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ISIR-Mediated Plasticity of IncL/M Plasmids Leads to the Insertion of *bla*_{OXA-48} into the *Escherichia coli* Chromosome

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The OXA-48 carbapenemase is mainly encoded by ~62-kb IncL/M plasmids. However, chromosome-mediated genes have been observed in *Escherichia coli* isolates. In this work, we investigated the genetic environment of OXA-48 in members of the family *Enterobacteriaceae* ($n = 22$) to understand how the OXA-48-encoding gene is transferred into the *E. coli* chromosome. The OXA-48-encoding gene was located within intact Tn1999.2 transposons in the ~62-kb plasmids or within a truncated variant of Tn1999.2 for the OXA-48-encoding genes located in the chromosomes of *E. coli* bacteria. The analysis of the Tn1999.2 genetic environment revealed an inverted orientation of the transposon in five ~62-kb plasmids (5/14 [35%]) and in all chromosome inserts ($n = 8$). The sequencing of pRA35 plasmid showed that this orientation of Tn1999.2 and the acquisition of an ISIR insertion sequence generated a 21.9-kb ISIR-based composite transposon encoding OXA-48 and designated Tn6237. The sequencing of a chromosomal insert encoding OXA-48 also revealed this new transposon in the *E. coli* chromosome. PCR mapping showed the presence of this element in all strains harboring an OXA-48-encoding chromosomal insert. However, different insertion sites of this transposon were observed in the *E. coli* chromosome. Overall, these findings indicate a plasticity of the OXA-48 genetic environment mediated by ISIR insertion sequences. The insertion sequences can induce the transfer of the OXA-48-encoding gene into *E. coli* chromosomes and thereby promote its persistence and expression at low levels.

Carbapenems are broad-spectrum antibiotics that constitute the last-line therapeutic option available to treat infection caused by multidrug-resistant members of the family *Enterobacteriaceae* (1). However, owing to the emerging resistance to carbapenems worldwide, the antimicrobial activity of these drugs is no longer guaranteed (2). An underlying mechanism is the acquisition of carbapenem-hydrolyzing β -lactamases encoded by genes located on mobile genetic supports, which facilitate their diffusion among bacteria (3).

The OXA-48 carbapenemase was initially identified from a *Klebsiella pneumoniae* isolate in Turkey (4). It then spread to various *Enterobacteriaceae* species especially throughout the southern Mediterranean area and in Europe (4, 5). The OXA-48-encoding genes (*bla*_{OXA-48}) are mostly found in *K. pneumoniae* and, to a lesser extent, in *Escherichia coli* and *Enterobacter* spp. (5). Their emergence is mediated by the rapid spread of strains containing broad-host-range conjugative IncL/M plasmids (6–8) harboring *bla*_{OXA-48} located within the Tn1999-type composite transposon (6, 9–11). The plasmid implicated in the diffusion of *bla*_{OXA-48} was a widespread 62-kb IncL/M plasmid (6). Derivatives have also been reported (7, 8). However, *bla*_{OXA-48} was observed in *E. coli* chromosomes, including in a strong killer strain (12).

The aim of this work was to investigate the genetic environment of *bla*_{OXA-48} in order to understand how this gene is transferred into the *E. coli* chromosome.

MATERIALS AND METHODS

Bacterial isolates. Twenty-two *Enterobacteriaceae* strains with reduced susceptibility or resistance to ertapenem and/or imipenem were used in the study (Table 1). Nineteen strains were recovered from clinical samples of patients hospitalized in Nini Hospital, Tripoli, Lebanon, between January 2008 and December 2012. The other three strains were collected in December 2012 during a study investigating intestinal carriage in healthy children. The strains were identified using the matrix-assisted

laser desorption ionization–time of flight (MALDI-TOF) mass spectrometry (MS) Vitek microbial identification system (bioMérieux, La Balme, France). Rifampin-resistant *E. coli* C600 was used for mating-out assays.

