Characteristics of the Conjugative Transfer System of the IncM Plasmid pCTX-M3 and Identification of Its Putative Regulators

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ABSTRACT Plasmid conjugative transfer systems comprise type IV secretion systems (T4SS) coupled to DNA processing and replication. The T4SSs are divided into two phylogenetic subfamilies, namely, IVA and IVB, or on the basis of the phylogeny of the VirB4 ATPase, into eight groups. The conjugation system of the IncM group plasmid pCTX-M3, from Citrobacter freundii, is classified in the IVB subfamily and in the MPFI group, as are the conjugation systems of IncI1 group plasmids. Although the majority of the conjugative genes of the IncM and IncI1 plasmids display conserved synteny, there are several differences. Here, we present a deletion analysis of 27 genes in the conjugative transfer regions of pCTX-M3. Notably, the deletion of either of two genes dispensable for conjugative transfer, namely, orf35 and orf36, resulted in an increased plasmid mobilization efficiency. Transcriptional analysis of the orf35 and orf36 deletion mutants suggested an involvement of these genes in regulating the expression of conjugative transfer genes. We also revised the host range of the pCTX-M3 replicon by finding that this replicon is unable to support replication in Agrobacterium tumefaciens,Ralstonia eutropha, and Pseudomonas putida, though its conjugation system is capable of introducing plasmids bearing oriT pCTX-M3 into these bacteria, which are representatives of Alpha-, Beta-, and Gammaproteobacteria, respectively. Thus, the conjugative transfer system of pCTX-M3 has a much broader host range than its replicon.

IMPORTANCE Horizontal gene transfer is responsible for rapid changes in bacterial genomes, and the conjugative transfer of plasmids has a great impact on the plasticity of bacteria. Here, we present a deletion analysis of the conjugative transfer system genes of the pCTX-M3 plasmid of the IncM group, which is responsible for the dissemination of antibiotic resistance genes in Enterobacteriaceae. We found that the deletion of either of the orf35 and orf36 genes, which are dispensable for conjugative transfer, increased the plasmid mobilization efficiency. Real-time quantitative PCR (RT-qPCR) analysis suggested the involvement of orf35 and orf36 in regulating the expression of transfer genes. We also revised the host range of pCTX-M3 by showing that its conjugative transfer system has a much broader host range than its replicon.

KEYWORDS IncM group, conjugative transfer, plasmid analysis, plasmid mobilization

Conjugative transfer is a prevalent phenomenon among bacteria; this phenomenon is crucial for horizontal gene transfer in the biosphere and is a major contributor to the rapid variability of bacterial genomes. In the process of conjugative transfer, DNA (a conjugative plasmid, a conjugative transposon, or an integrative conjugative element [ICE]) is transferred from a donor to a recipient cell after physical contact between the cells is established. The process may be regarded as a combination of DNA processing functions coupled to a type IV secretion system (T4SS; also known as mating pair formation [Mpf]) by a dedicated protein (coupling protein [CPI]). The DNA process-
ing functions are provided by the DNA transfer and replication (Dtr) system, also called the relaxosome complex (1). The T4SSs of Gram-negative bacteria are classified into two large phylogenetic groups, namely, IVA and IVB: the Agrobacterium tumefaciens VirB/D4 secretion system and the conjugation systems of the IncF and IncP plasmids are classified as type IVA (T4ASS) (2, 3), whereas type IVB (T4BSS) is represented by secretion systems found in Legionella pneumophila (Dot/Icm) and in other important pathogens (4, 5). The majority of the Dot/Icm proteins share homology with the constituents of the conjugation system of the R64 plasmid of the IncI group. Despite an increasing amount of information becoming available in recent years on the organization and regulation of T4BSSs, they are still less thoroughly characterized than T4ASSs.

The canonical T4SS is represented by the Agrobacterium tumefaciens VirB/D4 secretion system responsible for transfer DNA (T-DNA) transfer to plant cells during infection. This T4SS consists of 11 proteins (VirB1 to VirB11) and the coupling protein VirD4 (for review, see references 5 to 8). The translocation channel comprises the VirB3, B6, B7, B8, B9, and B10 proteins. In the translocation channel, three components form the core channel complex in the outer membrane (OM), also called outer membrane complex (OMC): VirB9, the pore-forming protein; VirB7, a small lipoprotein; and VirB10, the protein spanning both the inner membrane (IM) and the outer membrane (OM). Interactions of the OMC with the inner membrane complex (IMC), which comprises the VirB3, VirB6, and VirB8 proteins, and with the ATPases VirB4 and VirB11 result in the formation of a pore. The extracellular structure important for the establishment of contact between mating cells, namely, the T-pilus, is composed of the major subunit VirB2 and the minor component VirB5 localized at the tip. VirB3, the least-characterized Mpfs component, is also necessary for T-pilus assembly. Finally, VirB1 shows homology to a lytic transglycosylase that cleaves peptidoglycan (3, 9). The system is energized by three cytoplasmic ATPases: VirB4, VirB11, and the coupling protein VirD4. Of all of the Vir proteins listed above, VirB4 is the only component present in every T4SS described so far (10). The universal presence of VirB4 enabled all known Mpfs of both Gram-negative and Gram-positive bacteria and archaea to be divided into eight groups on the basis of VirB4 phylogeny (11). The A. tumefaciens VirB/D4 system and the conjugation system of the IncP plasmids are now classified in the MPFT group (11). The conjugation system of the F plasmid, one of the most well-known plasmids, belongs to the MPFF group (11). In addition, the IncI1 plasmid R64 codes for TraU, which is a distant VirB4 homologue, and constitutes the prototype of the MPF, group. The R64 conjugative transfer system is encoded by 22 transfer genes, namely, traE-traY, three traB-C genes, and the nuc gene, 16 of which have been shown to be indispensable for conjugation (12). The homology of R64 to the VirB/D4 system of A. tumefaciens is rather low. However, TraO displays homology to VirB10 (13), and TraM is distantly homologous to VirB8, TraJ to the VirB11 ATPase, and TraQ and TraR to the pilin subunit VirB2 (11).

Another plasmid encoding an MPF, conjugation system that displays homology to the T4BSS systems of the IncI1 plasmids is pCTX-M3 (accession no. AF550415), a member of the IncM incompatibility group (14). Plasmid pCTX-M3 was isolated from a clinical Citrobacter freundii strain in Poland in 1996 as a vector of the blaCTX-M-3 gene (14, 15). It is noteworthy that members of the IncM group are closely related not only to each other but also to plasmids of the IncL group (16), with which they were earlier classified jointly as the IncL/M group (17). IncL and IncM plasmids are widespread in Enterobacteriaceae (18) and are responsible for the dissemination of antibiotic resistance genes. These genes include blaCTX-M-3, which encodes an extended spectrum β-lactamase, blaNDM-1, which encodes a metallo-β-lactamase, blaOXA-48, which encodes a carbapenem-hydrolyzing enzyme (19, 20), blaKPC-4, which codes for the Klebsiella pneumoniae carbapenemase (21), blaIMP-4, which codes for imipenemase (22), and the aminoglycoside resistance gene armA (23).

