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Extended-Spectrum-Beta-Lactamases, AmpC Beta-Lactamases and Plasmid Mediated Quinolone Resistance in *Klebsiella* spp. from Companion Animals in Italy

Valentina Donati¹, Fabiola Feltrin¹, Rene S. Hendriksen², Christina Aaby Svendsen², Gessica Cordaro¹, Aurora García-Fernández³, Serena Lorenzetti¹, Raniero Lorenzetti¹, Antonio Battisti¹, Alessia Franco¹

1 Istituto Zooprofilattico Sperimentale delle Regioni Lazio e Toscana, Rome, Italy, 2 Technical University of Denmark, National Food Institute (DTU-Food), Kongens Lyngby, Denmark, 3 Istituto Superiore di Sanità, Department of Infectious, Parasitic and Immune-Mediated Diseases, Rome, Italy

**Abstract**

We report the genetic characterization of 15 *Klebsiella pneumoniae* (KP) and 4 isolates of *K. oxytoca* (KO) from clinical cases in dogs and cats and showing extended-spectrum cephalosporin (ESC) resistance. Extended spectrum beta-lactamase (ESBL) and AmpC genes, plasmid-mediated quinolone resistance (PMQR) and co-resistances were investigated. Among KP isolates, ST101 clone was predominant (8/15, 53%), followed by ST15 (4/15, 27%). ST11 and ST340, belonging to Clonal Complex (CC11), were detected in 2012 (3/15, 20%). MLST on KP isolates corresponded well with PFGE results, with 11 different PFGE patterns observed, including two clusters of two (ST340) and four (ST101) indistinguishable isolates, respectively. All isolates harbored at least one ESC or AmpC gene, all carried on transferable plasmids (IncI, IncFII, IncI1, IncN), and 16/19 were positive for PMQR genes (*qnr* family or *aac(6’)-Ib-cr*). The most frequent ESBL was CTX-M-15 (11/19, 58%), detected in all KP ST101, in one KP ST15 and in both KP ST340. *blaCTX-M-15* was carried on IncI plasmids in all but one KP isolate. All KP ST15 isolates harbored different ESC resistance genes and different plasmids, and presented the non-transferable *blaSHV-28* gene, in association with *blaCTX-M-15*, *blaCTX-M-1* (on IncI, or on IncN), *blaOXA-25* (on IncI) or *blaOXA-1* (on IncI). KP isolates were positive for *blaCTX-M-9* gene (on IncHI2), or for the *blaSHV-12* and *blaOXA-1* genes (on IncL/M). They were all positive for *qnr* genes, and one also for the *aac(6’)-Ib-cr* gene. All *Klebsiella* isolates showed multiresistance towards aminoglycosides, sulfonamides, tetracyclines, trimethoprim and amphenicols, mediated by *strA/B, aadA2, aadB, ant (2’)-Ia, aac(6’)-Ib, sul, tet, dfr* and *cat* genes in various combinations. The emergence in pets of multidrug-resistant *Klebsiella* with ESBL, AmpC and PMQR determinants, poses further and serious challenges in companion animal therapy and raise concerns for possible bidirectional transmission between pets and humans, especially at household level.

**Materials and Methods**

**Origin of ESC-resistant Klebsiella**

Between 2006 and 2012, the Istituto Zooprofilattico Sperimentale delle Regioni Lazio e Toscana (IZSLT) investigated samples from 1555 dogs and 429 cats of clinical cases and necropsy specimens with suspicious bacterial infections, submitted by veterinarians practising mainly in central Italy, and some practising in northern Italy. Presumptive positive *Klebsiella* isolates were identified using the API 20E identification system (bioMérieux, Graponne, France). For species-level identification of isolates,

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*E-mail: antonio.battisti@izslt.it*
with phenotypic inconclusive results 16S rDNA sequencing technique was employed, by means of the MicroSeq Full Gene system (Applied Biosystems, USA) as described previously [13].

Genotypic characterization
Multilocus Sequence Typing on KP isolates was performed as previously described [14], and interpreted according to the KP MLST database (www.pasteur.fr/mlst).

In addition, all isolates were genotyped by PFGE using XbaI according to the previously published protocol [15].

