

Abstracts book



9th Symposium on
Antimicrobial
Resistance in
Animals and the
Environment

2023JULY 3-5
TOURS









O27

High occurrence of carbapenemase- and extended-spectrum-β-lactamaseproducing *Escherichia coli* and *Klebsiella pneumoniae* from migratory birds (*Ciconia ciconia*) with detection of high-risk clones

<u>Sandra MARTÍNEZ-ÁLVAREZ</u>¹ Pierre CHÂTRE², Úrsula HÖFLE³, Carla Andrea ALONSO⁴, Pauline FRANÇOIS², Teresa CARDONA-CABRERA², Myriam ZARAZAGA¹, Jean-Yves MADEC², Marisa HAENNI², Carmen TORRES¹

¹Area of Biochemistry and Molecular Biology, OneHealth-UR Research Group, University of La Rioja - Logroño (Spain)

²ANSES – Université de Lyon, Unité Antibiorésitance et Virulence Bactériennes - Lyon (France)

³Health and Biotechnology (SaBio) Research Group, Institute for Game and

Wildlife Research IREC (CSIC-UCLM) - Ciudad Real (Spain)

⁴Hospital San Pedro, Logroño (Spain)

Background: The presence of Enterobacterales producing carbapenemases (CP) and extended-spectrum β -lactamases (ESBL) is closely monitored in humans and animals, but the potential of migratory birds as carriers of resistance genes remains poorly understood. The aim of the study was to detect and characterize CP- and ESBL-producing *E. coli* (EC) and *K. pneumoniae* (KP) obtained from storks feeding on two landfills in Spain.

Methods: ESBL and CP-producing EC/KP were isolated from 211 stork faecal samples using chromogenic culture media, and collected isolates were sequenced using NovaSeq6000 (Illumina) and MinION (Oxford Nanopore) technologies. Resistome, virulome, sequence types (ST) and replicon profiles were determined using bioinformatic tools. Localization of ESBL/CP genes was performed using Southern blots on S1-PFGE gels.

Results: ESBL-EC/KP were detected in 71 samples (33.6%; 28.4%-EC and 5.2%-KP), while 28 samples (13.3%) contained CP-producing EC/KP (11.8%-EC and 1.4%-KP). Different sequence types (ST) (EC, n=33; KP, n=3) were identified, including high-risk clones associated with humans (EC: ST131, ST58 or ST69; KP: ST307), potential high-risk clones (EC: ST10 and ST48) and more ubiquitous clones (EC: ST46, ST155, ST117, ST617). A wide range of ESBL/pAmpC-conferring genes (blactx-m-1/blactx-m-1/blactx-m-15/blactx-m-27/blactx-m-32/blactx-m-55/blactx-m-65/blashv-12/blacmy-2/bladha-1) was identified, as well as a large variety of CP-genes (blakpc-2, blakpc-3, blandm-1, blandm-7, blaoxa-48, blavim-1 and blages-7), including in some cases a combination of up to three types of CP (blakpc-2, blandm-7 and blavim-1) and/or with co-detection of ESBL/pAmpC genes. Likewise, a variety of ESBL/pAmpC- and CP-carrying plasmids were identified, such as IncY (n=30), IncF (n=60), IncX3 (n=25) IncN (n=19) or IncL (n=8). The genetic characterization of these plasmids is ongoing.