The pCTX-M3 plasmid can be transferred by approximately 10% of cells in an Escherichia coli donor population under optimal conditions (14). Bacteria bearing
pCTX-M3 can also conjugate in liquid culture; however, in contrast to IncI1 plasmids, which require type IV pili for conjugation in liquid media, pCTX-M3 does not encode additional pili (14, 24, 25). The conjugative transfer genes of pCTX-M3 are localized in two separate regions with predicted operon structures, namely, \( \text{tra} \) and \( \text{trb} \) (14), and these genes do not exhibit substantial sequence similarity to genes with ascribed functions available in public databases apart from those encoded by IncI1 and IncL/IncM plasmids (Fig. 1). The \( \text{tra} \) and \( \text{trb} \) genes of IncL/IncM plasmids such as pCTX-M3 encode proteins that exhibit 39% to 65% similarity to those encoded by IncI1 plasmids (such as R64 and ColIb-P9) (Table 1); however, a number of genes from each system do not have counterparts in the other system (14). The pCTX-M3 plasmid can mobilize plasmids that contain the heterologous \( \text{oriT} \) (ColIb-P9) (from the IncI1 plasmid ColIb-P9), and plasmids with \( \text{oriT} \) pCTX-M3 can be mobilized by a ColIb-P9-derived plasmid (14); both of these mobilizations occur at low frequencies.

Here, we present a deletion analysis of the \( \text{tra} \) and \( \text{trb} \) genes potentially involved in pCTX-M3 conjugative transfer. We found that the deletion of either of the \( \text{orf35} \) and \( \text{orf36} \) genes, both of which are dispensable for pCTX-M3 transfer, increases the mobilization efficiency of \( \text{oriT} \) pCTX-M3-bearing plasmids into \( \text{E. coli} \) and \( \text{A. tumefaciens} \). The deletion of these genes also affected the transcription of other conjugative transfer genes. In addition, we verified the host range of the pCTX-M3 conjugation system and found that the host range of its replicon, reported previously to comprise \( \text{Alpha-} \), \( \text{Beta-} \), and \( \text{Gammaproteobacteria} \) (26), is in fact much narrower than previously believed and is limited to \( \text{Enterobacteriaceae} \).

RESULTS AND DISCUSSION

Organization of the regions coding for the conjugative transfer systems of the pCTX-M3 and R64 plasmids. In pCTX-M3, both the \( \text{tra} \) and \( \text{trb} \) regions coding for the conjugative transfer system display extensive conservation of synteny with the conjugation system genes of the IncI1 plasmids R64 and ColIb-P9 (Fig. 1) (14). However, there are certain differences. pCTX-M3 has no homologues of the \( \text{traEFG} \), \( \text{traST} \), and \( \text{traV} \) genes. Neither \( \text{traEFG} \) nor \( \text{traS} \) is required for the conjugative transfer of R64, whereas \( \text{traT} \) and \( \text{traV} \) are indispensable (12). Moreover, in pCTX-M3, the \( \text{nuc} \) gene, which encodes a nuclease, is located at a distance from the \( \text{tra} \) region. In addition, the single \( \text{orf38} \) located between \( \text{traR} \) and \( \text{traU} \) replaces \( \text{traST} \), while \( \text{orf36} \), which is found only in IncI and IncM plasmids, separates \( \text{traL} \) and \( \text{traM} \). Furthermore, the \( \text{trbN} \) gene of

FIG 1 Gene organization of type IV secretion systems: IVA, the \( \text{A. tumefaciens virB/virD4} \) region, and IVB, the transfer regions of pCTX-M3, R64, and the dot or icm region of \( \text{L. pneumophila} \). Open reading frames (ORFs) are represented by arrows indicating their orientation. The ORFs homologous in all systems are shown in the same colors (except black, gray, and white). The ORFs homologous in the type IVB systems are indicated by identical patterns. The ORFs homologous in both plasmids are shown in gray, and those specific for either system are shown in white. The bars under the genes (pCTX-M3, R64, and dot or icm) indicate homology of single genes. The lines above the genes indicate the gene designation, and the corresponding letter is indicated above each gene. Dotted lines indicate small regions of unrelated genes, and line breaks indicate large gaps. The ORFs not related to the T4SS are shown as black arrows.
pCTX-M3, which encodes a putative lytic transglycosylase, has no homologues in R64. The homologue of orf46 is also absent in R64. However, the major difference between these plasmids concerns the position of their oriT regions with the nikAB genes. In R64, oriT lies far from the tra genes in a tail-to-tail orientation with respect to the trb operon. In pCTX-M3, the oriT region, along with the nikAB genes, is situated immediately upstream of the tra genes, and the nikAB and tra genes are predicted to constitute a single operon (Fig. 1) (14).

Identification of genes necessary for conjugation. A systematic deletion analysis was performed for the pCTX-M3 genes in the tra and trb regions. For this purpose, a collection of 27 derivatives with a deletion in each of the genes in the tra and trb

