Emergence of a Novel Plasmid-Mediated Tigecycline Resistance Gene Cluster, \textit{tmexCD4-toprJ4}, in \textit{Klebsiella quasipneumoniae} and \textit{Enterobacter roggenkampii}

Xun Gao, Chengzhen Wang, Luchao Lv, Xiaotong He, Zhongpeng Cai, Wanyun He, Tong Li, Jian-Hua Liu

\textsuperscript{a}College of Veterinary Medicine, Key Laboratory of Zoonosis of Ministry of Agricultural and Rural Affairs, Guangdong Provincial Key Laboratory of Veterinary Pharmaceutics Development and Safety Evaluation, South China Agricultural University, National Risk Assessment Laboratory for Antimicrobial Resistant of Microorganisms in Animals, Guangzhou, China

\textsuperscript{b}Guangdong Laboratory for Lingnan Modern Agriculture, Guangzhou, China

Xun Gao and Chengzhen Wang contributed equally to this work. Author order was determined by contribution.

\textbf{ABSTRACT} The occurrence of transferable tigecycline resistance determinants, \textit{tmexCD1-toprJ1}, \textit{tmexCD2-toprJ2}, \textit{tmexCD3-toprJ1b}, and multiple \textit{tet(A)} and \textit{tet(X)} variants, presents an unprecedented challenge to clinical therapeutic options. \textit{tmexCD-toprJ} gene clusters can mediate multidrug resistance and have been detected in a variety of bacteria. Here, we characterized the fourth \textit{tmexCD-toprJ} gene cluster, \textit{tmexCD4-toprJ4}, identified on untypeable plasmids of \textit{Klebsiella quasipneumoniae} and \textit{Enterobacter roggenkampii} isolated from chicken meat and environmental samples from farm markets, respectively. \textit{TMexCD4-ToprJ4} was closely related (92 to 99% amino acid identity) to \textit{TMexCD1-ToprJ1}, \textit{TMexCD2-ToprJ2}, and \textit{TMexCD3-ToprJ1}. Phylogenetic analysis revealed that \textit{tmexCD4-toprJ4} was not in the same branch as the other three variants. Expression of \textit{tmexCD4-toprJ4} increased tigecycline efflux in \textit{Escherichia coli} and resulted in a 4- to 8-fold increase in MICs of tigecycline in \textit{E. coli} and \textit{Klebsiella pneumoniae}. Moreover, \textit{tmexCD4-toprJ4} can act synergistically with its upstream gene \textit{tet(A)} to reduce the susceptibility of \textit{E. coli} and \textit{K. pneumoniae} strains to tigecycline. The \textit{tmexCD4-toprJ4}-containing plasmid is a novel plasmid type and can be transferred to \textit{E. coli} and \textit{K. pneumoniae} only via electrottransformation. The increasing emergence of plasmid-mediated tigecycline resistance gene clusters suggests that the spread of \textit{tmexCD-toprJ} gene clusters requires widespread attention.

\textbf{IMPORTANCE} The plasmid-mediated tigecycline resistance gene cluster \textit{tmexCD1-toprJ1} and other variants have been detected in a variety of strains from multiple sources, including human-derived strains. In addition to tigecycline, these \textit{tmexCD-toprJ} gene clusters reduce susceptibility of the host strain to many other antimicrobials. Here, we identified \textit{tmexCD4-toprJ4} in \textit{K. quasipneumoniae} and \textit{E. roggenkampii}, suggesting that this gene cluster is already present in the human-associated environment and the risk of transmission to humans is increased. Monitoring tigecycline-resistant Gram-negative bacteria is essential for understanding and addressing the spread of this gene cluster in agriculture and health care.

\textbf{KEYWORDS} RND efflux pump gene cluster, tigecycline, plasmid, Enterobacteriaceae

Tigecycline is a last-resort antibiotic used to treat severe clinical infections caused by multidrug-resistant (MDR) bacteria (1). However, the widespread use of tigecycline has led to the evolution of resistance, which reduces therapeutic effectiveness. Notably, the recent emergence of plasmid-mediated \textit{tet(X)} variants (2) and transferable resistance-nodulation-division (RND) family efflux pump gene clusters (3) that confer resistance to tigecycline has increased the challenges of controlling tigecycline resistance. Since the emergence of the RND family efflux pump resistance gene cluster \textit{tmexCD1-toprJ1} in \textit{Klebsiella pneumoniae},
two additional tmexCD-toprJ-like gene clusters, tmexCD2-toprJ2 (4) and tmexCD3-toprJ1b (5), have also been identified.