Determination of genetic relatedness and *E. coli* phylogroup. The genetic relatedness of the strains was determined by multilocus sequence typing (MLST). Allelic profiles and sequence types (STs) were assigned using the MLST schema of Diancourt et al. (13) and Wirth et al. (14) for *K. pneumoniae* and *E. coli* isolates, respectively.

Molecular characterization of *bla*_{OXA-48} and the associated transposon. The OXA-48-encoding genes were detected by PCR amplification as previously reported (12). The genetic environment of the *bla*_{OXA-48} gene was further investigated by PCR and sequencing using specific primers to map the transposon composites Tn1999 to Tn1999.4 (6, 9–11) (Table 2). The orientation of these transposons in their genetic environments was investigated by PCR and sequencing using combinations of primers targeting *bla*_{OXA-48} (5'-AATACACGCATAACGTCCCC-3' and 5'-GCCATCACAAAAGAAGTGCTC-3'), *lysR* (5'-GCTGCAAATAGCATCATA CC-3'), *orf67* (5'-CAGCCAGAATAAGAGCAATC-3'), and *pemI* (5'-TG ACCATGCCAACTTCATT-3'). The sequences of PCR products were established on both strands by dideoxy chain terminator using an Applied Biosystems sequencer.

Plasmid analysis. The transferability of the *bla*_{OXA-48} gene was studied by a mating-out assay. Selection was performed on agar plates supplemented with ticarcillin (32 μ g/ml) and rifampin (300 μ g/ml). The plasmid content of bacteria and the size of plasmids were determined using plasmid DNA extracted by the method of Kado and Liu (15) with the plasmids Rsa (39 kb),

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TABLE 1 Phenotypic and genotypic characteristics of OXA-48-producing isolates

Isolate	ST ^a (<i>E. coli</i> phylogroup)	Clinical sample ^b (yr)	OXA-48 genetic environment				Genetic support ^d
			Transposon	PCR result ^c			
				<i>parA</i>	<i>repA</i>	<i>traU</i>	
<i>E. coli</i> isolates							
EC37	88 (A)	Stool ^e (2012)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC267	88 (A)	Sputum (2012)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC264	227 (A)	Urine (2012)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC15	127 (B2)	Sputum (2011)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC8	131 (B2)	Urine (2012)	Inverted Δ Tn1999.2	+	+	+	Chrom
EC254	38 (D)	Urine (2012)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC265	38 (D)	Pus (2012)	Inverted Δ Tn1999.2	+	–	–	Chrom
EC49	38 (D)	Stool ^e (2012)	NT ^f	+	–	–	Chrom
EC253	617 (A)	Urine (2012)	Tn1999.2	+	+	+	62kb
EC269	617 (A)	Urine (2012)	Tn1999.2	+	+	+	62kb
EC119	3877 (A)	Stool ^e (2012)	Tn1999.2	+	+	+	62kb
EC260	1711 (B1)	Urine (2012)	Tn1999.2	+	+	+	62kb
<i>K. pneumoniae</i> isolates							
KP78	1157	Blood (2012)	Tn1999.2	+	+	+	62kb
KP87	1156	Pus (2012)	Tn1999	+	+	+	62kb
KP34	45	Urine (2011)	Tn1999.2	+	+	+	62kb
KP9	866	Urine (2009)	Inverted Tn1999.2	+	+	+	62kb
KP26	866	Urine (2008)	Tn1999.2	+	+	+	62kb
<i>E. aerogenes</i> EA2		Pus (2012)	Inverted Tn1999.2	+	+	+	62kb
<i>E. cloacae</i> isolates							
ECL6		Urine (2010)	Inverted Tn1999.2	+	+	+	62kb
ECL8		Urine (2009)	Tn1999.2	+	+	+	62kb
<i>C. koseri</i> CK8		Urine (2011)	Inverted Tn1999.2	+	+	+	62kb
<i>R. planticola</i> RA35		Pus (2011)	Inverted Tn1999.2	+	+	+	62kb

^a Sequence types for *E. coli* and *K. pneumoniae* isolates.

