Colonic transit time relates to bacterial metabolism and mucosal turnover in the human gut

Roager, Henrik Munch; Hansen, Lea Benedicte Skov; Bahl, Martin Iain; Frandsen, Henrik Lauritz; Carvalho, Vera; Gøbel, Rikke J.; Dalgaard, Marlene Danner; Sparholt, Morten; Vestergaard, Henrik; Hansen, Torben; Sicheritz-Ponten, Thomas; Nielsen, Henrik Bjørn; Pedersen, Oluf; Lauritzen, Lotte; Kristensen, Mette; Gupta, Ramneek; Licht, Tine Rask

Publication date:
2016

Document Version
Peer reviewed version

Citation (APA):
Colonic transit time relates to bacterial metabolism and mucosal turnover in the human gut

Henrik M. Roager\textsuperscript{1}, Lea B. S. Hansen\textsuperscript{2}, Martin I. Bahl\textsuperscript{1}, Henrik L. Frandsen\textsuperscript{1}, Vera Carvalho\textsuperscript{1}, Rikke J. Gøbel\textsuperscript{3}, Marlene D. Dalgaard\textsuperscript{2}, Morten H. Sparholt\textsuperscript{4}, Henrik Vestergaard\textsuperscript{1}, Torben Hansen\textsuperscript{3}, Thomas S. Pontén\textsuperscript{2}, H. Bjørn Nielsen\textsuperscript{2}, Oluf Pedersen\textsuperscript{3}, Lotte Lauritzen\textsuperscript{5}, Mette Kristensen\textsuperscript{5}, Ramneek Gupta\textsuperscript{2}, Tine R. Licht\textsuperscript{1}.

\textsuperscript{1}National Food Institute, Technical University of Denmark, Denmark
\textsuperscript{2}Department of Systems Biology, Technical University of Denmark, Denmark
\textsuperscript{3}The Novo Nordisk Foundation Center for Basic Metabolic Research, Section of Metabolic Genetics, University of Copenhagen, Denmark
\textsuperscript{4}Department of Radiology Bispebjerg and Frederiksberg Hospitals
\textsuperscript{5}Department of Nutrition, Exercise and Sport, University of Copenhagen, Denmark

Little is known about how colonic transit time relates to human colonic metabolism, and its importance for host health, although stool consistency, a proxy for colonic transit time, has recently been negatively associated with gut microbial richness. To address the relationships between colonic transit time and the gut microbial composition and metabolism, we assessed the colonic transit time of 98 subjects using radiopaque markers, and profiled their gut microbiota by 16S rRNA gene sequencing and their urine metabolome by ultra performance liquid chromatography mass spectrometry. Based on correlation analyses, we show that colonic transit time is associated with overall gut microbial composition, diversity and metabolism. A relatively prolonged colonic transit time associates with high microbial species richness and a shift in colonic metabolism from carbohydrate fermentation to protein catabolism as reflected by microbial metabolites in urine. This results in a number of potentially deleterious protein-derived metabolites. Additionally, longer colonic transit time correlates with metabolites likely reflecting reduced renewal of the colonic mucosa. Together, this suggests that a high gut microbial richness does not \textit{per se} imply a healthy gut microbiota, and contributes to the understanding of the pathophysiology of diseases where increased transit time is a risk factor. Finally, our findings highlight the colonic transit time as an important physiological variable, which should be considered in gut microbiota and metabolomics studies.