tmexCD1-toprJ1, which confers multidrug resistance, has been detected in K. pneumoniae (1), Klebsiella quasipneumoniae (6), Raoultella planticola (1), Klebsiella oxytoca (1), and Enterobacter cloacae and might be transferred by site-specific integranes (1) or IS26 (6). tmexCD1-toprJ1 has been detected in humans, animals, food, and sewage. tmexCD2-toprJ2 has been found in K. pneumoniae, Citrobacter freundii, Raoultella omithinolytica, Klebsiella variicola, K. quasipneumoniae, and Klebsiella michiganensis (7) obtained from environmental or clinical samples (4). tmexCD2-toprJ2 may be mobilized by the XerD-like recombinase system (4). Subsequently, tmexCD3-toprJ1b was found in Pseudomonas aeruginosa and Proteus mirabilis isolated from chicken fecal samples and is located in an integrative and conjugative element (ICE) (5). The three gene clusters could mediate the 4- to 16-fold decrease in tigecycline susceptibility of the host bacteria. In this study, we report a novel plasmid-mediated tmexCD-toprJ-like gene cluster, tmexCD4-toprJ4, in K. quasipneumoniae obtained from chicken meat and Enterobacter roggenkampii obtained from a farm market environment in China.

From May 2019 to July 2021, 109 chicken meat samples and 128 environmental samples were collected from farm markets in Guangzhou, China. All samples were selected on MacConkey agar plates supplemented with 4 mg/L tigecycline. Colonies with different morphologies were selected and screened for tmexCD1-toprJ1 by PCR using specific primers (see Table S1 in the supplemental material). Nine (8.2%) tmexCD1-toprJ1-positive strains and two tmexCD1-toprJ1-like strains (GLW9C22 and GD21SC1505) were identified. GLW9C22 was isolated from a chicken meat sample and showed resistance to florfenicol, apramycin, and several tetracyclines including tigecycline (32 μg/mL) (Table S2). GD21SC1505, which was recovered from an environmental sample, showed resistance to florfenicol, colistin, and tigecycline (16 μg/mL). Meanwhile, the MIC of tigecycline decreased to 0.5 μg/mL for both strains in the presence of the efflux pump inhibitor 1-[(1-naphthylmethyl)-piperazine (NMP) (Table 1).

The complete genomic DNA data of GLW9C22 and GD21SC1505 were generated by the combination of Illumina HiSeq platforms and Nanopore MinION, followed by assembly using Unicycler version 0.4.3.8. The results revealed that GLW9C22 belonged to K. quasipneumoniae and harbored a 5,129,993-bp chromosome and three plasmids. Moreover, GLW9C22 carried several known acquired antibiotic resistance genes, including aac(3)-IV, aph(4)-Ia, aph (3’)-Ia, flor, aadA2, aph(3’)-la, and tet(A). GD21SC1505 belonged to E. roggenkampii and harbored a 4,780,540-bp chromosome and five plasmids (Table 1). GD21SC1505 harbored the resistance genes blomir, qnsFI, tet(A), and florR. Although GD21SC1505 showed resistance to colistin, this strain did not carry colistin resistance-related genes, and no mutations in related genes were detected on the chromosome; therefore, the reason for resistance needs to be further investigated. K. quasipneumoniae branched from K. pneumoniae as a new bacterial species in 2014 (8). This species can persist in hospitalized patients and in the hospital environment for a long time and can spread between patients and sink drains. Moreover, K. quasipneumoniae has acquired multiple clinically important resistance genes, including carbapenem and colistin resistance genes (9). E. roggenkampii is a type of E. cloacae complex species (10) that frequently carries mcr-10 and has been detected in humans (11), chickens (12), and dogs (13).