^b Clinical sample that the strain was isolated from.

^c PCR result for “pOXA-48a genes.”

^d Chrom, chromosome-mediated OXA-48; 62kb, ~62-kb IncL/M plasmid encoding OXA-48.

^e Fecal carriage in children having no known contact with a hospital.

^f NT, nontypeable Tn1999-type environment.

TP114 (61 kb), pCFF04 (85 kb), and pCFF14 (180 kb) as standards. Plasmid restriction analysis was performed from DNA extracted by alkaline lysis and digested with EcoRI and HindIII restriction endonucleases (Boehringer Mannheim, Meylan, France). PCR-based replicon typing (PBRT) was used to identify plasmid incompatibility groups in transconjugants (16). The *repA*, *traU*, and *parA* genes were detected by PCR to relate the OXA-48-encoding plasmids to the pOXA-48a IncL/M plasmid as previously described (6).

Chromosome analysis. The chromosomal location of the *bla*_{OXA-48} gene was investigated by pulsed-field gel electrophoresis (PFGE) using the I-CeuI endonuclease and hybridization with probes specific for the 16S rRNA genes, *bla*_{OXA-48} gene and IS1999, as previously described (17). The chromosomal insertion of *bla*_{OXA-48} was further mapped by PCR using *bla*_{OXA-48}-containing DNA fragments purified from pulsed-field gels and primers specific for pOXA-48a. Seventeen primer pairs (Table 2) covering pOXA-48a plasmid backbone were used for PCR mapping; the targets were *trbN*, *pemI*, *mucA*, *nuc*, *orf22*, *orf23*, *orf24*, *korC*, *orf28*, *orf29*, *orf30*, *mobA*, *traJ*, *traU*, *repA*, and *parA*.

Amplification by nested TAIL-PCR. The sites flanking the pOXA-48 insertion site in the *E. coli* chromosome were amplified by a nested thermal asymmetric interlaced PCR (TAIL-PCR) approach. The primer targeting the pOXA-48 insert was ORF-25-R1 (R stands for reverse) (5'-GCCAGC GAGAAGCGAACAAAACG-3'), and the arbitrary degenerate primers were TAIL-PCR-F1 (F stands for forward) (5'-GTAATACGACTCACTAT AGGGCAGCGTGGTNTCGASTWTSWGTGTT-3'), TAIL-PCR-F2 (5'-GTAATACGACTCACTATAGGGCAGCGTGGTNGTCASWGANA

WGAA-3'), and TAIL-PCR-F3 (5'-GTAATACGACTCACTATAGGGCA CGCGTGGTWTGNAGWANCANAGA-3'). The amplifications comprised 15 standard PCR cycles, including 10 high-stringency cycles (annealing temperature, 62°C) and then 5 low-stringency cycles (annealing temperature, 35°C). The PCR product (1 μ l) was then amplified under high-stringency conditions by nested PCR (annealing temperature, 62°C) using the primer ORF25-R2 (5'-CGCAACATCAAACGAGCTCC-3') and a primer targeting the adaptor part of the degenerated primers (5'-G TAATACGACTCACTATAGGGC-3').

Plasmid and chromosome sequencing. Plasmid DNA of *Raoultella planticola* strain RA35 was purified with the NucleoBond Xtra Midi kits (Macherey-Nagel) following the manufacturer's instructions. After fragmentation, DNA was sequenced with Roche 454 GS-FLX system. The genomic DNA of *E. coli* EC15 was extracted and purified using the Genra Puregene Yeast/Bact kit (Qiagen, Valencia, CA) and sequenced using Illumina sequencing technology with 500-bp paired-end libraries.

