

## Analysis of ColE1 MbeC Unveils an Extended Ribbon-Helix-Helix Family of Nicking Accessory Proteins<sup>∇</sup>

Athanasia Varsaki,<sup>1</sup>†‡ Gabriel Moncalián,<sup>2</sup>† Maria del Pilar Garcillán-Barcia,<sup>2</sup>  
Constantin Drainas,<sup>1</sup> and Fernando de la Cruz<sup>2\*</sup>

Sector of Organic Chemistry and Biochemistry, Department of Chemistry, University of Ioannina, Ioannina, Greece,<sup>1</sup> and  
Departamento de Biología Molecular, Universidad de Cantabria, and Instituto de Biomedicina y Biotecnología de  
Cantabria (CSIC-UC-IDICAN), Santander, Spain<sup>2</sup>

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**MbeC is a 13-kDa ColE1-encoded protein required for efficient mobilization of ColE1, a plasmid widely used in cloning vector technology. MbeC protein was purified and used for in vitro DNA binding, which showed that it binds specifically double-stranded DNA (dsDNA) containing the ColE1 *oriT*. Amino acid sequence comparison and secondary structure prediction imply that MbeC is related to the ribbon-helix-helix (RHH) protein family. Alignment with RHH members pointed to a conserved arginine (R13 in MbeC) that was mutated to alanine. The mutant MbeC(R13A) was unable to bind either single-stranded DNA or dsDNA. Limited proteolysis fragmented MbeC in two stable folding domains: the N-terminal domain, which contains the RHH motif, and the C-terminal domain, which comprises a signature shared by nicking accessory proteins. The results indicate that MbeC plays a similar role in conjugation as TraY and TrwA of plasmids F and R388, respectively. Thus, it appears that an extended, possibly universal mechanism of DNA conjugative processing exists, in which *oriT*-processing is carried out by relaxases assisted by homologous nicking accessory proteins. This mechanism seems to be shared by all major conjugative systems analyzed thus far.**

Conjugation is one of the mechanisms through which horizontal gene transfer occurs in bacteria. The appearance of multiresistant bacteria brought horizontal gene transfer to the attention of scientists worldwide, since the dissemination of antibiotic resistance genes by conjugative plasmids poses serious health problems. ColE1 is mobilized by a wide range of conjugative plasmids and is the prototype of a family that comprises mobilizable plasmids found in gram-negative and gram-positive bacteria (14). Its mobilization region contains five genes (*mbeA*, *mbeB*, *mbeC*, *mbeD*, and *mbeE*), with two of them (*mbeB* and *mbeD*) entirely overlapping *mbeA* (Fig. 1A). Although the first report on ColE1 mobilization dates back almost 40 years (10), little is known regarding its mobilization properties. Other than the fact that *mbeA* encodes the ColE1 relaxase (38), only few experimental data are available about the role of the other four *mob* gene products. It was proposed that MbeB and MbeC are constituents of the ColE1 relaxosome (23); *mbeD* codes for ColE1 “entry exclusion” protein (40), whereas *mbeE* is not essential for plasmid mobilization (4). Although *mbeA* and *mbeC* genes are present in all ColE1 family plasmids, *mbeB*, *mbeD*, and *mbeE* are not conserved. This observation led to the suggestion of a common ancestral mobilization region containing only *mbeA* and *mbeC* (14). Along this line of reasoning, *mbeC* should play a pivotal role in ColE1 conjugative mobilization.