The tmexCD1-toprJ1-like gene cluster was identified in plasmids pHNLW22-2 and pHN21SC1505-3. Compared with tfnxB1-tmexCD1-toprJ1, this gene cluster lacks the upstream regulator gene tfnxB, thereby forming the structure of tmexC-tmexD-toprJ1. At the nucleotide level, it displays high homology with tmexCD1-toprJ1, tmexCD2-toprJ2, and tmexCD3-toprJ1b (Table S3). Moreover, at the amino acid level, the proteins encoded by this gene cluster shared 98.97%, 96.84%, and 92.26% identity with TMexC1, TMexD1, and TOprJ1; 97.68%, 98.56%, and 92.47% identity with TMexC2, TMexD2, and TOprJ2; and 98.20%, 98.09%, and 92.26% identity with TMexC3, TMexD3, and TOprJ1, respectively. Phylogenetic analysis of this gene cluster with related nucleotide sequences revealed that it belongs in a separate branch (Fig. 1A). Therefore, we assigned this gene cluster to a new allele, tmexCD4-toprJ4.
| Strain      | Bacterial species | Plasmid or chromosome | Plasmid type | Size (bp) | Resistance gene(s) | MIC of tigecycline (+ NMP) (mg/L) |
|------------|-------------------|-----------------------|--------------|-----------|--------------------|-----------------------------------|
| GLW9C22    | K. quasipneumoniae | Chromosome            |              | 5,129,993 | aqxAB, fosA         | 32 (0.25)                         |
|            |                   | pHNWL22-1             | IncFIB(K)    | 259,761   | aac(3)-IV, aph(4):la, aaph(3 ’)-la, floR, aada2, aph(3 ’)-la, tet(A) |                    |
|            |                   | pHNWL22-2             |              | 34,729    | tmexCD4-toprJ4, tet(A), floR |                    |
|            |                   | pHNWL22-3             |              | 4,150     |                    |                                    |
| GD21SC1505 | E. rogenkampii    | Chromosome            |              | 4,780,540 | blaoAB-6           | 16 (0.25)                         |
|            |                   | pHN21SC1505-1         | IncFII       | 90,290    |                    |                                    |
|            |                   | pHN21SC1505-2         |              | 40,695    | qns1, tet(A)       |                                    |
|            |                   | pHN21SC1505-3         |              | 34,729    | tmexCD4-toprJ4, tet(A), floR |                    |
|            |                   | pHN21SC1505-4         |              | 27,664    |                    |                                    |
|            |                   | pHN21SC1505-5         |              | 5,027     |                    |                                    |
| BW25113    | E. coli           | pHNWL22-2             |              | 34,729    | tmexCD4-toprJ4, tet(A), floR | 0.125 (0.125)                   |
| BW25113-pHNLW22-2 | E. coli       | pHNWL22-2             |              | 34,729    | tmexCD4-toprJ4, tet(A), floR | 4 (0.25)                         |
| BW25113-pHSG575 | E. coli        | pHSG575                |              | 90,290    |                    | 0.125 (0.125)                   |
| BW25113-pHSG575-tet(A) | E. coli | pHSG575-tet(A)        |              | 40,695    | tet(A)             | 0.25 (0.25)                     |
| BW25113-pHSG575-tmexCD4-toprJ4 | E. coli | pHSG575-tmexCD4-toprJ4 |              | 34,729    | tmexCD4-toprJ4     | 1 (0.25)                        |
| BW25113-pHSG575-tet(A)-tmexCD4-toprJ4 | E. coli | pHSG575-tet(A)-tmexCD4-toprJ4 |              | 27,664    | tmexCD4-toprJ4, tet(A) | 2 (0.25)                        |
| AH58I      | K. pneumoniae    | pHNWL22-2             |              | 34,729    | tmexCD4-toprJ4, tet(A), floR | 0.03 (0.03)                    |
| AH58I-pHNLW22-2 | K. pneumonia   | pHNWL22-2             |              | 34,729    | tmexCD4-toprJ4, tet(A), floR | 0.03 (0.03)                    |
| AH58I-pHS575 | K. pneumonia   | pHS575                |              | 90,290    |                    | 0.06 (0.06)                     |
| AH58I-pHS575-tet(A) | K. pneumonia | pHS575-tet(A)        |              | 40,695    | tet(A)             | 1 (0.25)                        |
| AH58I-pHS575-tmexCD4-toprJ4 | K. pneumonia | pHS575-tmexCD4-toprJ4 |              | 34,729    | tmexCD4-toprJ4     | 1 (0.25)                        |
| AH58I-pHS575-tet(A)-tmexCD4-toprJ4 | K. pneumonia | pHS575-tet(A)-tmexCD4-toprJ4 |              | 27,664    | tmexCD4-toprJ4, tet(A) | 2 (0.125)                       |
To verify the role of tmexCD4-toprJ4 in antimicrobial resistance, a recombinant plasmid, pHSG575-tmexCD4-toprJ4, was constructed. Compared to the empty vector pHSG575, it showed an 8-fold increase in the MICs of tigecycline in both host strains, *Escherichia coli* BW25113 and *K. pneumoniae* AH58I. Reduced susceptibility was also observed for ciprofloxacin (4-fold increase), cefquinome (8-fold increase), and apramycin (2-fold increase). Therefore, like tmexCD1-toprJ1, tmexCD4-toprJ4 is also a multidrug resistance gene cluster. However, the MIC of tigecycline conferred by this recombinant plasmid was 8-fold lower in recipient *K. pneumoniae* AH58I than in the wild-type strain GLW9C22 (Table 1). Further sequence analysis revealed that a major facilitator superfamily transporter gene, *tet*(A), was present upstream of tmexCD4-toprJ4. Hence, to investigate the effect of Tet(A) on the function of TMexCD4-TOprJ4, two recombinant plasmids, pHSG575-tet(A)-tmexCD4-toprJ4 and pHSG575-tet(A), were constructed by cloning and ligating the *tet*(A) sequence as well as the intergenic sequence between tmexCD4-toprJ4 and tet(A). Relative to the strain with the empty vector, pHSG575-tet(A) and pHSG575-tet(A)-tmexCD4-toprJ4 increased tigecycline MIC by 2-fold and 16-fold in BW25113 and AH58I, respectively. To further determine the antimicrobial-resistant phenotype mediated by TMexCD4-TOprJ4, an extremely sensitive host strain, BW25113ΔacrAB (an acrAB knockout mutant [14]), was used to express tmexCD4-toprJ4 and tet(A). The results also revealed that TMexCD4-TOprJ4 mediated tigecycline resistance with a 32-fold increase in tigecycline MIC. Expression of both tmexCD4-toprJ4 and tet(A) resulted in a 64-fold increase in tigecycline MIC, suggesting a synergistic effect of TMexCD4-TOprJ4 and TetA. Therefore, the
coexistence of tmexCD4-toprJ4 and tet(A) allows the host strain to survive better under drug selection pressure. To further check the expression of tmexCD4-toprJ4 and tet(A), the transcriptional levels of tmexC4, tmexD4, toprJ4, and tet(A) were measured. Among strains carrying tet(A), tmexCD4-toprJ4, or tet(A)-tmexCD4-toprJ4, the expression of tmexCD4-toprJ4 and that of tet(A) were at a similar level. However, expression levels of these genes in BW25113ΔacrAB were higher than those in BW25113 (Fig. S1), which might be due to the changes of the acrAB-related regulators caused by the acrAB deletion (15). But the specific reasons need to be further analyzed.