Bioinformatic analyses. The *de novo* assemblies were performed with MIRA package (mimicking intelligent read assembly implemented in Linux) (18). The contiguous sequences (contigs) were then aligned with the pOXA-48a plasmid sequence (JN626286) using the BLAST algorithm (19). The gaps and insertions between contigs were closed or checked by PCR and Sanger sequencing. Open reading frames (ORFs) were predicted and annotated using RATT (20). The resulting annotation was manually checked and curated using Artemis (21). The resulting DNA sequences were compared with the pOXA-48a sequence using Easyfig (22).

TABLE 2 Primers used in this study

Target	Primer sequence (5' to 3' end)	Positioning of primers in pOXA-48a sequence	Reference
IS1999	CAGCAATTCTTTCTCCGTG CAAGCACAAACATCAAGCGC	2.780–2.798/7.520–7502 ^a 3.764–3.746/6.536–6.554 ^a	6
bla _{OXA-48}	AATACACGCATAACGTCGCC GCCATCACAAAAGAAGTGCTC	6.232–6.251 5.489–5.469	32
lysR	GCTAGTGCCAATCTTACAGG GCTGCAAATAGCATCATACC	4.983–5.002 4.262–4.243	10
pemI	TGACCATGCCAACTTCATT AAACATACCAGCCTTTAACG CCCTCCTGACCAGCCGCC	8.022–8.004 7.869–7.850 8.163–8.149	This study
trbB	CACCGCTTCACCAGAAAA	61.698–61.715	This study
trbA	GAAGCGTTCCGAGAATTTG	519–501	This study
trbN	CAGAGAATTTTGGTAAACC CCGCCCTCAATCCCGCTGC	1.314–1.332 1.688–1.670	This study
mucA	CGTTCACTGTCATGAGACAC GGATCAATACCTTCTCCG	8.995–9.014 10.203–10.187	This study
parA	GCAGTGAAAACGTTGATCAG GATCGCAATGCGTCTTGGTG	18.746–18.765 19.277–19.258	6
nuc	GATAAGAATCTGGAAGAAAC GAGCCTTGATAATACGACG	20.047–20.066 20.745–20.727	This study
orf22	CCTGCAGGGCTATTTATTC	22.482–22.500	This study
korC	TTTTGCGCTGCTATTTC	23.859–23.842	This study
orf25	GGAATAGCAGCGCAAAA CCATAACATCGCCATCAT ATGATGGCGATGTTATGG GGTTCGATGCCCGTATTG	23.842–23.859 24.048–24.031 24.031–24.048 25.237–25.220	This study
orf28	GACTGGCGCAGGAGTAAA	28.046–28.063	This study
orf30	GTGGTTCAGGGTGAACAG	29.678–29.661	This study
mobB	TATCTGATGAGCCGTAGC	33.189–33.206	This study
mobA	CGCTCTCCATCGACTTC	34.487–34.470	This study
traJ	GCACCAGGAAGTTTGAT CGGGTCTTTTCAGTTGCA	37.849–37.866 38.251–38.234	This study
traU	ATCTCACGCAATCTTACGTC TCGCGTCATGCGTGATCTTC	48.129–48.148 48.705–48.686	6
repA	GACATTGAGTCAGTAGAAGG CGTGCAGTTCGTCTTTCGGC	56.427–56.446 57.351–57.332	6

^a Positioning of primers corresponding to the two copies of IS1999.