Antimicrobial susceptibility testing
Antimicrobial susceptibility testing was performed as minimum inhibitory concentrations (MIC) by micro-broth dilution in 96-well microtitre plates (Trek Diagnostic Systems, Westlake, OH, USA). The following antimicrobials were tested: ampicillin, cefotaxime, cefazidime, ciprofloxacin, chloramphenicol, florfenicol, gentamicin, kanamycin, nalidixic acid, streptomycin, sulfonamides, tetracycline, and trimethoprim. The results were interpreted according to the European Committee on Antibiotic Susceptibility Testing (EUCAST) epidemiological cut-offs (www.eucast.org) and to Clinical Laboratory Standard Institute [16] or EUCAST clinical breakpoints for those drugs for which epidemiological cut-offs have not been made available (kanamycin, chloramphenicol, sulfamethoxazole, trimethoprim). For streptomycin, a cut-off of 16 mg/L was used, according to EUCAST MIC distributions.

Confirmatory test for the detection of ESBLs were performed on isolates resistant to cefotaxime or ceftazidime according to Clinical Laboratory Standard Institute (CLSI) recommendations [16].

Detection of genes encoding beta-lactamase and PMQR
For the confirmed ESBL-producing isolates, the encoding genes belonging to the beta-lactamase and PMQR families were further analyzed for the presence of blaCTX-M [17], blaSHV [18], blaTEM [19], blaOXA [20], bla AmpC families [21], as well as for genes of the qnr family, qep-A, and acc(6')-Ib-cr encoding for PMQR [22], [23], [24], [25], [26]. The isolates were further screened by PCR for genes encoding carbapenemases [27]. Amplicons were sequenced by BigDye Terminator chemistry (Applied Biosystems, Foster City, CA, USA) and migrated with an automated sequencer (ABI Prism 310; Applied Biosystems). Sequence data analysis was performed using CLC DNA workbench software version 5.7.1 (CLC Bio, Aarhus, Denmark) and evaluated against the GenBank nucleotide databases.

Detection of plasmid replicons
Identification of plasmids was performed by PCR-based replication typing as previously described [28], [29], [30], and using the PBRT kit (Diatheva, Fano, Italy).

Plasmid analysis
Plasmid DNA preparations were performed using the NucleoSpin Plasmid/Plasmid (NoLid) kit (Macherey-Nagel, Düren, Deutschland) and used to transform MAX Efficiency DH5α Competent Cells (Invitrogen, Life Technologies, USA). In order to identify the plasmid carrying the ESBLs and AmpC genes, the selection of the transformants was performed on LB agar plates containing 100 mg/ml ampicillin.

Additionally, the isolates were tested according to the manufacturer’s instructions using an array hybridization kit for DNA-based detection of the most common resistance genes, and for the integrase gene (insI) of class 1 integrons of Gram negative bacteria (Alere Technologies GmbH, Jena, Germany) and the results interpreted by the ArrayMate, Alere.

Results

Isolation rates
The samples (n = 1984; dogs and cats) yielded a total 70 (3.53%, 95% CI: 2.72%–4.34%) KP and 23 (1.16%), 95% CI: 0.69%–1.63%) KO among the isolates, respectively. Of these, 15 (21.4%) KP and four (17.4%) KO revealed resistance to ESC and were investigated in this study.

Genetic relatedness
The 15 KP isolates investigated by MLST were assigned to four different Sequence Types (ST): ST11 (n = 1), ST340 (n = 2), ST101 (n = 8), and ST15 (n = 4) (Figure 1). ST11 and its single-locus (tonB) variant (SLV) ST340 (3/15, 20%), both belonging to CC11, were detected in 2012. The separation of the isolates based on MLST corresponded well with PFGE results grouping the same isolates (Figure 1). A total of 11 different PFGE patterns were observed including two clusters of two and four indistinguishable isolates, respectively (Figure 1). The cluster of the two isolates both belonged to ST340 and was related (80% similarity) to a single isolate exhibiting a unique PFGE pattern and belonging to ST11. The other cluster of four indistinguishable isolates was highly related (from 99% to 80% similarity) to additional four isolates within the same PFGE group, all belonging to ST101 (Figure 1). No clustering was observed related to time, animal origin, or infection, but some to the presence of resistance genes (Table 1). No MLST was assigned to the four KO isolates. However, the four isolates revealed three different PFGE patterns of which one was a cluster of two identical isolates (Figure 2). The three patterns seemed not to be related, indicating a similarity of 45% and 55% to the pattern of the two clustering isolates. Interestingly, the two isolates of the same PFGE pattern were both from dogs and isolated within the same year, but it could be the result of a random effect (Figure 2).