Conjugative DNA processing involves two main groups of proteins: relaxases and nicking accessory proteins. Although the need for nicking accessory proteins has been described in most conjugative and mobilizable plasmids, no sequence relatedness was appreciated for these genes as a class, despite their functional similarity. The best-characterized accessory proteins are the MOB<sub>F</sub> proteins TraY of plasmid F (TraY<sub>F</sub>) and TrwA of plasmid R388 (TrwA<sub>R388</sub>). TraY<sub>F</sub> enhances the activity of TraI<sub>F</sub> relaxase (19, 29). It binds double-stranded DNA (dsDNA) in a way that bends F plasmid *oriT*, possibly creating a single-stranded DNA (ssDNA) conformation suitable for relaxase binding (24, 39). TraY<sub>F</sub> is monomeric in solution (28, 33) and contains two tandem nonidentical ribbon-helix-helix (RHH) motifs, relating it to the RHH proteins Arc and Mnt (3). Similarly, TrwA<sub>R388</sub> enhances the TrwC<sub>R388</sub> relaxase function, as well as participating in the control of the *trwABC* operon transcription (26), and also belongs to the RHH family (27). Besides MOB<sub>F</sub> accessory proteins, the MOB<sub>P</sub> proteins NikA<sub>R64</sub>, TraJ<sub>RP4</sub>, and MobC<sub>pC221</sub> and the MOB<sub>O</sub> protein MobC<sub>RSF1010</sub> have been characterized biochemically. NikA<sub>R64</sub> binds the inverted repeat (IR) proximal to its *nic* site and induces DNA bending (15). Likewise, TraJ<sub>RP4</sub> binds within RP4 *oriT* (43) and enhances *nic* cleavage performed by TraI<sub>RP4</sub> relaxase (30). MobC<sub>pC221</sub> is essential for relaxosome formation and plasmid mobilization (35) and enhances the MobA<sub>pC221</sub> relaxase activity by binding to two sites close to *nic* (7). The MOB<sub>O</sub> protein MobC<sub>RSF1010</sub> assists in *oriT* strand opening, extending the separation on the DNA strands around *nic* and thereby increasing the efficiency of cleavage by the relaxase (42).

We demonstrate here that MbeC<sub>ColE1</sub> is a MOB<sub>HEN</sub> accessory protein, possibly belonging to the RHH family of DNA-binding proteins. Sequence comparisons revealed that

\* Corresponding author. Mailing address: Departamento de Biología Molecular, Universidad de Cantabria, C/Herrera Oria s/n, 39011 Santander, Spain. Phone: 34 942 201942. Fax: 34 942 201945. E-mail: delacruz@unican.es.

† A.V. and G.M. contributed equally to this study.

‡ Present address: IBBS Biophysics Laboratories, School of Biological Sciences, University of Portsmouth, Portsmouth, United Kingdom.

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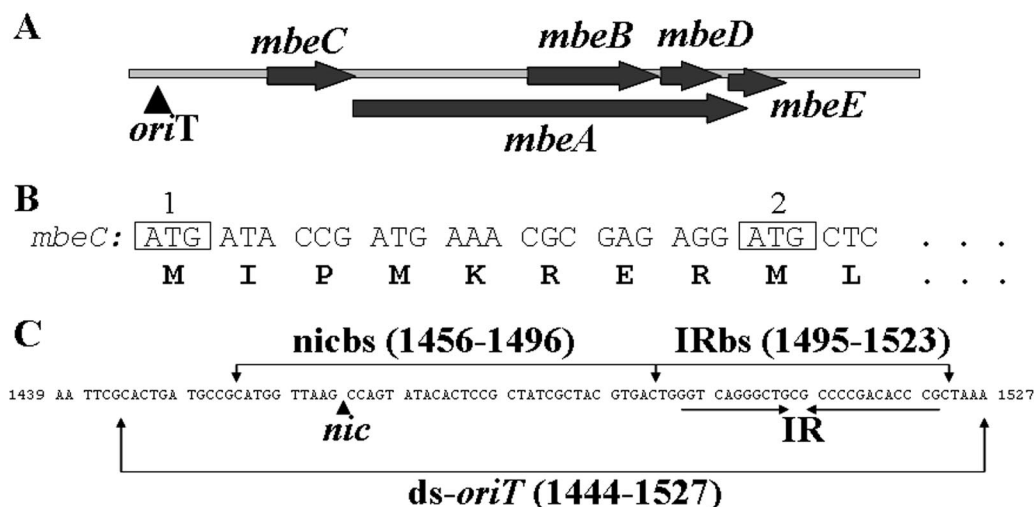


FIG. 1. (A) mobilization region of ColE1 plasmid. (B) Representation of the two AUG codons of *mbeC*. (C) *oriT* region of ColE1 plasmid. The characteristic IR is illustrated by arrows below the nucleotide sequence. The position complementary to the *nic* cleavage site is indicated with a vertical arrow, since the sequence shown is the strand complementary to the one that is nicked. The different fragments used in the EMSA (Fig. 3) are also shown.