RESULTS

Genetic background of bacteria. The 22 isolates included in this study produced the carbapenemase OXA-48. The isolates belonged to the *E. coli* (clinical strains [$n = 9$]; colonizing strains [$n = 3$]), *K. pneumoniae* ($n = 5$), *Enterobacter cloacae* ($n = 2$), *Enterobacter aerogenes* ($n = 1$), *Raoultella planticola* ($n = 1$), and *Citrobacter koseri* ($n = 1$) species. The genetic background of *K. pneumoniae* and *E. coli* producing OXA-48 was investigated by MLST (Table 1). Three *K. pneumoniae* isolates (KP9, KP26, and KP112) belonged to sequence type 866 (ST866). The three other

K. pneumoniae isolates were assigned to ST45 (KP34), ST1156 (KP78), and ST1157 (KP87). Seven STs were represented among *E. coli* isolates (ST227 [$n = 2$], ST617 [$n = 2$], ST38 [$n = 3$], ST88 [$n = 2$], ST131 [$n = 1$], ST1711 [$n = 1$], and ST127 [$n = 1$]), and they covered all *E. coli* phylogroups (Table 1).

Location of bla_{OXA-48} in plasmids or chromosome. OXA-48-encoding transconjugants were obtained from 14 strains with OXA-48-encoding genes. Plasmids from natural isolates and their transconjugants were extracted and hybridized with a bla_{OXA-48} probe. The results showed that the bla_{OXA-48} gene was located on

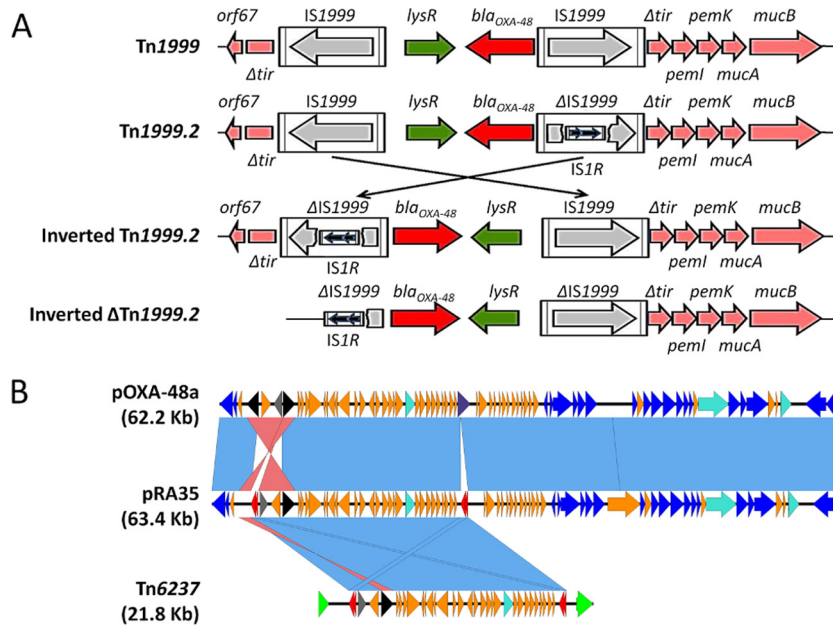


FIG 1 Comparison of pRA35 plasmid and *bla*_{OXA-48} chromosome-mediated insert (Tn6237) with pOXA-48a plasmid. (A) Schematic diagram showing the rearrangements in Tn1999-type transposons encoding OXA-48. (B) Global comparison performed using the bioinformatics tools Easyfig. The transposases of insertion sequences are represented by red arrows for IS1R and black arrows for IS1999, *bla*_{OXA-48} by gray arrows, genes implicated in transfer and mobility (*mob* and *tra*) by navy blue arrows, genes specific to pOXA-48a-like plasmids (*parA*, *traU*, and *parA*) by light blue arrows, and *E. coli* chromosome genes by green arrows.

~62-kb plasmids (64% [14/22] [Table 1]). Further analysis of these plasmids from transconjugants revealed similar restriction profiles (data not shown). None of the plasmids could be assigned to an incompatibility group by the PBRT method. However, the *repA*, *traU*, and *parA* genes of pOXA-48a were detected in all plasmids, suggesting that the plasmids had an IncL/M pOXA-48a-like backbone.