### Table 1: Bioinformatics analysis of the predicted proteins encoded in the transfer region of pCTX-M3

| Protein | Size (aa) | Mol wt (kDa) | pl | SP | TMH | Motif found/putative function | Homologues in IncI1 plasmid and Legionella pneumophila dot or icm (aa [% aa sequence identity/% aa sequence similarity]) |
|---------|-----------|--------------|----|----|-----|--------------------------------|----------------------------------------------------------------------------------------------------------------------------------|
| Orf55   | 121       | 9.0          | 13.57 | No | No | Helix-turn-helix motif/nickase accessory protein | 6–108 (32/53) to YggA of R64 |
| NikA    | 105       | 11.6         | 9.41  | No | No | Relaxase domain/nickase (Rlx) | 3–101 (30/62) to NikA of R64 |
| NikB    | 658       | 75.1         | 9.53  | No | No | Lipoprotein/CTS              | 70–157 (45/65) to TraH of ColIb-P9; 29–258 (48/65) to TraH of ColIb-P9; 3–258 (28/46) to DotC of L. pneumophila |
| TrH     | 166       | 18.7         | 9.77  | Yes| 1  | TraH-like (TivB10)           | 11–369 (40–63) to TraP of ColIb-P9; 11–377 (31/51) to DoTb of L. pneumophila |
| Trl     | 259       | 29.2         | 9.33  | Yes| 1  | NA/CTS                       | 4–419 (30/45) to TraO of ColIb-P9; 208–390 (45/41) to DotG (IcmE) of L. pneumophila |
| Trj     | 387       | 43.3         | 6.05  | No | No | GSP/PIE domain with Walker motifs A and B/ATPase VirB11-like (TivB11) | 6–82 (41/61) to TraK of ColIb-P9; 6–80 (25/45) to IcmT of L. pneumophila |
| Trk     | 86        | 10.3         | 10.55 | No | 2  | NA/pilin                     | 8–114 (33/58) to TraL of ColIb-P9 |
| Pri     | 1070      | 117.6        | 5.00  | No | No | NA/primease (Pri)            | 53–254 (26/53) to TraM of ColIb-P9; 73–257 (31/53) to Dott (IcmL) of L. pneumophila |
| TraL    | 170       | 17.9         | 6.27  | Yes| 1  | NA/TrbC-like (TivB8)         | 3–101 (30/62) to NikA of ColIb-P9 |
| Orf56   | 221       | 25.0         | 9.41  | No | No | NA/CT5                       | 119–382 (45/58) to TraN of ColIb-P9; 47–381 (29/44) to DotH (IcmK) of L. pneumophila |
| TrM     | 260       | 29.5         | 5.72  | No | 1  | NA/CTA                       | 3–258 (48/65) to TraH of ColIb-P9; 29–258 (48/65) to TraH of ColIb-P9; 3–258 (28/46) to DotC of L. pneumophila |
| Trn     | 383       | 40.7         | 5.97  | Yes| 1  | VirB10 domain/CTS VirB10-like (TivB10) | 22–234 (30/47) to TraP of ColIb-P9 |
| TraO    | 449       | 47.6         | 5.90  | No | No | NA/CTS                       | 3–258 (48/65) to TraH of ColIb-P9; 29–258 (48/65) to TraH of ColIb-P9; 3–258 (28/46) to DotC of L. pneumophila |
| TraP    | 234       | 24.7         | 9.20  | No | No | NA/PII                       | 3–258 (48/65) to TraH of ColIb-P9; 29–258 (48/65) to TraH of ColIb-P9; 3–258 (28/46) to DotC of L. pneumophila |
| TraQ    | 176       | 18.5         | 8.49  | No | 3  | NA/pilin                     | 22–234 (30/47) to TraP of ColIb-P9 |
| TraR    | 129       | 13.5         | 9.72  | Yes| 3  | NA/pilin                     | 12–127 (26/47) to TraP of ColIb-P9 |
| Orf58   | 164       | 18.9         | 6.96  | No | No | NA/pilin                     | 8–114 (33/58) to TraL of ColIb-P9 |
| TraU    | 1016      | 114.1        | 6.24  | No | 3  | Walker motifs A and B/ATPase VirB4-like (TivB4) | 1–1,006 (35/56) to TraU of ColIb-P9; 27–1,008 (28/47) to DotO (IcmB) of L. pneumophila |
| TraW    | 402       | 42.3         | 8.58  | Yes| 3  | NA/entry exclusion system    | 472–506 (37/55) to TraY of ColIb-P9; 4–217 (25/45) to TraQ of ColIb-P9; 5–172 (25/43) to DoT of IcmC of L. pneumophila |
| TraX    | 216       | 24.1         | 9.23  | No | No | NA/entry exclusion system    | 55–132 (31/47) to ExcA of ColIb-P9 |
| TraY    | 726       | 78.3         | 5.58  | Yes| 3  | NA/entry exclusion system    | 3–258 (48/65) to TraH of ColIb-P9; 29–258 (48/65) to TraH of ColIb-P9; 3–258 (28/46) to DotC of L. pneumophila |
| ExcA    | 217       | 25.4         | 9.28  | Yes| 3  | NA/pilin                     | 55–132 (31/47) to ExcA of ColIb-P9 |
| Orf46   | 169       | 19.2         | 9.79  | No | No | Lytic transglycosylase signature/Slt | None |
| TrbC    | 695       | 79.5         | 5.08  | No | 3  | Walker motifs A and B/CP (Cpl) | 29–609 (34/52) to TrbC of ColIb-P9; 107–599 (30/50) to DotL (IcmO) of L. pneumophila |
| TrbB    | 338       | 37.4         | 9.10  | No | 1  | Thioredoxin-like domain/NA   | 121–256 (37/47) to TrbB of ColIb-P9 |
| TrbA    | 435       | 49.1         | 6.43  | No | 3  | NA/CP (Cpl)                  | 13–124 (41/57) to TrbA of ColIb-P9; 80–368 (24/44) to DotM (IcmP) of L. pneumophila |
| TrbN    | 131       | 14.7         | 9.14  | No | No | Lytic transglycosylase signature/Slt | None |

- **a**Calculated with the use of the ProtParam tool of Expasy.
- **b**SP, signal peptide determined with the use of SignalP v.2.0.
- **c**TMH, transmembrane helices determined with the use of TMPred.
- **d**Determined with MotifScan. NA, not available; CP, coupling protein; CTS, core transmembrane subcomplex.
- **e**Determined with BLASTp.
- **f**Protein name according to reference 51.
- **g**Determined with LipoP.
regions, as well as orf35 from the pCTX-M3 leading region and orf46, which is located downstream of trbC, was constructed (see Table S1 in the supplemental material) by replacing a given gene with the cat gene (27). To avoid a difference in expression levels depending on the position of the gene in the operon, the cat gene with its own promoter sequence was inserted in the opposite orientation to that of the tra or trb genes. For each pCTX-M3 deletion derivative, the frequency of conjugative transfer (conjugation efficiency) from E. coli BW25113 cells to the recipient E. coli JE2571Rifr cells, both in liquid and on filters, was determined (Fig. 2A). Liquid mating occurred at a lower efficiency than filter mating; therefore, further conjugative transfer analysis of the pCTX-M3 derivatives was performed using filter mating only. All the plasmids displayed a diminished or completely inhibited conjugative transfer ability except for pCTX-M3 orf35:cat, pCTX-M3 orf36:cat, and pCTX-M3 orf46:cat; these plasmids showed the same conjugation efficiency as the parental pCTX-M3 plasmid, which was approximately \(10^{-7}\) for filter mating (Fig. 2B). We therefore concluded that orf35, orf36, and orf46 are dispensable for conjugative transfer.

**Complementation of the deleted tra and trb genes.** To exclude the possibility that the reduced conjugative transfer of the pCTX-M3 deletion derivatives was caused by a polar effect, each mutated plasmid was complemented with an appropriate gene cloned into pMT5 or pAL3 under the control of the \(P_{lac}\) promoter (Table S1). For the
The majority of the pCTX-M3 derivatives, the complementation fully or at least partially restored the conjugation efficiency ($10^{-4}$ to $10^{-1}$ per donor cell). However, for the *nikB*, *traM*, *traN*, *traU*, *traW*, and *traY* deletion derivatives, the presence of the complementing gene rescued the conjugation efficiency only to levels less than $10^{-5}$; for *traP* and *pri*, the complementation had no detectable effect (Fig. 2B). Although several independent *traP* and *pri* deletion mutants were investigated, none regained the conjugative transfer ability upon complementation with pMT*straP* or pAL*pri*, respectively.

In R64, the disruption of *nikB*, which encodes nickase (relaxase), abolished conjugative transfer; however, *nikB* can be complemented well (28). Furthermore, the disruption of *traM*, *traN*, *traW*, and *traY* was detrimental to the transfer of R64, but these genes were able to be complemented by plasmids expressing the relevant gene to the wild-type (WT) transfer efficiency or, for *traY*, 10× lower than the WT transfer efficiency (12). In R64, disruptions of *traU* are detrimental to transfer. Similar to the effect of limited complementation of pCTX-M3*traU::cat*, the transfer efficiency of R64 that expressed a *traU* gene disrupted close to its start site but that was complemented by a plasmid bearing *traU* was $9 \times 10^{-7}$ (12). Taking into account the differences in complementation between the respective mutants of pCTX-M3 and R64, the *nikB*, *traM*, *traN*, *traW*, *traY*, *pri*, *traP*, and *traU* mutants of pCTX-M3 were analyzed further. First, the expression of the complementing genes was verified by real-time quantitative PCR (RT-qPCR). The transcript levels of all these genes were at least 10-fold higher than those found in the strain bearing pCTX-M3 (Fig. 3A).

