

First report of *bla*_{OXA-181}-carrying IncX3 plasmids in multidrug-resistant *Enterobacter hormaechei* and *Serratia nevei* recovered from canine and feline opportunistic infections

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ABSTRACT Whole-genome sequence analysis of six *Enterobacter hormaechei* and two *Serratia nevei* strains, using a hybrid assembly of Illumina and Oxford Nanopore Technologies sequencing, revealed the presence of the epidemic *bla*_{OXA-181}-carrying IncX3 plasmids co-harboring *qnrS1* and Δ *ere(A)* genes, as well as multiple multidrug resistance (MDR) plasmids disseminating in all strains, originated from dogs and cats in Thailand. The subspecies and sequence types (ST) of the *E. hormaechei* strains recovered from canine and feline opportunistic infections included *E. hormaechei* subsp. *xiangfangensis* ST171 ($n = 3$), ST121 ($n = 1$), and ST182 ($n = 1$), as well as *E. hormaechei* subsp. *steigerwaltii* ST65 ($n = 1$). Five of the six *E. hormaechei* strains harbored an identical 51,479-bp *bla*_{OXA-181}-carrying IncX3 plasmid. However, the *bla*_{OXA-181} plasmid (pCUVET22-969.1) of the *E. hormaechei* strain CUVET22-969 presented a variation due to the insertion of *ISKpn74* and *ISSbo1* into the *virB* region. Additionally, the *bla*_{OXA-181} plasmids of *S. nevei* strains were nearly identical to the others at the nucleotide level, with *ISEc11* inserted upstream of the *qnrS1* gene. The *E. hormaechei* and *S. nevei* lineages from canine and feline origins might acquire the epidemic *bla*_{OXA-181}-carrying IncX3 and MDR plasmids, which are shared among Enterobacterales, contributing to the development of resistance. These findings suggest the spillover of significant OXA-181-encoding plasmids to these bacteria, causing severe opportunistic infections in dogs and cats in Thailand. Surveillance and effective hygienic practice, especially in hospitalized animals and veterinary hospitals, should be urgently implemented to prevent the spread of these plasmids in healthcare settings and communities.

IMPORTANCE *bla*_{OXA-181} is a significant carbapenemase-encoding gene, usually associated with an epidemic IncX3 plasmid found in Enterobacterales worldwide. In this article, we revealed six carbapenemase-producing (CP) *Enterobacter hormaechei* and two CP *Serratia nevei* strains harboring *bla*_{OXA-181}-carrying IncX3 and multidrug resistance plasmids recovered from dogs and cats in Thailand. The carriage of these plasmids can promote extensively drug-resistant properties, limiting antimicrobial treatment options in veterinary medicine. Since *E. hormaechei* and *S. nevei* harboring *bla*_{OXA-181}-carrying IncX3 plasmids have not been previously reported in dogs and cats, our findings provide the first evidence of dissemination of the epidemic plasmids in these bacterial species isolated from animal origins. Pets in communities can serve as reservoirs of significant antimicrobial resistance determinants. This situation places a burden on antimicrobial treatment in small animal practice and poses a public health threat.

KEYWORDS *bla*_{OXA-181}, carbapenemase, cats, dogs, Enterobacterales, MDR, WGS

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Carbapenem-resistant Enterobacterales are Gram-negative bacteria of major public health concern, causing a wide range of opportunistic infections in hospitals. Alongside *Escherichia coli* and *Klebsiella pneumoniae*, the *Enterobacter cloacae* complex (ECC) and *Serratia* species are increasingly recognized as causing various opportunistic diseases in hospitalized patients (1, 2) and are occasionally associated with opportunistic infections in dogs and cats (3, 4). In addition to their intrinsic resistance, they can become multidrug-resistant (MDR) by acquiring antimicrobial resistance genes (ARGs) on mobile genetic elements, limiting therapeutic options (5–7). Carbapenems have been off-label prescribed in the treatment of small animals in veterinary medicine (8). In recent years, several publications have reported the dissemination of carbapenemase-encoding genes associated with specific plasmid families in ECC and *Serratia* spp. in humans, companion animals, and veterinary settings, but so far, none of the veterinary strains were found to carry *bla*_{OXA-181} (7, 9, 10).

