curricula to ensure we provide young scientists with consistent and synoptic information? Should future scientists be encouraged to use antibiotics more responsibly and sparingly in both educational and research settings?

Methods

The roundtable discussion will debate whether we give conflicting messages to young scientists and will examine how learning opportunities encompassing antibiotics and antibiotic resistance questions are provided. Should we begin to practice what we preach and develop alternative practical classes where antibiotics and antibiotic resistance are not treated as disposable teaching tools?

Conclusions
Self scrutiny of our own educational practices will give us a stronger voice when we ask society to use antibiotics more responsibly and may encourage development of new technologies offering similar benefits and functionality as current tool set that is based on antibiotic resistance.
ACTIVITIES OF THE EDUCATION & COMMUNICATION DIVISION OF THE SPANISH SOCIETY FOR MICROBIOLOGY (D+D SEM): FOCUS ON YOUTH

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Background
The Spanish Society for Microbiology (Sociedad Española de Microbiología, SEM; www.semicrobiologia.org) has 11 specialised Divisions. Among them, the most recently created and transversal is the Education and Communication Division (Docencia y Difusión, D+D SEM; www.semicrobiologia.org/ddm). It hosts 180 members committed to Education in Microbiology at several educational levels within Spain and to promote our Science to Spanish-speaking people worldwide via the internet and social networks.

Objectives
D+D SEM aims to cover different degrees of formation:

Methods

First, a work team establishes links with professionals at primary and secondary schools, offering assessment and support to Science teachers about microbiological subjects, and aiming to the content of official teaching modules and text books. D+D SEM has recently published a book of educative short stories about Microbiology aimed to children.

Second, D+D SEM promotes the elaboration and open exchange of materials and resources for Microbiology Education for University degrees, including videos, podcasts, games, blogs, interactive on-line tools, webpages and innovative approaches for Microbiology teaching. Remarkably, our Division organizes yearly workshops on “Initiation to Research in Microbiology”, in which 20 brilliant and motivated University students selected nationwide share with senior Microbiologists their passion for research.

Third, at the postgraduate level, young researchers can find support and information on Master degrees, training courses or job opportunities at the Group of Young Researchers (sites.google.com/site/jovenesinvestigadoressem), hosted by D+D SEM.

Conclusions

We seek to share experience and coordinate our effort with similar initiatives in other countries.
WORKSHOPS ORGANISED BY STUDENTS FOR STUDENTS: AN ORIGINAL MULTIDISCIPLINARY APPROACH FOR TEACHING MICROBIOLOGY AND INFECTIOUS DISEASES.

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Background

During the last decades, efforts have been made for the development of innovative teaching in medical studies.

Objectives

To describe an Innovative Teaching Unit (ITU) implemented for the students being in the 4th year of medical studies in Angers University, France.

Methods

This ITU includes 20 original topics. Each year, each topic is attributed to 4 groups containing 4 students. After choosing an issue in the field of their topic, they have to organise and conduct a 3-hour workshop, to present a poster or an oral communication in a local congress, and to produce a written report. Each topic is supervised by two teachers from different medical specialties to ensure a multidisciplinary approach. The topic called ‘Major sanitary crises’ (MSC) was included in the ITU 4 years ago.

Conclusions

Two or three meetings between 2 teachers and students are necessary to prepare the workshops. In 8 of the 12 sessions of MSC that have been already completed, the students chose a topic in the field of microbiology and infectious risk (bioterrorism, Ebola virus, risks of waterborne diseases after a hurricane, and the use of antibiotics in intensive livestock farming). Students used original supports to animate workshops (homemade videos, street interviews, live staging, interventions of professionals and quiz). This ITU, combining problem-based learning and project-based learning, allows a total implication of students in the organization of workshop and obtaining original
teaching and supports on subjects that are not namely taught during their curriculum and prepare them early to scientific communications
Background

"Manimal" is a pioneering interdisciplinary training programme (ITP) (Master level) based on the “One World, One Health” concept promoted by international organizations (WHO, OIE, FAO), with an interdisciplinary approach for the management of global health and nutritional safety.