Given that a polar effect of the deletion of those genes was not strictly excluded, the transcription of genes located directly downstream of the target genes was quantified by RT-qPCR (Fig. 3B). Indeed, the transcript levels of each of the downstream genes analyzed, except for those of the *traN::cat* and *traU::cat* derivatives, were lower than those found in the strain bearing the intact pCTX-M3 (Fig. 3B). The complementation of the analyzed deletions with the appropriate genes restored the expression of the downstream genes to the levels observed in the strain with pCTX-M3 for the *pri*, *traM*, and *traP* deletion mutants and nearly restored expression for the *traU* deletion mutant, but complementation had only a small effect on the *nikB::cat* and *traW::cat* mutants (Fig. 3B). In the case of the *traY::cat* mutant, where no decrease in the expression of the downstream gene was found, the complementation had no effect on the transcript level of the downstream gene.

The results so far showed that five deletion mutants of pCTX-M3 behaved in an unexpected manner: *traP* and *pri*, which could not be complemented; *traY*, where the

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**FIG 3** Transcript levels of genes in the pCTX-M3 plasmid deletion derivatives. (A) Transcript levels of the complementing genes (in bold) in the respective deletion derivatives. (B) Transcript levels of genes (in bold) located directly downstream of the deleted gene in the respective pCTX-M3 deletion derivatives (light gray) complemented with the appropriate plasmid (dark gray). Each result is the mean value from biological triplicates normalized to the transcript level of the appropriate gene in the BW25113(pCTX-M3) strain. Error bars indicate the SDs.
Complementation was very poor; and nikB and traW, where the downstream genes, traH and traX, respectively, showed low expression in addition to poor complementation. To determine whether the behavior of these genes was due to the deletion of the genes in question or was connected with the presence of the cat gene in their loci, the cat gene was eliminated from the mutants with the use of the pCP20-encoded FLP recombinase. The ability of the constructed plasmids (pCTX-M3ΔtraP, pCTX-M3Δpri, pCTX-M3ΔtraW, pCTX-M3ΔtraY, and pCTX-M3ΔnikB) to undergo conjugative transfer was tested after complementation with a plasmid bearing the appropriate genes (dark gray) in filter mating. vec, empty vector; #, undetectable transfer (<10^-7). Each result is the mean from at least five experiments. Error bars indicate the SDs.

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However, the transfer efficiency of pCTX-M3Δpri after complementation was low (approximately 2 × 10^-7). The results obtained for the pri deletion in pCTX-M3 differ from those for the sog deletion in R64, where the deletion resulted in only a small drop in the transfer efficiency, from 10^-2 to 10^-3 (29). The sog gene encodes two proteins, the SogL primase (1,255 amino acids [aa]), which generates RNA primers for plasmid replication, and the shorter SogS (844 aa), which is a product of translational reinitiation within the sog reading frame (30). It was speculated that Sog proteins create a complex with DNA, coating the transferred single DNA strand to protect and stabilize it (31). Both proteins are transported from the donor to the recipient cell during conjugation, and the transport has been shown to rely on the pil genes encoding thin pili (32), which are absent in pCTX-M3 (14). The pri gene of pCTX-M3 also has the potential to encode two proteins, a primase (1,070 aa) and a putative 689-aa protein comprising the C-terminal moiety of the primase. The high level of the pri transcript (Fig. 3A) might result in the production of a large amount of these two proteins, which can coat the single-stranded DNA (ssDNA) of the transferred plasmid and block the transporter.
However, the reason for the lack of complementation of the pri-deficient pCTX-M3 is highly speculative, and the pCTX-M3 pri gene therefore needs further study.

The results presented above demonstrate that though the disruptions in the majority of the tra and trb genes of the pCTX-M3 plasmid, which were performed via cat insertion, do not prevent the expression of a functional conjugative transfer system, in some cases, the deletion of the inserted cat gene was required. In these cases, the observed decreased transcript levels and lower conjugation efficiencies of the mutated plasmids even in the presence of the appropriate complementing genes suggest a defective regulation of gene expression. The mechanism of the regulation of the pCTX-M3 tra and trb operons is unknown and needs further research.

Putative roles of the tra and trb genes of pCTX-M3. (i) Putative components of the T4CP subcomplex. The analysis of the trb region of pCTX-M3 showed that the deletion of either trbA, trbB, or trbC abolished conjugation. In R64, the trbA and trbC genes were found to be indispensable for conjugative transfer, while trbB was required for a high transfer efficiency (33). TrbC\textsubscript{pCTX-M3}, with its ATPase Walker motifs A and B, is homologous to TrbC\textsubscript{R64} and to DotL (IcmO) of \textit{L. pneumophila}, which acts as a type IV coupling protein (T4CP) (Table 1). In the Dot/Icm system, DotL forms the T4CP subcomplex along with other proteins that have no CP function: two inner membrane proteins, namely, DotM (IcmP) and DotN (IcmJ), and the secretion adapter proteins IcmS and IcmW (34). The DotM homologue of pCTX-M3 is encoded by the traB gene. Homologues of TrbB\textsubscript{pCTX-M3}, except for the TrbB proteins encoded by IncI plasmids, are putative disulfide bond isomerases. Therefore, the trbC gene of pCTX-M3 is likely to encode the coupling protein, while trbA codes for an element of the T4CP subcomplex, whose other components are to be characterized.

(ii) Putative components of the transmembrane subcomplex. The traN gene of both R64 (12) and pCTX-M3 encodes a homologue of DotH (IcmK) of \textit{L. pneumophila}. The proper localization of DotH in the OM is assisted by the lipoproteins DotC and DotD, which together form a pore similar to that formed by the VirB7/VirB9 proteins of \textit{A. tumefaciens} (13). DotC, DotD, and DotH, along with the IM proteins DotF (IcmG) and DotG (IcmE), have been found to form the core transmembrane subcomplex that bridges the IM and the OM in \textit{L. pneumophila} (13, 35). A homologue of \textit{L. pneumophila} DotC, encoded by traI, is indispensable for the conjugative transfer of pCTX-M3, while the disruption of traI\textsubscript{R64} led only to a reduction in the transfer efficiency (12). In pCTX-M3 and R64, TraH is a homologue of DotD of \textit{L. pneumophila}; however, in contrast to the deletion of traH\textsubscript{R64} (12), the disruption of traH\textsubscript{pCTX-M3} abolishes the transfer of pCTX-M3. The putative localization of TraH\textsubscript{pCTX-M3} in the cell membrane is supported by the presence of a predicted signal peptide and a lipid attachment motif in its sequence (Table 1). Distant homologues of DotF of \textit{L. pneumophila} are TraP\textsubscript{pCTX-M3} and TraP\textsubscript{R64}, which are necessary for the conjugative transfer of both plasmids. In the TraP\textsubscript{pCTX-M3} sequence, a single transmembrane helix was found, suggesting IM localization (Table 1). DotG of \textit{L. pneumophila} shares homology with TraO\textsubscript{R64} (13) and TraO\textsubscript{pCTX-M3} (Table 1) homologues of VirB10 of \textit{A. tumefaciens}. Interestingly, the deletion of the traO gene abolished the conjugative transfer of pCTX-M3, while in R64, traO deletion only reduced the transfer efficiency (12). Therefore, we propose that TraH, TraI, TraN, TraO, and TraP of both pCTX-M3 and R64 are components of the core transmembrane subcomplex. The different consequences of the deletion of traH, traI, or traO on the conjugative transfer of pCTX-M3 and R64 raise the possibility that the compositions of the core transmembrane subcomplexes encoded by these two plasmids are also different.