To confirm that TMexCD4-TmprJ4 acts as an efflux pump and acts synergistically with Tet(A), we determined the intracellular accumulation of tigecycline in E. coli carrying tet(A), tmexCD4-toprJ4, tet(A)-tmexCD4-toprJ4, or the control plasmid pHSG575. BW25113 and BW25113ΔacrAB were employed as sensitive hosts to assess efflux efficiency. Compared with bacterial cells carrying the empty plasmid pHSG575, both host strains expressing tmexCD4-toprJ4 significantly decreased the intracellular amount of tigecycline, whereas cells carrying tet(A) also showed slightly lower intracellular drug concentrations (Fig. 1B). It is worth noting that the intracellular drug concentration in BW25113ΔacrAB-tmexCD4-toprJ4 was similar to that in BW25113-tmexCD4-toprJ4, implying that TMexCD4-TmprJ4 independently functioned as a drug transporter. Furthermore, we observed an almost equal concentration of intracellular tigecycline accumulation in cells carrying tmexCD4-toprJ4 or tet(A)-tmexCD4-toprJ4 in both bacterial hosts, demonstrating the prominent role of TMexCD4-TmprJ4 in the efflux of tigecycline. According to these results, it could be found that the resistance phenotype and drug efflux capacity conferred by tmexCD4-toprJ4 in BW25113 or BW25113ΔacrAB are at similar levels, indicating that the function of tmexCD4-toprJ4 is independent of acrAB. However, there are differences in the drug efflux capacity between BW25113-pHSG575-tet(A) and BW25113ΔacrAB-pHSG575-tet(A), indicating that as previously reported, the tigecycline efflux function of tet(A) is acrAB dependent (16, 17).