MOB<sub>F</sub> and MOB<sub>P</sub> accessory proteins share with MOB<sub>HEN</sub> proteins the RHH DNA-binding domain. Thus, our analysis of MbeC<sub>ColE1</sub> provides a link between the different families of accessory proteins and suggests the existence of a potentially universal mechanism for accessory proteins in conjugative DNA processing.

#### MATERIALS AND METHODS

**Bacterial strains and growth conditions.** Strains used were *Escherichia coli* DH5 $\alpha$  [ $F^-$  *endA1* *hsdR17* *supE44* *thi-1* *recA1*  $\Delta$ (*argF-lacZYA*) *u160*  $\phi$ 80*dlacZ* $\Delta$ M15 *gyrA96*] (17) for standard plasmid maintenance, BL21(DE3) ( $F^-$  *ompT*  $r_B^-$   $m_B^-$ ) (36) for protein overproduction, and HMS174 (*recA1* *hsdR* Rif<sup>r</sup>) (6) as the recipient in conjugal mobilization experiments. For the maintenance and/or selection of plasmids, growth media were supplemented with antibiotics at the following concentrations: ampicillin (Am), 100  $\mu$ g/ml; kanamycin (Km), 50  $\mu$ g/ml; chloramphenicol (Cm), 25  $\mu$ g/ml; tetracycline (Tc), 5  $\mu$ g/ml; and rifampin (Rif), 100  $\mu$ g/ml.

**Plasmid constructions and genetic techniques.** Plasmids used in the present study are listed in Table 1. All PCR-generated fragments of ColE1 were synthesized by using plasmid pSU4601 DNA as a template. The identity of constructed plasmids was checked by DNA sequencing. Bacterial transformation of BL21(DE3) was carried as previously described (9). DH5 $\alpha$  competent cells were prepared (18) and transformed by electroporation (13). Mating experiments were performed as formerly reported (38).

**Oligonucleotides and labeling.** Oligonucleotides used in the present study are listed in Table 2. They were labeled at 5' end by using [ $\gamma$ -<sup>32</sup>P]ATP (300 Ci/mmol) and polynucleotide kinase (32). Unbound [ $\gamma$ -<sup>32</sup>P]ATP was eliminated from the mixture by using MicroSpin G-25 (GE Healthcare) column purification.

**Protein sequence analysis.** Protein sequences were aligned by using CLUSTAL W (22). Phylogenetic and molecular evolutionary analyses were conducted by using MEGA version 3.1 (21). The tree was inferred by using the neighbor-joining method and tested by bootstrap (1,000 replicates). The cutoff bootstrap value for condensed tree was set in 50%. Protein secondary structures were predicted by using PSIPRED (<http://bioinf.cs.ucl.ac.uk/psipred/>) (25), Jpred3 (<http://www.compbio.dundee.ac.uk/jpred>) (11), and GOR4 ([http://npsa-pbil.ibcp.fr/cgi-bin/npsa\\_automat.pl?page=npsa\\_gor4.html](http://npsa-pbil.ibcp.fr/cgi-bin/npsa_automat.pl?page=npsa_gor4.html)) (12).

**Purification of MbeC-His<sub>6</sub> and derivative mutant.** BL21(DE3) cells containing the appropriate plasmid (pUIV239 or pUIV262) were incubated in 2 liters of LB broth containing kanamycin using a micro-DCU fermentation system (B. Biotech International). Cultures were induced at an  $A_{600}$  of 0.6 with 1 mM IPTG (isopropyl- $\beta$ -D-thiogalactopyranoside). After 4 h of incubation, cells were pelleted, resuspended in buffer A (100 mM Tris-HCl [pH 7.5], 500 mM NaCl, 1 mM EDTA, 5 mM benzamidine, 0.5 mM phenylmethylsulfonyl fluoride), and lysed by sonication. No lysozyme was used, since it has a molecular weight similar to that

of the MbeC-His<sub>6</sub> and MbeC(R13A)-His<sub>6</sub> monomer when visualized after sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE). Cellular debris was eliminated after centrifugation at 45,000  $\times$  g for 30 min at 4°C. Supernatant was collected, adjusted to buffer B (50 mM Tris-HCl [pH 7.5], 200 mM NaCl, 1 mM EDTA, 20 mM imidazole), and applied to a HisTrap SP HP column (GE Healthcare). Proteins were eluted in a 20 to 500 mM imidazole gradient of buffer B. Protein samples were collected and dialyzed overnight against buffer C (50 mM Tris-HCl [pH 7.5], 1 mM EDTA, 200 mM NaCl) and further purified by HiTrap SP HP column (GE Healthcare) using a 0.2 to 1 M NaCl gradient of buffer C. Finally, proteins were purified by gel filtration through a Superdex 75 HR 10/30 (Pharmacia) using buffer D (50 mM Tris-HCl [pH 7.5], 150 mM NaCl, 1 mM EDTA). The yield was 10 mg of protein per liter of culture for both MbeC-His<sub>6</sub> and MbeC(R13A)-His<sub>6</sub> purified proteins.