(iii) Putative functions of other pCTX-M3-encoded proteins. The nikA and nikB genes encode components of the nickase complex (an auxiliary protein and a relaxase, respectively). The deletion of these genes completely abolishes the conjugative transfer of pCTX-M3 and R64 (36). The traJ gene encodes an ATPase homologous to VirB11 of \textit{A. tumefaciens} and DotB of \textit{L. pneumophila} (37) (Table 1). Its deletion abolishes the conjugative transfer of pCTX-M3, while in R64, traJ deletion only reduced the transfer efficiency (12). In turn, TraK\textsubscript{R64} and TraK\textsubscript{pCTX-M3} (Table 1) are homologues of the IcmT
protein of L. pneumophila, whose function is unknown (37). TraM_{R64} and TraM_{CTX-M3} display homology with DotI (IcmL) of L. pneumophila and with VirB8 of A. tumefaciens (11, 38). The traU gene encodes a putative ATPase that is a homologue of DotO (IcmB) of L. pneumophila (37, 39). The encoded protein is also distantly homologous to VirB4 (40), which is involved in pilus assembly (40) and is essential for the virulence of Agrobacterium (41).

The traR and traQ genes, which are indispensable for the conjugative transfer of pCTX-M3 and R64 (12), code for proteins distantly homologous to each other (Table 1) and that belong to the VirB2 family, which forms the major T-pilus subunit in the A. tumefaciens VirB/D4 system (11). The product of traY is a distant homologue of DotA of L. pneumophila (37) and, together with ExcA, builds the entry exclusion system of R64 (42) and of pCTX-M3 (17). The putative functions of the other proteins encoded in the tra and trb regions of pCTX-M3 remain unknown (Table 1).

**Putative regulators of the pCTX-M3 transfer genes.** To identify the pCTX-M3 genes affecting the mobilization efficiency of plasmids bearing oriT_{pCTX-M3}, we compared the ability of E. coli cells carrying pCTX-M3, pCTX-M3orf35::cat, pCTX-M3orf36::cat, or pCTX-M3orf46::cat plasmids to mobilize the pToriT plasmid into the recipient E. coli J2571Rfr^+ cells (Fig. 5A). The helper plasmid pCTX-M3orf35::cat was >200-fold more effective in aiding pToriT mobilization than the other plasmids (2.76 x 10^{-1} versus 1.25 x 10^{-3}, respectively; P = 0.0042). Surprisingly, pCTX-M3orf36::cat was approximately 5 times more effective than the WT plasmid (3.61 x 10^{-3} versus 1.25 x 10^{-3}, respectively; P = 0.0124). The protein product of orf35 of pCTX-M3 exhibits 44% amino acid sequence similarity with that of yggA, the first gene in the leading region of R64 (see Fig. S1), whose involvement in mobilization has not been studied.

With A. tumefaciens as the recipient, E. coli carrying pCTX-M3orf35::cat or pCTX-M3orf36::cat was >100- or 10-fold more efficient, respectively, as a pToriT donor than was E. coli bearing pCTX-M3 or pCTX-M3orf46::cat (Fig. 5B). However, only the increase in mobilization efficiency in the presence of pCTX-M3orf35::cat was statistically significant (P = 0.0024).

To further analyze the effects of orf35 and orf36 on plasmid mobilization, the deletions of these genes were complemented with appropriate plasmids, namely,
orf35 and orf36, respectively. The pABBoriT plasmid was mobilized to an E. coli recipient from E. coli DH5α/H9251 donors. The helper plasmids pCS, pC35S, and pC36S, which were derived from pCTX-M3 orf35::cat and pCTX-M3 orf36::cat, respectively, and lack kanamycin resistance, were generated for use with the Kmr pABBoriT plasmid (Table 2).

In the presence of pC35S and the complementing pAL orf35 plasmid, the mobilization efficiency of pABB20oriT was slightly reduced relative to that observed in the presence of pC35S and the pAL3 vector (Fig. 6A), but this reduction occurred only when freshly obtained transformants were used as donors and the experiment was performed at 28°C. It is worth noting that the growth of E. coli bearing both pCTX-M3 and pAL orf35 was disturbed, while the presence of pAL orf35 alone did not affect cell growth. This effect can result from the possible deregulation of the tra genes controlled by the product of the orf35 gene, especially when expressed from the two coresident plasmids.

The complementation of the orf36 mutation in pC36S by pAL orf36 decreased the pABB20oriT mobilization efficiency even below the level obtained with pCS as the helper plasmid (Fig. 6B). Interestingly, in the strain bearing both pCS and pAL orf36, the mobilization efficiency of pABB20oriT was reduced.

To address the question of the role of orf35 and orf36 in conjugative transfer, the transcript levels of the nikA, nikB, and traH genes, the first three genes of the tra operon, respectively.