Six carbapenemase-producing (CP) ECC and two CP *Serratia marcescens* strains, identified using matrix-assisted laser desorption ionization time-of-flight mass spectrometry (Bruker Daltonics GmbH, Germany), were isolated from canine and feline diagnostic samples sent to the Veterinary Diagnostic Laboratory, Faculty of Veterinary Science, Chulalongkorn University, Bangkok, Thailand, between 2018 and 2022. Whole-genome-based identification using BLAST-based average nucleotide identity through the online available service JSpeciesWS (<https://jspecies.ribohost.com/jspeciesws/>) reclassified the ECC as *Enterobacter hormaechei* subsp. *xiangfangensis* ($n = 5$) and *E. hormaechei* subsp. *steigerwaltii* ($n = 1$), and the *S. marcescens* as *Serratia nevei* (Table S1). The two CP *S. nevei* (CP-*Sn*) and three CP *E. hormaechei* (CP-*Eh*) strains were recovered from dogs, while the other three CP-*Eh* strains were isolated from cats. Carbapenem resistance and the presence of *bla*_{OXA-48-like} gene were first observed with imipenem minimum inhibitory concentration ≥ 16 $\mu\text{g}/\text{mL}$ in diagnostic results using Vitek 2 automated system (bioMérieux, France) and by multiplex PCRs, respectively (11, 12).

Antimicrobial susceptibility testing was further conducted by broth microdilution assay using Sensititre ASSECAF/ASSECB plates (Thermo Fisher Scientific, USA). Resistance was interpreted following the interpretive criteria of the Clinical and Laboratory Standards Institute, except for tigecycline, which was referred to the criteria of *E. coli* from the European Committee on Antimicrobial Susceptibility Testing (13–15). The two CP-*Sn* isolates exhibited resistance to all drugs tested. Antimicrobial resistance phenotypes of the six CP-*Eh* strains were variable (Table S2). However, all CP-*Eh* isolates exhibited resistance to ampicillin, meropenem, nalidixic acid, ciprofloxacin, and azithromycin but were still susceptible to colistin and tigecycline.

Whole-genome sequences of all strains were obtained using the hybrid assembly of sequence reads from both Illumina Nextseq 550 and Oxford Nanopore Technologies. Raw reads were qualified using the FastQC program v0.11.9 (<https://www.bioinformatics.babraham.ac.uk/projects/fastqc/>). Illumina reads and sequence adaptors were trimmed and removed using Trimmomatic v0.39. Assembly of the reads was done using Unicycler v0.4.8 generating contigs of circular complete genomes and plasmids in all strains (Table S3). The quality of the contigs was assessed using QUAST v5.0.2 (<https://github.com/ablab/quast>) before submission to the NCBI Prokaryotic Genome Annotation Pipeline for gene annotation (Table S3). Whole-genome sequence analysis revealed the presence of *bla*_{OXA-181} gene and additional ARGs (Table S1) in all strains by using NCBI AMRFinderPlus v3.11.17, Resfinder v4.1, and CARD v3.2.7. Plasmid incompatibility complex (Inc) groups and insertion sequence (IS) elements were identified by PlasmidFinder v2.1 and ISFinder tools, respectively. Three of the CP-*Eh* strains belong to the ST171 high-risk clone (16), while the others were ST65, ST121, and ST182. The core genome of CP-*Eh* was extracted using the Roary pipeline v3.13.0 before performing single-nucleotide polymorphism (SNP) analysis of the core gene alignment to obtain core genome single-nucleotide polymorphism (cgSNP) using SNP-sites v2.5.1. cgSNP-based phylogeny obtained using RaxML v8.2.11 revealed that, within the ST171 clades, CUVET21-1190 and CUVET21-1726 had identical core genes and differed from