Antimicrobial susceptibility testing
All isolates showed microbiological resistance to third-generation cephalosporins, and also clinical resistance, either when the MIC results were interpreted according to clinical breakpoints set by CLSI [16] or by EUCAST (e. g. MIC cefotaxime > 4 mg/L), except for 9KP (MIC 1 mg/L). The phenotype for PMQR was evident (ciprofloxacin MIC 0.25mg/L, nalidixic acid 8 mg/L) in KO isolates only, because in all KP isolates it was masked by concurrent genetic background conferring Mics of 8 and 128 mg/L, respectively.

Moreover, all isolates showed multidrug-resistance towards other classes of antimicrobials, such as aminoglycosides, sulfonamides, tetracyclines, dihydrofolate reductase inhibitors and amphenicols, mediated by strA/B, ada2, adaB, ant (2")-In, acc(6')-Ib, sul, tet, dfr and cat genes in various combinations, as reported in Table 1.

Genes encoding ESBL-, AmpC-, and PMQR
All Klebsiella isolates investigated showed the presence of at least one ESBL or AmpC gene encoding ESC resistance. Additionally, 16 out of 19 isolates harbored a PMQG gene (qnr family or acc(6')-Ib-cr, single or in combination). The most frequent ESC gene harbored by KP isolates was blaCTX-M-15 (n = 11, 58%), detected in all eight ST101, in one ST13 and in both two ST340 isolates, respectively. All four ST15 KP isolates carried the blaSHV-28 gene, single or in combination with the ESC resistance genes blaCTX-M-15.
Based on nemase genes, isolates under study were negative for the presence of carbapenemase genes. Presented the IncR plasmid carrying different ESBL genes such as blaCTX-M-15 and blaCMY-2 and the blaTEM-1 gene. Similarly, the ST340 and ST11 KP isolates were the only ESC-resistant KP harboring genes within the qnr family.

The ST15, ST101 and ST340 KP isolates presented the blaCTX-M-15 gene, which was mostly associated with the blaTEM-1, blaOXA-1 and the blaSHV-1, or blaSHV-11 or blaSHV-28 (Table 1).

Of the four KO isolates, two were positive for blaCTX-M-9 gene, both associated with the PMQR gene qnrS1, with the one the two also positive for the blaSHV-12, blaTEM-1, blaDHA-1 and qnrB4 genes. All isolates under study were negative for the presence of carbapenemase genes.

Plasmids analysis
Plasmids detected from all the strains and classified by the PCR-based replicon typing method, are reported in Table 1.

In our study, the blaCTX-M-15 gene was successfully transferred by transformation from several MLST prototypic strains, demonstrating its plasmid localization. All but one transformant strains were positive for plasmids belonging to the IncR incompatibility group, and in the 18KP transconjugant the blaCTX-M-15 gene was co-transferred with the blaTEM-1 gene. Other ST15 KP strains also presented the IncR plasmid carrying different ESBL genes such as blaSHV-2, and blaCTX-M-1. The 2KP ST15 strain harbored a IncI plasmid carrying the blaCMY-2 gene, while an IncN plasmid carrying both the blaCTX-M-1 and qnrS1 gene was detected in the ST11 16KP strain.

The KO transformants carried the ESBL genes, blaCTX-M-9 and blaSHV-12 in two incompatibility groups, IncHI2 and IncL/M, respectively (Table 1).

Discussion
The paper provides evidence of heterogeneity of determinants conferring ESC resistance in clinical Klebsiella isolates from dogs and cats in Italy. To the best of our knowledge, all the ESC-resistant Klebsiella investigated were from sporadic clinical cases, although two clusters of KO (n = 2) and KP ST101 (n = 4) showed 100% similarity. Apparently, isolates were epidemiologically unrelated, with the exception of 6KP and 1KP (both ST101), with a DNA restriction profile showing 80% similarity, isolated from two cases of necrospies requested by the same veterinary practitioner in 2010 and 2011, respectively.