### TABLE 2 Strains and plasmids used in the study

| Strain or plasmid | Relevant feature or construction description | Source or reference |
|-------------------|---------------------------------------------|---------------------|
| **Strains** | | |
| Escherichia coli | | |
| DH5α | ∆lacZΔM15 deoR endA1 gyrA96 hsdR17 recA1 relA1 supE44 thi-1 ΔlacZYA ArgF U169 | 52 |
| BW25113 | Δ[araD-araB]567 Δ[araD-araB]567 ΔlacZ4787::rrnB-3 hsdR174 rph-1 | 27, 53 |
| JE2571 | leu thi thi lacY thy pil fla | 25 |
| JE2571Rif | JE2571 selected on LB plus rifampin | This work |
| Agrobacterium tumefaciens | | |
| LBA1010 | Rif<sup>a</sup> | 54 |
| Pseudomonas putida | Rif<sup>a</sup> gfp Km<sup>a</sup> | 55 |
| KT2442 | | |
| Ralstonia eutropha JMP228 | Rif<sup>a</sup> | 56 |
| **Plasmids** | | |
| pCTX-M3 | IncM plasmid, 89,468 bp, Ap<sup>r</sup> Pf<sup>a</sup> Cz<sup>f</sup> Km<sup>r</sup> Gen<sup>r</sup> To<sup>r</sup> | 14, 15 |
| pACYC184 | Vector (ori<sub>V</sub><sup>p15A</sup> Tcr Cmr) | 57 |
| pAL3 | pUC18 BstUI fragment containing the lacZ gene and MCS, cloned into Scal-PvuII pACYC184 (ori<sub>VV</sub><sup>p15A</sup> Tcr) | This work |
| pBBR1 MCS-2 | Vector (ori<sub>VBBR1</sub><sup>pCTX-M3</sup> Cmr) | 58 |
| pKD3 | Template for generation of the PCR products used in gene disruption, pir-dependent replicon (ori<sub>V</sub><sup>pir</sup> Ap<sup>r</sup> Cm<sup>r</sup>) | 27 |
| pKD46 | Lambda Red recombinase expression plasmid, repA101(Ts) (ori<sub>VV</sub><sup>pir</sup> Ap<sup>r</sup>) | 27 |
| pCP20 | FLP recombinase expression plasmid, repA101(Ts) (ori<sub>VV</sub><sup>pir</sup> Ap<sup>r</sup> Cm<sup>r</sup>) | 59 |
| pMT5 | pACYC184 SspI-MstI fragment containing a gene for Tc<sup>r</sup> cloned into Dral-SspI pUC18 (ori<sub>V</sub><sup>pirB1</sup> Tc<sup>r</sup>) | This work |
| pUC18 | Cloning vector (ori<sub>V</sub><sup>pirB1</sup> Ap<sup>r</sup>) | 60 |
| pABB20 | Cloning vector (ori<sub>V</sub><sup>pirB1</sup> Km<sup>r</sup>) | 61 |
| pOrIT | ori<sub>TT</sub><sup>pCTX-M3</sup> (nucleotides 31,616–31,721) cloned into the pUC18-derived pM13 vector (ori<sub>V</sub><sup>pirB1</sup> Cm<sup>r</sup>) | 14 |
| pALorIT | pOrIT EcoRI-PstI fragment containing ori<sub>TT</sub><sup>pCTX-M3</sup> cloned into EcoRI-PstI pAL3 (ori<sub>V</sub><sup>pir</sup> Tc<sup>r</sup>) | This work |
| pBBortIT | pALorIT XbaI-PvuII fragment containing the tetracycline resistance gene and ori<sub>TT</sub><sup>pCTX-M3</sup> cloned into PvuI-XbaI pBBR1 MCS-2 (ori<sub>V</sub><sup>pirB1</sup> Tc<sup>r</sup>) | This work |
| pTorIT | pBBortIT derivative, fragment BsaI-Bst1107I with MOR<sup>pir</sup> removed (ori<sub>V</sub><sup>pirB1</sup> Km<sup>r</sup> Tc<sup>r</sup>) | This work |
| pABB20orIT | pOrIT BamHI-PstI (blunted) fragment containing ori<sub>TT</sub><sup>pCTX-M3</sup> cloned into BamHI-PstI pABB20 (ori<sub>V</sub><sup>pirB1</sup> Cm<sup>r</sup>) | This work |
| pHS11 | pCTX-M3 derivative containing SexAI-SnaBI (nucleotides 36,645–40,568) and NruI-HindIII (nucleotides 51,663–58,653) fragments | This work |
| pCS | pCTX-M3 largest SalI fragment (nucleotides 1–595,520 and 79,940–89,486) self-ligated, Cfl<sup>r</sup> Cm<sup>r</sup> | This work |
| pC35S | pCTX-M3orf35::cat largest SalI fragment self-ligated, Cfl<sup>r</sup> Cm<sup>r</sup> | This work |
| pC36S | pCTX-M3orf36::cat largest SalI fragment self-ligated, Cfl<sup>r</sup> Cm<sup>r</sup> | This work |

<sup>a</sup>Azt, aztreonam; Cft, cefotaxime; Caz, ceftazidime; Gen, gentamicin; Pi, piperacillin; To, tobramycin; (Ts), thermosensitive replication.
were determined in *E. coli* strains bearing either pCTX-M3*orf35*::cat or pCTX-M3*orf36*::cat and were compared with those in the control strain bearing intact pCTX-M3 (Fig. 6C and D). In the strain bearing pCTX-M3*orf35*::cat, the transcript levels of all three genes were elevated, approximately 40-, 23-, and 80-fold for *nikA*, *nikB*, and *traH*, respectively, relative to those in the control strain. In the strain bearing pCTX-M3*orf36*::cat, the levels of the *nikA* and *nikB* transcripts were unchanged, while the *traH* transcript was approximately 120-fold more abundant than in the control strain.

We propose that the pCTX-M3 *tra* operon, which encodes both the nickase complex and the T4SS, is subject to *orf35*-dependent repression. The effect of derepression of the *tra* operon in pCTX-M3*orf35*::cat cells would be visible for mobilizable multicopy plasmids, while the conjugation ability of the low-copy-number pCTX-M3*orf35*::cat would not benefit from the derepression of *tra* due to the limited number of accessible oriT-bearing plasmid molecules.

Similarly, the deletion of *orf36*, which is unique to IncL and IncM plasmids (14) and is dispensable for conjugation, increased the mobilization efficiency into *E. coli* and upregulated *traH* but not *nikA* or *nikB*. Moreover, the presence of additional copies of *orf36* significantly impaired mobilization even in donors bearing the native pCTX-M3 conjugative transfer region. We propose that the expression of *traH* and probably also that of the downstream genes encoding the T4SS are additionally regulated by the *orf36* product in a manner independent of *nikAB* transcription. The mechanism underlying the *Orf35*- and *Orf36*-dependent regulation is currently unknown and deserves further study, especially given that these predicted proteins do not contain known DNA-binding motifs (Table 1).

**FIG 6** Effect of *orf35* and *orf36* on the efficiency of pABB20oriT mobilization. (A and B) Mobilization efficiency of pABB20oriT into *E. coli* recipient cells by respective helper plasmids: pCS (pCTX-M3 devoid of all antibiotic resistance genes except *bla*TEM-1 and *bla*CTX-M-3) and its derivatives pC35S (A) and pC36S (B) in the presence of pAL3 (vector), pAL*orf35* (*orf35*), or pAL*orf36* (*orf36*). (C and D) Relative transcript levels of *nikA*, *nikB*, and *traH* in *E. coli* strains bearing pCTX-M3 deletion derivatives pCTX-M3*orf35*::cat (C) or pCTX-M3*orf36*::cat (D). The transcript levels were normalized to those in cells bearing intact pCTX-M3. Each result is the mean from at least four experiments. Error bars indicate the SDs.
Earlier studies (26) have demonstrated that pCTX-M3 can be transferred to *A. tumefaciens* via conjugation. Unexpectedly, despite a number of attempts, we were unable to transfer pCTX-M3 from *E. coli* to *A. tumefaciens* by mating (Fig. 5B). However, our mobilization experiments demonstrated that the conjugation system of pCTX-M3 is highly efficient in transferring the mobilizable broad-host-range (*oriV*<sub>pBBR1</sub>) plasmid pToriT, which contains *oriT*<sub>pCTX-M3</sub>, into *A. tumefaciens* (10<sup>5</sup>/H11002 transconjugants per donor after 30 min of mating).