Despite many attempts, no transconjugants or transformants were obtained from eight unrelated *E. coli* isolates belonging to the A, D, and B2 *E. coli* phylogroups. The insertion of *bla*_{OXA-48} into the bacterial chromosome was therefore investigated by PFGE after I-CeuI nuclease treatment and hybridization. Distinct DNA fragments, with a range of 600 to 880 kb, hybridized with the *bla*_{OXA-48} probe (data not shown), suggesting that the insertion of *bla*_{OXA-48} in *E. coli* chromosomes is not rare (8/22 [36%]) and may occur at different sites.

Genetic environment of *bla*_{OXA-48}. The *bla*_{OXA-48} gene was identified as part of the transposons designated Tn1999 to Tn1999.4 (6, 9–11). To characterize the genetic environment of *bla*_{OXA-48} in our strains, we mapped by PCR and sequencing the Tn1999-type transposons harboring *bla*_{OXA-48}. The *bla*_{OXA-48} gene was predominantly associated with Tn1999.2 (68% [15/22]). However, the eight *bla*_{OXA-48} genes on *E. coli* chromosomes (32% [7/22]) were located in incomplete Tn1999.2 elements, in which the IS1999 element flanking *bla*_{OXA-48} was truncated. Finally, *bla*_{OXA-48} was carried by Tn1999 in *K. pneumoniae* KP87, and we were unable to identify a Tn1999-type element in *E. coli* EC49.

PCR mapping covering the whole sequence of pOXA-48a was then performed by targeting 17 genes and the sequences overlapping Tn1999.2 edges. The complete backbone of pOXA-48a was observed in all bacteria, except for *E. coli* harboring a chromosome-mediated OXA-48 gene, in which we observed PCR prod-

ucts only for sequences located between *bla*_{OXA-48} and *orf25* of pOXA-48a. In addition, no amplification was obtained with the primers targeting the sequences overlapping Tn1999.2 edges for five plasmids and in all *E. coli* harboring a chromosome-mediated OXA-48 gene, suggesting a significant rearrangement in the vicinity of the transposon encoding OXA-48.

Sequence of pRA35 plasmid. To investigate the rearrangement of Tn1999.2, we sequenced the ~62-kb plasmid of *R. planticola* RA35(pRA35) using a next-generation sequencing approach. The sequencing reactions of pRA35 plasmid yielded 9,607 reads with an average length of 407 bases. A total of 9,303 reads with an average length of 355 bases were conserved (97%) after trimming, amounting to a total number of 3,302,032 bases of DNA. After contig assembling, gap closing, and repeat sequence mapping by Sanger sequencing, a total of 8,576 reads (92% of reads with an average length of 354 bases) totaling 3,037,308 bases were assembled to obtain the complete sequence of pRA35, resulting in an average of 47 \times sequence coverage (range, 38 \times to 121 \times). Overall, the complete sequencing of pRA35 showed an IncL/M circular plasmid of 63.434 kb with an average G+C content of 51.2%. The pRA35 and pOXA-48a nucleotide sequences shared overall 99% identity, and most of the synteny was conserved. Differences in the DNA sequence were observed in several genes or open reading frames; the most divergent genes were the nuclease-encoding gene and the DNA primase gene. However, the pRA35 plasmid differed from the reference plasmid pOXA-48a in three major respects (Fig. 1): (i) the insertion of an additional copy of the IS1R insertion sequence in *orf25*, (ii) the presence of the Tn1999.2 transposon, which differs from Tn1999 by the insertion of the IS1R insertion sequence in the IS1999 insertion sequence located upstream of *bla*_{OXA-48}, and (iii) an inverted orientation of Tn1999.2. The orientation of Tn1999.2 was investigated in all

OXA-48-encoding plasmids by PCR mapping and sequencing. The inverted orientation was detected in five plasmids (5/14 [35%]).