Further whole-genome sequencing analysis showed that pHNLW22-2 and pHN21SC1505-3 were both 34,729-bp in length and had a GC content of 60.04%, differing by only five single nucleotide polymorphisms (SNPs). Therefore, pHNLW22-2 was used as the test plasmid in subsequent experiments. A 15-day antibiotic-free stability test indicated that pHNLW22-2 was stable in the original strain as well as in E. coli BW25113 (Fig. S2), suggesting that this small plasmid can replicate and remain stable in subcultures. pHNLW22-2 failed to be transferred from donor strain GLW9C22 into recipient E. coli J53 via the conjugation assay at 30°C, 37°C, and 42°C because of the lack of a functional conjugative system (Fig. 1C). However, it was transformed into BW25113, BW25113ΔacrAB, and AH58I by electrotransformation, and the tigecycline MIC was increased by 32-fold, 128-fold, and 32-fold, respectively (Table 1). This plasmid was untypeable using PlasmidFinder, and the complete structure of pHNLW22-2 included a replication system (repA), a partitioning system (parA, parC), a mobilization system (mobABC), and an ~21-kb variable region (Fig. 1C). BLASTn results showed that the backbone of pHNLW22-2 was highly similar to the untypeable plasmid pRSB101, with 90% coverage and 99.05% identity, which were identified from uncultured bacteria in sludge. It also exhibited high levels of homology to the backbone of blaGES-2Kα-carrying plasmid pSTW0522-60-3 (Enterobacter kobei, Japan, sewage, AP022449) and blaKPC-2-carrying plasmid pKPC-038c (Aeromonas sp., United States, wastewater, CP026230), with 81% coverage and 98.8% identity (Fig. 1C). The main differences among these plasmids are the different resistance genes or heavy metal resistance genes and the diversity of insertion sequences carried in the variable region. Previous studies indicated that the replication region and partitioning system of pRSB101 (pHNLLW22-2-like plasmid) were highly similar to those of plasmids isolated from environmental bacteria and phytopathogenic bacteria (such as Xanthomonas and Aeromonas), suggesting that pHNLLW22-2-like plasmids likely originated from environmental bacteria (18). In addition, the pRSB101 plasmid can be mobilized to the recipient strain by a self-transmissible helper plasmid (IncP-1α plasmid RP4), and pHNLLW22-2 and pHNLLW22-2-like plasmids have spread to various bacterial species and may belong to broad-host-range plasmids (18). Therefore, the broad host range and transferable characteristics of pHNLLW22-2 will accelerate the transmission of tet(A)-tmexCD4-toprJ4 gene clusters in different bacterial
species. In addition to tmexCD4-toprJ4, more antimicrobial resistance genes originating from environmental bacteria might be captured by the pHNLW22-2-like plasmid and then transferred to other bacteria, further exacerbating the increasingly serious challenges of antimicrobial resistance.

Further genetic context analysis of tmexCD4-toprJ4 revealed a TnAs1-hp-tet(A)-tmexC4-tmexD4-toprJ4-IS26 structure (Fig. 1C), which was different from the genetic structures of three previously reported tmexCD-toprJ-like gene clusters associated with site-specific integrases, IS26 (6) or ICE (5). Although two mobile elements, TnAs1 and IS26, were present on the flanks of tmexCD4-toprJ4, there was no evidence that TnAs1 and IS26 could mediate the transfer of tmexCD4-toprJ4. Therefore, we hypothesized that tmexCD4-toprJ4 may have been transferred into this plasmid through multiple insertional recombination events.

In conclusion, we are the first to report a novel plasmid-mediated RND family efflux pump, tmexCD4-toprJ4, carried by an untypable plasmid in K. quasipneumoniae and E. rogenkampii isolated from chicken meat and environmental samples from farm markets. tmexCD4-toprJ4 significantly differs from the previously reported tigecycline resistance efflux pump clusters at both the nucleotide and amino acid levels, as well as in the genetic structure. TMexCD4-ToprJ4 mediated low-level tigecycline resistance and exhibited a synergistic effect with Tet(A) in reducing tigecycline susceptibility. Therefore, screening for the tmexCD-toprJ-like gene cluster should be urgently included in the surveillance of tigecycline-resistant Gram-negative pathogens in animals, humans, and the environment.

Data availability. The complete sequences of the GLW9C22 chromosome and four plasmids and the GD21SC1505 chromosome and five plasmids were submitted to GenBank with accession numbers CP089441 to CP089444 and CP091081 to CP091086, respectively.

SUPPLEMENTAL MATERIAL
Supplemental material is available online only.

SUPPLEMENTAL FILE 1, PDF file, 0.3 MB.

ACKNOWLEDGMENTS
This work was supported in part by the Laboratory of Lingnan Modern Agriculture Project (NT2021006), the National Natural Science Foundation of China (no. 32141002 and 31625026), and the Local Innovative and Research Teams Project of Guangdong Pearl River Talents Program (2019B020N05).

X.G. was responsible for most of the experiments as well as data analysis and article writing, C.W. was responsible for drug accumulation assay, data analysis and article writing.

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