We also document the novel finding of the co-existence of the ESBL blacmym-2 and the AmpC blacmy-2 gene in one KP ST15 isolate from a cat and the first report of qnrS and qnrA and -Ib-cr PMQR associated with Klebsiella infections in companion animals from Europe. Klebsiella with ESBL phenotype were described in dogs and cats from China [12], although it was mediated in those isolates by the presence of the CTX-M-9 and CTX-M-15 group beta-lactamases, while a CTX-M-15 positive ST15 KP clone was reported from hospital-acquired infections in pets from France [31]. In our study, the ESBL gene blacx-m-15 accounts for the majority of CTX-M genes detected in KP, but it was harbored mainly by the predominant KP ST101 lineage.

Interestingly, all KP ST15 isolates showed the association of the blacx-m-15, blacx-m-15, or blacx-m-2 genes with the blashv-28 gene, a blashv-1 mutant detected for the first time in China in 2002 (GenBank AF538324), and previously reported to encode for an ESBL phenotype [32]. Co-presence of blacx-m-15 and blashv-28 has been reported for the first time in the human KP ST15 epidemic clone by Nielsen et al., in 2011 [33]. Our transformation experiments did not succeed in transferring any of the blashv-28 genes from positive KP isolates.

The multidrug-resistant CTX-M-15-producing KP isolates are an of important concern in the nosocomial infections and the IncFII-type plasmid is the main vehicle of blacx-m-15 transmission in human isolates [34]. However, in our study, this ESBL in KP from pets was mostly carried by IncR plasmids. The association of blacx-m-15 -IncR replicons in KP was documented for the first time by Coelho et al. in 2010 [35], in human clinical isolates, and also reported in the KP clone causing hospital-acquired infections in pets in France [31], and in Spain, associated to qnrB4, bladha-1 and armA genes. [36]

Interestingly, the 17KP transformant only presented the blacx-m-15 gene and the qnrS1 gene both located in a IncFII plasmid. In our study, almost all the blacx-m-15 – positive isolates were also positive for IncFII replicons: although they have been specifically described in KP [30], these plasmids were never transferred in our experiments. Interestingly, in 2012 we have reported for the first

Figure 1. Dendrogram showing the genotypic relatedness of ESC-resistant Klebsiella pneumoniae (KP) isolates from dogs and cats based on XbaI-PFGE fingerprints, and comparison with Multilocus Sequence Typing classification.
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<table>
<thead>
<tr>
<th>Sequence Type</th>
<th>Key</th>
<th>Antimicrobial Resistance profile</th>
<th>Plasmid</th>
<th>ESBL and AmpC genes</th>
<th>PMQR genes</th>
<th>Other resistance and Integron genes</th>
</tr>
</thead>
<tbody>
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<td>9KP</td>
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<td>13KP</td>
<td>AMP, CTX, CFT, NAL, CIP, STR, KAN, GEN, CLO, SULFA, TRI, TET</td>
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<td></td>
<td>aadA2; aadA4; ada17; ada18; sul1; tet(A); intI1</td>
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<td>10KP</td>
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<td>aac&lt;sub&gt;6’&lt;/sub&gt;-Ib-cr; qnrB; qnrS1</td>
<td>aac&lt;sub&gt;6’&lt;/sub&gt;-Ib; ada14; dfrA17; sul2; intI1</td>
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<tr>
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<td>3KP</td>
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<tr>
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<td>1KP</td>
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<td>IncH2; IncR; IncFIIk, IncFII</td>
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<td>aadA2; ant26a; sul; intI1</td>
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</tbody>
</table>

Legend:
NA: Not Applicable; AMP = Ampicillin; CFT = Ceftazidime; CIP = Ciprofloxacin; CLO = Chlomphenicol; CTX = Cefotaxime; GEN = Gentamicin; KAN = Kanamycin; NAL = Nalidixic Acid; STR = Streptomycin; SULFA = Sulfamethoxazole; TET = Tetracycline; TRI = Trimethoprim.

Note: When underscored, plasmids and their content of beta-lactamase and PMQR genes were detected in transformant strains.
time in clinical cases of pets from Italy the clone KP ST11 and its SLV ST340, harboring ESC and qnrS-PMQR resistance. Among these CC11 isolates, the ST11 harbored IncN plasmid, which has been frequently involved in the transmission of the $bla_{CTX-M-1}$ gene, a feature suggesting an animal reservoir for this ESBL, since this Inc plasmid types have been demonstrated to be highly prevalent in zoonotic enterobacterial pathogens [29]. The same animal origin reservoir is proposed for the IncH1 plasmids harboring the $bla_{CMY-2}$ gene found in E. coli avian commensal strains [34].