Sequence of the chromosome insert containing the bla_{OXA-48} gene. To understand the insertion of bla_{OXA-48} into the *E. coli* chromosome, the bla_{OXA-48} environment was sequenced in *E. coli* strain EC15 using a next-generation sequencing approach. The sequencing reactions yielded 18,710,176 reads with an average length of 101 bases. A total of 18,441,693 reads of 93-base average lengths were conserved (98.6%) after trimming. Three contigs containing sequences corresponding to pOXA-48a were obtained by *de novo* assembling. After gap closing and repeat sequence mapping by Sanger sequencing, we obtained a contiguous sequence of 27.6 kb, which contained a 21.8-kb central fragment related to the pOXA-48a sequence flanked by 2.7-kb and 3.1-kb *E. coli* chromosome sequences. A total of 112,059 reads (0.6%) with an average length of 93 bases totaling 10,415,879 bases were aligned to 100% of this sequence, resulting in an average of 377× sequence coverage (range, 62× to 795×).

The sequence of the bla_{OXA-48} chromosomal insert shared 100% identity with the pRA35 sequence and identical synteny (Fig. 1). The insert was flanked by two copies of IS_{IR}, the first provided by the truncated and inverted Tn1999.2 transposon, and the second inserted into *orf25*, as observed in the pRA35 plasmid. The deletion in Tn1999.2 was due to the loss of residual IS₁₉₉₉ sequences upstream of Tn1999.2-located IS_{IR}. Between the IS_{IR} insertion sequences, after the truncated Tn1999.2 transposon, we observed 9 genes (*pemI*, *pemK*, *muca*, *mucB*, *resD*, *para*, *parB*, *nuc*, and *korC*) and 18 hypothetical ORFs, as previously observed in pRA35 and pOXA-48a. These genes and ORFs, associated with the flanking IS_{IR} insertion sequences, formed an IS_{IR}-based composite transposon designated Tn6237. The flanking regions of this element corresponded to the hypothetical gene *orf33* belonging to the II₅₃₆ pathogenicity island, which has been observed in the reference uropathogenic strain *E. coli* 536 (23). Twelve-base imperfect direct repeats (GATCACTTAGAT and GATCTTTAGAT) were observed at the site of inversion, suggesting that the insertion of Tn6237 into the *E. coli* chromosome is the result of a transposition event. PCR mapping of Tn6237 in *E. coli* showed that Tn6237 is conserved in all strains harboring a chromosome-mediated OXA-48-encoding gene. However, sequencing of nested TAIL-PCR products revealed distinct insertion sites. In the EC264 strain (ST227, phylogroup A), the bla_{OXA-48} insert was detected in the *yjfF* gene, which encodes a putative quinol oxidase subunit. In *E. coli* EC8 (ST131, phylogroup B2), the bla_{OXA-48} insert was detected in an intergenic region ~375 bp at the 5' site of the *pgi* gene (glucose-6-phosphate isomerase). Overall, these data showed that the chromosome-mediated bla_{OXA-48} inserts exhibited similar organization in all strains and corresponded to a novel composite transposon, which can be inserted in different sites of the *E. coli* chromosome.

DISCUSSION

Mobile genetic elements, including large broad-host-range plasmids, transposons, and insertion sequences (IS) have played a part in the evolution and generation of diversity of bacterial populations. They performed a critical role in the horizontal and vertical transfer and rapid spread of virulence factors and drug resistance genes in *Enterobacteriaceae*. IncL/M-type plasmids are currently detected worldwide in *Enterobacteriaceae* isolates of different ori-

gin and are considered to be epidemic resistance plasmids (24). They were implicated in the large dissemination of specific genes encoding extended-spectrum β-lactamases (ESBL), in particular the bla_{CTX-M-3} gene. IncL/M-type plasmids have also been reported to contribute to the diffusion of the carbapenemase-encoding genes bla_{NDM-1} and bla_{OXA-48}.