**Limited trypsin digestion.** MbeC-His<sub>6</sub> protein at a concentration of 0.25 mg/ml (9  $\mu$ M) in buffer D (18  $\mu$ M) were incubated with different amounts of trypsin for 15 min at room temperature. The reaction was stopped by adding 200  $\mu$ M phenylmethylsulfonyl fluoride, and samples were stored at -80°C. Cleavage was verified by SDS-PAGE and Coomassie brilliant blue staining. Molecular masses of intact and protease-digested MbeC-His<sub>6</sub> were determined by mass spectrometry. Liquid chromatography-mass spectrometry data were acquired by using a Q-ToF micro-mass spectrometer (Waters) interfaced with a CapLC capillary chromatograph (Waters). Then, 5  $\mu$ l of each sample was loaded onto a Symmetry 300 C<sub>18</sub> NanoEase Trap precolumn (Waters) and washed with 0.1% formic acid for 5 min at a flow rate of 20  $\mu$ l/min. The precolumn was connected to an Atlantis dC18 NanoEase column (75  $\mu$ m by 150 mm; Waters) equilibrated in 5% acetonitrile and 0.1% FA. A flow splitter was used to decrease the flow rate to 0.2  $\mu$ l/min, and peptides were eluted with a 30-min linear gradient of 10 to 60% acetonitrile directly onto a NanoEase Emitter (Waters). Obtained spectra were manually analyzed by using MassLynx 4.1 software (Waters).

**DNA electrophoresis mobility shift assay (EMSA).** Increasing amounts of MbeC-His<sub>6</sub> were incubated with 10 nM radiolabeled ssDNA or dsDNA in buffer E (20 mM Tris-HCl [pH 7.5], 50 mM NaCl) and in the presence of 1 nM bovine serum albumin (BSA). Reactions (20  $\mu$ l) were carried out for 30 min at 37°C. Samples were loaded onto a 5% native polyacrylamide Bio-Rad minigel and run for 60 min at 100 V in 0.5 $\times$  TBE buffer (45 mM Tris-borate, 1 mM EDTA [pH 8.2]). Gels were vacuum dried for 2 h at 80°C and quantified with a Molecular Imager FX System (Bio-Rad). The radiolabeled dsDNA was formed by mixing equal concentration of 5'-end <sup>32</sup>P-labeled oligonucleotides with their unlabeled complementary and incubated at 95°C for 15 min. The incubator was then turned off, and the mixture was left in the incubator and allowed to cool slowly to room temperature. The dsDNAs used were ds-*oriT* (84 bp, formed by using oligo-plus and oligo-minus), nickbs (41 bp, formed by using *nic*-plus and *nic*-minus), and IRbs (29 bp, formed by using IR-plus and IR-minus) (Fig. 1C). The ssDNAs used were the oligonucleotides oligo-plus and oligo-minus (Table 2). For the competition experiments, MbeC-DNA complexes were formed as described above.