The finding that pCTX-M3 is not transferred into *A. tumefaciens* is inconsistent with previous results (26) showing that the conjugative transfer efficiency of the entire pCTX-M3 into Alpha-, Beta-, and Gammaproteobacteria was on the order of 10<sup>−5</sup> transconjugants per donor cell after 24 h of mating. To investigate this discrepancy, we performed a 24-h mating experiment using *E. coli* DH5α(pCTX-M3, pToriT) as the donor and *E. coli*, *A. tumefaciens*, *R. eutropha*, and *P. putida* as recipients. In such a system, the transfer of pCTX-M3 during mating reflects the host range of both its conjugation system and the IncM replicon, while the transfer of pToriT, the mobilizable broad-host-range plasmid containing *oriT*<sub>pCTX-M3</sub>, reflects the host range of the conjugation system of pCTX-M3 only. Transconjugants carrying pCTX-M3 were selected on plates containing gentamicin and rifampin, while those with pToriT were selected on plates with tetracycline and rifampin (*A. tumefaciens, R. eutropha, and P. putida*) or with kanamycin and rifampin (*A. tumefaciens* and *P. putida*). As shown in Fig. 7, the pCTX-M3 conjugation system is highly efficient in transferring pToriT into *A. tumefaciens* and *R. eutropha* (2 × 10<sup>−3</sup> and 3 × 10<sup>−4</sup> transconjugants per donor cell, respectively) and, with a lower efficiency, also into *P. putida* (3 × 10<sup>−6</sup> per donor cell). In contrast, pCTX-M3 transconjugants were obtained only in the *E. coli* recipient. Thus, pCTX-M3 itself, when transferred, cannot be established in *A. tumefaciens, R. eutropha*, or *P. putida*. These results indicate that the host ranges of the pCTX-M3 conjugative transfer system and its replicon differ markedly; the former shows a broad range comprising *Alpha-, Beta-, and Gammaproteobacteria*, and the latter is restricted to *Enterobacteriaceae*. A similar observation concerning the differences between the host ranges of the conjugation system and the replicon has been reported previously for the narrow-host-range mobilizable *Klebsiella pneumoniae* plasmids pGMS31 and pGMS32 (43).
plasmids replicate only in *Gammaproteobacteria*, but their mobilization systems enable the conjugative transfer of a heterologous replicon into several *Alphaproteobacteria* hosts by the RK2 (IncP1α) conjugation system (43). In addition, recently, different host ranges of the conjugative and the replicative systems have been shown for the self-transferable *P. putida* plasmid NAH7 of the IncP-9 group (with the MPF3 system) (44).

Concluding remarks. Although the conjugation system of pCTX-M3 belongs to the MPF1 group, it differs from the one encoded by IncI1 plasmids. Therefore, it would be valuable to reevaluate the mobilization host range of MPF1 systems. It has been shown that ssDNA transiently generated during conjugative transfer triggers the SOS response in recipient cells unless the plasmid codes for an anti-SOS factor (45). As a consequence, homologous recombination and integron integrase genes are induced, leading to DNA rearrangements (45). Therefore, plasmids with an MpF host range broader than their replicon host range, such as pCTX-M3, which does not code for an anti-SOS factor but does bear an integron, may have a greater impact on the adaptability of bacterial populations than previously appreciated.

**MATERIALS AND METHODS**

**Bacterial strains, plasmids, and growth conditions.** The strains used in this work are listed in Table 2. *E. coli* DH5α was used as the host strain for DNA cloning. In the mating experiments, *E. coli* strain BW25113 or, where stated, strain DH5α bearing pCTX-M3 and its derivatives, was used as the donor, and *E. coli* strain JE2571RIF was used as the recipient. In transspecies matings, *Pseudomonas putida*, *Ralstonia eutropha*, or *Agrobacterium tumefaciens* was used as the recipient. The bacteria were cultured with agitation in LB medium (Biocorp, Warsaw, Poland) or on agar-solidified LB plates (46) at either 37°C (*E. coli* and *P. putida*) or 28°C (*A. tumefaciens* and *R. eutropha*). When required, antibiotics were added to the medium at the following final concentrations (μg/ml): ampicillin, 100; chloramphenicol, 20; gentamicin, 50; kanamycin, 50; rifampin, 100; tetracycline, 20 or 6 (for pTorIT selection).

**Cloning and DNA manipulation.** Plasmid DNA was isolated by alkaline lysis using Plasmid Mini or Plasmid Midi kits (A&A Biotechnology, Gdańsk, Poland) according to the manufacturer’s instructions. DNA cloning was performed according to standard protocols (46). All the enzymes used for cloning were from MBi Fermentas/Thermo Scientific (Vilnius, Lithuania).

**Plasmid constructions.** The plasmids that were constructed and used in this study are listed in Table 2 and Table S1 in the supplemental material. pCTX-M3 derivatives with deletions of genes from the tra and trb regions were constructed through lambda Red-mediated recombination (27). First, the BW25113(pKD46) strain was electrotransformed with pCTX-M3 and was selected on LB agar plates with gentamicin at 28°C. PCR products comprising the cat gene sequence with extensions homologous to the gene to be replaced on the pKD3 plasmid template were obtained using the primers listed in Table S2. Then, BW25113(pCTX-M3, pKD46) cells were electrotransformed with these DpnI-treated PCR products and were selected on LB agar plates with chloramphenicol at 37°C (to avoid the propagation of pKD46, which shows temperature-sensitive replication). Single colonies were isolated by the streak plate method at 37°C, and the correct integration of the cat gene into the target gene was verified by PCR (35 cycles) with the primer pairs listed in Table S3 and S4. The integration of cat in the four longest genes (nikB, pri, trbU, and trbR) was further analyzed by multiplex PCR with three primers (see Table S5); (i) catReVer, which anneals to the cat gene, (ii) a primer located upstream of the deleted gene (nikAF, priUVer, traUsU, or traYsU, respectively), and (iii) a primer that anneals to the gene to be deleted (nikBDVer, priDVer, traUDVer, or traYDVer, respectively). The primers were designed so that the expected products were smaller than 1 kb and enabled discrimination between the native pCTX-M3 and the appropriate mutant plasmid. All mutated plasmids were verified by sequencing with the catU142 primer. The loss of pKD46 was checked by multiplex PCR with the repKD46F and repKD46R primers, which were designed to amplify the repA101 (thermosensitive replication) gene fragment, and with the TEMfor and TnTEMrev primers (for amplification of the baαTnα Rev gene, which is present in both pCTX-M3 and pKD46) as an internal PCR control (see Table S4). The constructed plasmid derivatives are listed in Table S1.

The cat gene was eliminated from six pCTX-M3 derivatives, namely, pCTX-M3::cat, pCTX-M3::trbR::cat, pCTX-M3::trbU::cat, pCTX-M3::trbA::cat, and pCTX-M3::trbY::cat, using pCP20, which encodes FLP recombinase, as a helper plasmid. In this process, the strain DH5α(pCP20) was electrotransformed with the appropriate pCTX-M3 derivative, and transformants were selected on LB agar with gentamicin at 28°C. Then, the transformants were streaked in parallel on LB with both chloramphenicol and gentamicin and were grown at 37°C. After colony purification, the clones that were chloramphenicol sensitive and gentamicin resistant were verified by PCR with the primers listed in Tables S3 and S5. The loss of pCP20 was verified by PCR with the repKD46F and repKD46R primers. The plasmids were then introduced into the BW25113 strain.