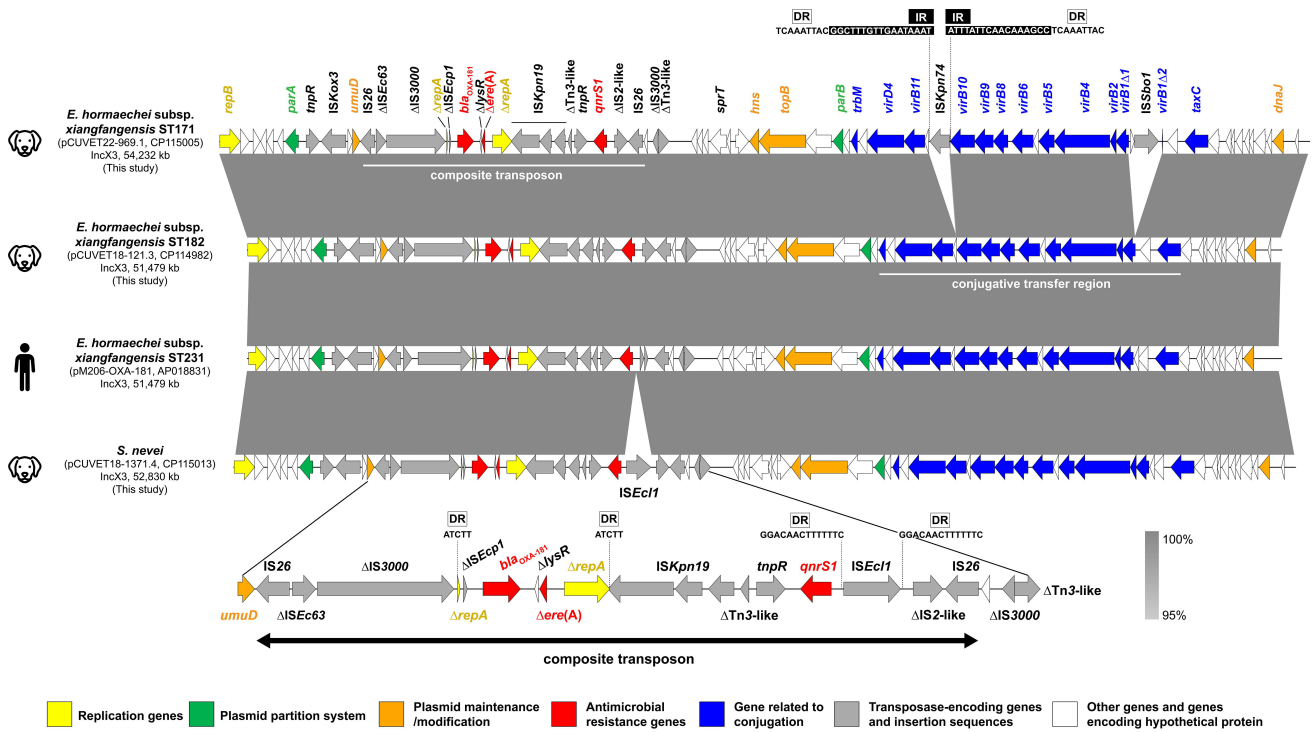


FIG 2 Sequence comparison of the *bla*_{OXA-181}-carrying IncX3 plasmids in *Enterobacter hormaechei* and *Serratia nevei* isolated from dogs in Thailand with pM206-OXA181 (GenBank accession number AP018831) harbored by *E. hormaechei* ST231 isolated from a patient in Myanmar. The same sequence structure of IncX3 plasmid in *E. hormaechei* ST182 (pCUVET18-121.3) was found in pCUVET19-891.3, pCUVET21-1190.1, pCUVET21-1726.1, and pCUVET22-793.2, all exhibiting 100% nucleotide identity. The IncX3 plasmids (pCUVET18-1371.4 and pCUVET18-1784.4) in two *S. nevei* strains were identical and had an additional insertion of *ISEc1* upstream of *qnrS1*, flanked by two 14-bp DRs. The genetic environment of *bla*_{OXA-181} flanked by two copies of IS26 composite transposon is demonstrated at the bottom. The 5-bp AT-rich duplicate sequences (ATCTT) of the extremities of *ΔrepA* suggested insertion and subsequent recombination between two copies of *ISEcp1* (20, 22). The arrows indicate the orientation and length of genes. The color of each arrow represents the gene type or function as given in the legend below. The dark gray shading indicates 100% sequence similarity. The symbol Δ indicates truncation.