Objectives

Microbiology is a topic of great importance for global health and food safety and quality. Our objective was to determine the place of microbiology in this ITP.

Methods

This international ITP is organised in 5 thematic teaching units (TU): epidemiology, health: organisation, management and communication, biological and chemical risks, production chains, nutritional contamination and transfers. We particularly identified the interactions developed between microbiology teaching and other TUs in an interdisciplinary perspective.

Conclusions

Traditional lectures (face teaching or e-learning sessions) about microbiology are included in the “biological and chemical risks” TU. However, microbiology is also involved in interdisciplinary trainings via student groups around case scenarios (problem-based learning) and visits of production sites to understand the control measures implemented against the development of *Listeria monocytogenes*.
(interactions with the production chains TU). Two case scenarios are proposed: "Listeriosis and dairy industry" (interactions with the nutritional contamination TU) and "Antimicrobial resistance" (interactions with all other TUs). Groups containing 4 or 5 students with various initial educations (doctors, pharmacists, veterinarians, biologists, and agricultural engineers) and from different geographic origins (Asia, Africa, Europe) are constituted with the aim to make them solve the problems together. We envisage now to introduce microbiology in tutored projects (project-based learning) and to diversify the microbiological topics in the case scenarios.
THE SMALL WORLD INITIATIVE AT MCGILL: ENGAGING UNDERGRADUATES IN SCIENTIFIC DISCOVERY AND CROWDSOURCING THE SEARCH FOR NEW ANTIBIOTICS

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Background
The Small World Initiative (SWI), spearheaded by the Yale Center for Scientific Teaching, is an international, multi-institutional initiative which brings authentic scientific research addressing the real-world problem of antibiotic resistance to undergraduate life science students. SWI students culture soil microbes, screen for antibiotic activity against safe relatives of ESKAPE pathogens, and characterize their positive isolates by classical methods and 16s rRNA gene sequencing.

Objectives
McGill launched the SWI program in a sophomore Microbiology laboratory course of 105 students in fall 2014.

Methods
During the fall 2014 semester, the students collected soil samples from urban and suburban Montreal and cultured 2,500 bacterial isolates. Following activity screening, 104 isolates were prioritized for further characterization using classical biochemical and microscopy methods and molecular characterization.

Conclusions
Of the 104 isolates selected for further characterization, 94 gave positive results in 16S PCR, and of these, 71 yielded high quality 16S sequences. The isolates were equally divided between Gram-positive and Gram-negative, with a wide variety of genera represented. The students were highly engaged throughout the process and reported a high level of enthusiasm in course evaluations.
THE DEVELOPMENT OF ‘VIRUSES: A PRACTICAL RESOURCE FOR POST-16 BIOLOGY TEACHERS’

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Background
Practical science activity in the classroom is an essential part of science education in secondary schools. Microbiology, the study of microorganisms (including bacteria, fungi, algae and viruses) relies heavily on practical activity in its teaching. Viruses have significant impact on human activity, encompassing human disease and epidemiology, molecular biology research, treatment of disease and phage-related damage to bacterial starter cultures in food production. These phenomena relate to many aspects of the school teaching specification.

Objectives
To develop a new resource containing background information on viruses including real world case studies, three practical activities about viral infection and molecular techniques for diagnosis of viral infection, and guides for teachers, technicians and students.

Methods
The booklet underwent extensive formative evaluation in preparation for launch, including trials with students and teachers as well as considerable development in a laboratory setting, in and out of the classroom environment.

Conclusions
Subsequently, ‘Viruses: a practical resource for post-16 biology teachers’ was launched (2013) as an 80-page colour educational resource aimed at post-16 biology education. The resource is freely available to Society for General Microbiology members and is about to undertake extensive summative evaluation.