To construct the plasmids carrying individual genes from the tra and trb regions for use in the complementation experiments (Table S1), specific genes were amplified by PCR using Pfu DNA polymerase (Thermo Fisher Scientific, Waltham, MA) with the primers listed in Table S6 and were cloned into the pMTrS or pAL3 vectors, as described in Table S1. Genes which, probably due to the harmful effects of high-level expression, were not able to be cloned into the multicopy plasmid pMT5 (oriVAMB), were
cloned into the low-copy-number vector pAL3 (oriV3tica). Only the pALpri plasmid was constructed without a PCR amplification step, as indicated in Table S1. The cloned genes were verified by sequencing (primers listed in Tables S3 and S4). The expression of the complementing genes cloned into the pAL3 and pMTS vectors is driven by the lactose operon promoter (Plac).

Plasmids bearing oriTCTX-M-3 were obtained by cloning the oriT sequence into appropriate plasmids, as described in Table 2.

The plasmids pCS, pC35S, and pC36S (Table 2) were obtained by the digestion of pCTX-M3, pCTX-M3oriF33:cat, and pCTX-M3oriF36:cat, respectively, with Sall and the recircularization of the largest DNA fragment. Thus, these plasmids are devoid of all resistance genes except the blaTEM.1 and blaCTX-M-3 genes present in pCTX-M3.

PCR conditions. PCR was performed in a Veriti thermal cycler (Applied Biosystems, Foster City, CA) using DreamTaq or Pfu DNA polymerase with the supplied buffers (Thermo Fisher Scientific), deoxyribonucleotide triphosphate (dNTP) mixture, template DNA (purified DNA or bacterial colonies), and the appropriate primer pairs listed in Tables S2 to S6, according to the manufacturer’s recommendations.

DNA sequencing. The sequencing was performed in the DNA Sequencing and Oligonucleotide Synthesis Laboratory at the Institute of Biochemistry and Biophysics, Polish Academy of Sciences, using a dye terminator sequencing kit and an automated sequencer (ABI 377; PerkinElmer, Waltham, MA).

Real-time quantitative PCR. RNA was isolated from cells of BW25113 strains bearing a specific pCTX-M3 deletion derivative alone or in combination with a plasmid carrying a appropriate complementing gene in the late exponential phase of growth (optical density at 600 nm [OD600] of 0.8 to 1) using a GeneJET RNA purification kit (Thermo Fisher Scientific) according to the manufacturer’s protocol. RNA quality and integrity were checked by agarose gel electrophoresis, and the concentration was estimated using a NanoDrop ND-1000 spectrophotometer (Thermo Fisher Scientific). Three biological replicates were analyzed.

Reverse transcription was performed with random hexamer primers using the Maxima First Strand cDNA synthesis kit for RT-qPCR with dsDNase (Thermo Fisher Scientific). The specific qPCR primers used to amplify the reference (the repA gene encoding the replication initiator protein of the pCTX-M3 plasmid) and target genes are listed in Table S7. Real-time PCR was carried out using Real-Time SYBR master mix SYBR (A&A Biotechnology) in a final volume of 10 µl in the LightCycler 480 system (Roche Life Sciences, Penzberg, Germany) with an initial denaturation at 95°C for 5 min followed by 40 cycles of amplification (95°C for 10 s and 50°C for 10 s). The relative gene expression in the deletion strains was calculated and normalized to the value obtained for a strain carrying the native pCTX-M3 plasmid (47).

Plasmid conjugative transfer. Cultures of the donor and recipient strains (approximately 10⁶ CFU ml⁻¹) were grown to stationary phase in LB to stationary phase with LB medium, and the donor strain was resuspended in a volume of LB equal to the initial volume of the culture, while the recipient strain was resuspended in one-fourth of the initial culture volume. Then, 0.5 ml of the donor and recipient suspensions was mixed and filtered through a sterile Millipore HA 0.45-µm filter (Millipore, Billerica, MA). The filter was incubated on an LB plate for 24 h at 37°C or 28°C (A. tumefaciens) or 28°C (A. tumefaciens and E. coli). For the pCTX-M3 host range tests, the filter was incubated on an LB plate for 24 h at 37°C (P. putida and E. coli) or 28°C (A. tumefaciens and R. eutropha). Bacteria were washed from the filter with 1 ml of a sterile 0.85% NaCl solution. Conjugation was stopped by vigorously vortexing the mating mixture for 30 s and then placing it on ice. Serial dilutions of the mixture of donor, recipient, and transconjugant cells were plated on LB agar supplemented with the appropriate selection antibiotics. As a control, the donor and recipient cells were plated on LB supplemented with antibiotics for the transconjugant selection. Mating in liquid was performed as described above but without the use of the filter: the mating mix was incubated for 30 min at 37°C, and conjugation was stopped by vortexing for 30 s. The conjugative transfer efficiency is equivalent to the number of transconjugants per donor cell.

Bioinformatics analysis. Nucleotide sequences were analyzed using Clone Manager 9 professional edition. Sequence similarity searches were performed using the BLAST programs (48) provided by the National Center for Biotechnology Information (NCBI) (http://blast.ncbi.nlm.nih.gov/Blast.cgi) to search against the NCBI “nr” (nonredundant) DNA or protein database with standard parameters; no filters or masks were applied. The molecular weight and theoretical pl were calculated with the use of the ProtParam tool of Expasy (http://web.expasy.org/protparam/) (48). Motifs in protein sequences were searched with the use of MotifScan (http://myhits.isb-sib.ch/cgi-bin/motif_scan). SignalP v.2.0 (http://www.cbs.dtu.dk/services/SignalP-2.0/) was used to search for signal peptides, TMpred (http://www.ch.embnet.org/software/TMPRED_form.html) was used to determine the presence of transmembrane helices (TMH) (49), and LipidP (http://www.cbs.dtu.dk/services/LipoP/) was used to predict lipid attachment motifs (50).

Statistical analysis. Data concerning the plasmid conjugative transfer frequencies are presented as the means ± standard deviations (SDs). The differences between the mobilization efficiencies of pTorIT by the different plasmids (Fig. 5) were tested for statistical significance using the t test (Prism 6; GraphPad Software, Inc., La Jolla, CA). P values of <0.05 were considered statistically significant.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at https://doi.org/10.1128/JB.00234-18.

SUPPLEMENTAL FILE 1, PDF file, 0.6 MB.
ACKNOWLEDGMENTS

We thank D. Bartosik (Warsaw, Poland), C. M. Thomas (Birmingham, UK), and K. Smalla (Braunschweig, Germany) for providing bacterial strains (via G. Jagura-Burdzy) and J. Cieś (IBB PAS) for help in the RT-qPCR analysis. We acknowledge the technical assistance of M. N. Bulka (Warsaw University of Technology) and E. Bodo (Warsaw University of Life Sciences–SGGW).

This work was supported by grants PBZ-MNiSW-04/I/2007 and N N401 534640 to I.K.-Z.

We declare no conflicts of interest.

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September 2018 Volume 200 Issue 18 e00234-18