In this study, most bla_{OXA-48} genes were also located in ~62-kb IncL/M plasmids related to pOXA-48a, as previously observed in Europe and the Mediterranean region (25). However, most plasmids harbored bla_{OXA-48} inserted into the Tn1999.2 transposon and not into the Tn1999 transposon, as previously observed in pOXA-48a. In addition, 67% (8/12) of OXA-48-producing *E. coli* strains investigated in this study harbored a chromosome-mediated gene encoding OXA-48. The chromosomal location of bla_{OXA-48} is therefore not rare and is observed in unrelated *E. coli* isolates. This chromosomal location of bla_{OXA-48} was associated with the presence of a truncated form of Tn1999.2 and significant rearrangement in the vicinity of this element.

To investigate the mechanism underlying the chromosomal location of the OXA-48-encoding gene, we sequenced plasmid pRA35, the chromosomal insert encoding bla_{OXA-48} in *E. coli* EC15 and flanking regions of two additional chromosomal inserts. We observed a 21.9-kb bla_{OXA-48}-encoding and composite transposon designated Tn6237 in the *E. coli* chromosome and pRA35. This element was detected in all strains harboring an OXA-48-encoding chromosomal insert. It forms an IS_{IR}-based composite transposon harboring bla_{OXA-48}, which is responsible for its insertion into the *E. coli* chromosome.

IS₁ is one of the smallest bacterial insertion sequences isolated so far. It is 768 bp long, includes two ~23-bp imperfect inverted repeats (IRL [inverted repeat left] and IRR [inverted repeat right]) located at its ends, and two partly overlapping open reading frames (*insA* and *insB'*) located in the 0 and -1 relative translational phases, respectively (26). IS₁ transposition may appear as a simple insertion or as a replicative fusion event (27, 28). The origin of the second copy of IS_{IR} observed in pRA35 and Tn6237 may therefore be the result of the replicative transposition of the IS_{IR} insertion sequence located in Tn1999.2.

The IS₁ transposition can occur in both orientations, and the direction of IS₁ is not an important factor in transposition. Reports have mentioned that the gene being mobilized should be flanked by short terminal inverted repeats within the IS₁ sequence regardless of the orientation of the two IS₁ elements (29). The orientation of the IS_{IR} elements in Tn6237 and pRA35 is therefore compatible with the formation of a functional composite transposon. IS₁-based composite transposons (elements Tn9 and Tn1681) have already been reported (30). In Tn1681, the mobile element is flanked by two inverted repeats of IS₁, which bracket the *E. coli* heat-stable toxin. In Tn9 and derivatives, a gene encoding chloramphenicol acetyltransferase is bracketed by two direct repeats of the IS₁ insertion sequence, as observed in our bla_{OXA-48}-encoding transposon. In the *E. coli* EC15 chromosome, this IS_{IR}-based transposable element is flanked by direct repeats of the insertion site, which may be marks of a transposition event.

Investigation of the target specificity of the IS₁ element indicated that acquisition of IS₁ is generally located within or directly adjacent to the A+T-rich content segments (31). The Tn6237 insertion site in the EC15 strain is the *orf33* gene of the pathogenicity island (PAI) II₅₃₆, which has been previously reported in the uropathogenic *Escherichia coli* 536 strain (23). The *orf33* gene (1.668

kb) and surrounding sequence are characterized by a very low-GC percentage (~33%), as at the insertion sites in *E. coli* EC264 (~44%) and EC8 (~37%). Likewise, the *IS1R* insertion sites in the *orf25* gene in pRA35 and Tn6237 are identical and characterized by a low GC percentage (40%), which makes a putative hot spot of *IS1R* insertion in the backbone of pOXA-48a-like IncL/M plasmids and therefore a region likely to be variable.

In this work, we provide new insights into the *bla*_{OXA-48} genetic environment and describe its high plasticity, which is promoted by Tn1999.2 inversion and *IS1R* elements responsible for the transposition of the OXA-48-encoding fragment derived from pOXA-48 in *E. coli* chromosomes.

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We declare that we have no conflicts of interest.

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