It is noteworthy that ST11 and ST340 carried transferable ESBL resistance but not resistance to carbapenems. ST(CC)11 and ST15 and ST101 are among human epidemic clones, carrying both ESBLs and carbapenemases, which have been increasingly detected worldwide, in Europe and in Italy in the last years [36], [37], [38], [39], [40].

These infections are worrisome, since the antimicrobial treatment options for these multidrug-resistant strains are very limited. In Italy, during the last years the rapid emergence of the ESBL resistance but not resistance to carbapenems. ST(CC)11 and ST15, and predominantly to a single Sequence Type ST258, has become a serious problem in health-care settings [41], [42], [43].

As for CTX-M and SHV-12 ESBLs in Italy, a high occurrence in KP isolated from humans has been demonstrated, being the ST15, ST37, ST147 and ST273 the prevalent clones [44], [45], [46], [47].

In two KO isolates, the ESBL-encoding $bla_{SHV-12}$ gene co-existed with the AmpC gene $bla_{SHV-1}$ in accordance with phenotype of resistance to cephalotaxine and cefoxitin observed in the ESBL phenotypic confirmatory test. These two isolates also carried the PMQR gene $qnrB1$. In these two KO transformants the IncL/M plasmid harbored both the $bla_{SHV-12}$ and the $bla_{SHV-1}$ genes but not the $qnrB1$ gene. To our knowledge, this feature has never been described before.

The other KO presented the $bla_{CTX-M-9}$ gene, located in an IncHI2 plasmid as described worldwide but associated to a $qnrA$ gene, a feature previously described in Spain in E. coli and KP of human origin [48], [49] and in KO in clinical specimens from Japan [50].

Similarly to what has been observed in other human and canine KP isolates [34], [12], the association of $bla_{CTX-M}$ genes with the the $aac(6’)-Ib-cr$ encoding an aminoglycoside acetyl transferase determining PMQR, was demonstrated in all ST101 and ST340 isolates, but only in one out of four ST15, but these PMQR genes were not located in the same plasmid in our strains. Conversely, PMQR encoded by different $qnr$ genes of the $qnrA$ or $qnrB$ groups were observed in all the KO isolates studied (Table 1). In the case of the $qnrA1$ gene, the two KO isolates also harbored the ESBL $bla_{CTX-M-9}$ gene, a feature reported previously in association with $bla_{ST-1}$ and IncHI2 plasmids in KO of human origin [51].

Multidrug-resistance in the ESC resistant and PMQR isolates studied is of further concern from a therapeutic perspective, for a possible impact on clinical outcome of affected animals. In many isolates, the demonstration of the integrase intI1, accounts for the presence of resistance gene cassettes with $aad$, $ant$, $cat$, $dfr$, genes, associated with Class I integrons, similarly to what has been described in KP of human origin [52], [53]. As for streptomycin resistance, MIC>16 mg/L correlated in 100% isolates with the presence of $strA/B$ genes.

Fortunately, the absence of carbapenemases offers so far a better scenario for antimicrobial therapy in companion animals, although a possible circulation, within a short time, of these carbapenemase-producing epidemic strains, is of concern also in veterinary medicine.

In conclusion, monitoring and characterization of multidrug-resistant Klebsiella in companion animals by means of phenotypic and molecular methods proved to be useful for providing a picture of mechanisms of resistance that may further spread clonally or by horizontal gene transfer, at regional or even at international level. Sharing this kind of information appears essential for building awareness in companion animal therapy, also in view of preventing and controlling the spread of multidrug-resistant strains in veterinary hospital settings. Indeed, the bi-directional exchange between owners and pets of Klebsiella carrying resistance to critically important antimicrobials for human health, raise some concerns also for the possibility of a spill back to humans, especially at household level.

The emergence of the PMQR and, above all, the emergence of concurrent transferable cephamycin, oxymino-cephalosporin, and beta-lactamase inhibitor resistance in multidrug-resistant Klebsiella isolates in pets, a recent issue even in human therapy [54], may pose in the next future more and serious therapeutic challenges also in bacterial infections of companion animals.

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**Author Contributions**

Conceived and designed the experiments: AF AB RSH FF AGF. Performed the experiments: VD SL FF GC RL CAS AGF. Contributed reagents/materials/analysis tools: AB. Analyzed the data: FD FF AB RL RSH CAS AGF. Contributed: AB FF RSH AGF. Wrote the paper: AF AB RSH AGF.
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