

O.438. Prevalence and control of Shiga toxin-producing *E. coli*: from diversity in dairy cattle to phage therapy

Carla Dias¹, Andressa Ballem², Ana Pinto³, Gonçalo Almeida², Pablo Fuciños⁴, Carina Almeida², Maria José Saavedra^{3,5}, Filipe Silva⁵, José Almeida⁵, Jorge Azevedo⁵, Maria José Gomes⁵, Hugo Oliveira¹

¹Centre of Biological Engineering (CEB), University of Minho, Campus de Gualtar, Braga, Portugal

²National Institute for Agrarian and Veterinary Research (INIAV), Vila do Conde, Portugal.

³Veterinary Science Department, University of Trás-os-Montes and Alto Douro (UTAD), Vila Real, Portugal

⁴International Nanotechnology Laboratory (INL), Braga, Portugal

⁵Animal Science Department, and Animal Science Research Center (CECAV), University of Trás-os-Montes and Alto Douro (UTAD), Vila Real, Portugal

Shiga toxin-producing *E. coli* (STEC) strains are important foodborne pathogens worldwide, transmitted from ruminant to humans through contaminated food. Their control is still a challenge as most *E. coli* in nature are commensal and, thus, controlling strategies should target only pathogenic strains/serotypes. Bacteriophages (bacterial viruses) can cope with this challenge by allowing a tailored intervention. We performed an epidemiological study of STEC at 21 milk farms across the Northern region of Portugal and evaluated the potential of bacteriophage therapy to control the well-known O157 STEC serotype. From 409 dairy cattle analyzed, STEC strains were more prevalent in heifers (45 %) than in lactating cows (16 %). STEC isolates with several *stx1* and *stx2* subtypes were identified and they belonged to 73 different O:H serotypes. Regarding bacteriophage therapy evaluation, an O157-specific phage (CBA120), was tested *in vitro* and *in vivo*. The bacteriophage reduced STEC in contaminated ruminant fluids of rumen and intestine (>4 logs) as well as STEC biofilms adhered with intestinal mucosa (>2 logs). Moreover, bacteriophage treatments significantly reduced *E. coli* O157:H7 numbers (1 log) in artificially contaminated sheep, comparatively with the mocked-treated group. Overall, results suggest the potential use of bacteriophages to control STEC *in vivo*.

Acknowledgement

This study was supported by project PhageSTEC (POCI-01-0145-FEDER-029628) funded by FEDER through COMPETE2020 (Programa Operacional Competitividade e Internacionalização) and by National Funds through FCT (Fundação para a Ciência e a Tecnologia).