by *ΔlysR* and *Δere(A)*, which were flanked by fragments of *repA*. A 5-bp AT-rich sequence repeat (ATCTT) was present at the extremities of *ΔrepA*, suggesting the insertion and subsequent recombination between two copies of *ISEcp1* of the ColKP3 replicon into this region (20, 22). The *qnrS1* was separately located downstream of *ΔrepA* at the right extremity of the transposon. Additionally, these plasmids were mobilizable, as they had conjugative regions consisting of a relaxase gene (*taxC*), a gene encoding type IV coupling protein (*virD4*), and *virB* gene cluster of type IV secretion system (Fig. 2) (18). The broad host range ability of IncX3 plasmids has contributed to the widespread dissemination of carbapenemase-encoding genes, including *bla*_{OXA-181} and *bla*_{NDM-5}, in Enterobacterales (23). Identical or highly similar IncX3 plasmids have also been identified in *E. coli*, *K. pneumoniae*, and *S. marcescens* (20, 21, 24, 25). The detection of this resistance plasmid in *Enterobacter* and *Serratia* is relatively rare, as these genera causing opportunistic diseases are less commonly found when compared to *E. coli* and *K. pneumoniae* (2, 17).

The canine and feline *E. hormaechei* and *S. nevei* strains were MDR and carried multiple ARGs on both the chromosome and plasmids. Two additional MDR plasmids were detected in CP-*Eh* ST182 strain, including IncFIB and IncR plasmids, which contained 13 and four ARGs, respectively (Table S1). The CP-*Eh* ST65 strain carried a 283,562-bp IncHI2/2A plasmid (pCUVET22-793.1) containing *mcr-9.1*, but the strain was susceptible to colistin. This plasmid was similar to pC45_001 (GenBank accession number CP042552) and pCM18-216 (GenBank accession number CP050312) of *E. hormaechei* ST133 and ST110, respectively, which were isolated from a patient and

hospital environment in Australia (Fig. S2). This *mcr-9.1*-carrying IncHI2/2A plasmid has also been found in other Enterobacterales (7, 26, 27). However, the expression of *mcr-9.1* in this pCUVET22-793.1 plasmid likely did not occur due to the absence of *qseB*–*qseC* two-component regulatory genes (28). Additionally, two CP-*Sn* strains harbored identical IncA/C2 and IncFII MDR plasmids, each containing eight and six ARGs, respectively (Table S1).

In summary, our study represents the first detection and description of *E. hormaechei* and *S. nevei* carrying the epidemic *bla*_{OXA-181}-bearing IncX3 plasmids originating from animals. This also indicates that animals and animal environment contribute to the dissemination of MDR and carbapenemase-encoding plasmids among different species of opportunistic Enterobacterales. Given their proximity to humans and communities, monitoring the spread of clonal lineages and plasmids, especially in hospitalized animals and veterinary hospitals, should be further implemented.

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DATA AVAILABILITY

The genome sequences of six *E. hormaechei* strains and two *S. nevei* strains are deposited in the NCBI database under BioProject accession number [PRJNA912905](https://www.ncbi.nlm.nih.gov/bioproject/PRJNA912905).

ADDITIONAL FILES

The following material is available [online](#).

Supplemental Material

Supplemental material (Spectrum03589-23-S0001.pdf). Tables S1 to S3; Fig. S1 and S2.

Open Peer Review

PEER REVIEW HISTORY (review-history.pdf). An accounting of the reviewer comments and feedback.

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