TABLE 1. Plasmids

Plasmid	Antibiotic selection	Description <sup>a</sup>	Size (bp)	Source, reference, or construction strategy
pCR2.1	Km, Am	Cloning vector	3,900	Invitrogen
pCR-blunt	Km	Cloning vector	3,500	Invitrogen
pET29c(+)	Km	Expression vector	5,372	Novagen
pUC19	Am	Cloning vector	2,686	Bioline
pSU18	Cm	Cloning vector	2,300	2
pSU4601	Km	ColE1::kan	7,930	5
R64drd-11	Tc, Sm	R64 derepressed for transfer	56,700	20
pUIV201	Am	pUC19::oriT(ColE1)	2,886	38
pUIV206	Km	pET29c::mbeA	6,974	38
pUIV230	Cm, Mob+	pSU18::mob(ColE1)	4,700	38
pUIV235	Km, Am	pCR2.1::mbeC*	4,239	This study; the 339-bp PCR fragment using the primers mobC-plus and mobC-minus cloned at pCR2.1
pUIV236	Km	pET29c::mbeC*	5,591	This study; the 317-bp NdeI-BamHI fragment from pUIV235 cloned at the corresponding sites of pET29c
pUIV238	Km, Am	pCR2.1::mbeC†	4,268	This study; the 368-bp PCR fragment using the primers mobC-P1-plus and mobC-minus cloned at pCR2.1
pUIV239	Km	pET29c::mbeC†	5,615	This study; the 341-bp NdeI-BamHI fragment from pUIV238 cloned at the corresponding sites of pET29c
pUIV241	Km, Am	pCR2.1::89-bp oriT(ColE1)	4,016	This study; the 116-bp PCR fragment using the primers plus-oriT and minus-oriT cloned at pCR2.1
pUIV244	Km	pCR-blunt::(1389-1855)ColE1	3,967	This study; the 467-bp PCR fragment using the primers plus-ΔmbeC and minus-ΔmbeC cloned at pCR-blunt
pUIV245	Cm	pSU18::89-bp oriT(ColE1)	2,338	This study; the 89-bp EcoRI-HindIII fragment from pUIV241 cloned at the corresponding sites of pSU18
pUIV247	Km	pET29c::mob(ColE1)ΔmbeC	7,267	This study; the 446-bp NdeI-XbaI fragment from pUIV244 cloned at the corresponding sites of pUIV206
pUIV248	Cm	pSU18::mob(ColE1)ΔmbeC	4,310	This study; the 2,037-bp XbaI-EcoRI fragment from pUIV247 cloned at the corresponding sites of pSU18
pUIV261	Km	pCR-blunt::mbeC(R13A)	3,866	This study; the 366-bp PCR fragment using the primers mbeC-mutR and mobC-minus cloned at pCR-blunt
pUIV262	Km	pET29c::mbeC(R13A)	5,615	This study; the 341-bp NdeI-BamHI fragment from pUIV261 cloned at the corresponding sites of pET29c
pUIV264	Km	pCR-blunt::mob(ColE1)ΔoriT	5,441	This study; the 1,941-bp PCR fragment using the primers mbeC-P1-plus and minus-ab cloned at pCR-blunt
pUIV265	Cm	pSU18::mob(ColE1)ΔoriT	4,201	This study; the 1,941-bp BamHI-EcoRI fragment from pUIV264 cloned at the corresponding sites of pSU18
pUIV266	Km	pET29c::mob(ColE1)ΔoriT	7,215	This study; the 1,941-bp NdeI-BamHI fragment from pUIV264 cloned at the corresponding sites of pET29c
pUIV267	Km	pCR-blunt::oriT(ColE1)ΔIR	3,549	This study; hybridizing the ΔIR-plus and ΔIR-minus primers and cloning the dsDNA in pCR-blunt

<sup>a</sup> \*, *mbeC* cloned from the second ATG (coordinates 1867 to 1869); †, *mbeC* cloned from the first ATG (coordinates 1843 to 1845).

After the formation of the complexes, either 38 nM (1 μg) unlabeled salmon sperm DNA (nonspecific competitor) or various amounts of unlabeled *ds-oriT* (specific competitor) was added, and the samples were incubated for 30 min at 37°C. Samples were subjected to electrophoresis and analyzed as described above.

## RESULTS

**Purification of MbeC protein.** For the *mbeC* gene, two putative AUG start codons were proposed (Fig. 1B) (8). In order to determine the actual start codon, we constructed two clones: pUIV239 (*mbeC* cloned in pET29c using the first AUG codon) and pUIV236 (*mbeC* cloned in the pET29c using the second AUG codon) (Fig. 1B). Both plasmids were tested for their ability to complement pUIV248, a Δ*mbeC* ColE1-derived plasmid. BL21(DE3) cells carrying (pUIV248 plus pUIV239) or (pUIV248 plus pUIV236) were used as donors, DH5α/R64drd-11 was used as a helper, and HMS174 was used as the recipient. Mating experiments revealed that pUIV239 complemented pUIV248 10,000 times better than pUIV236 (Table 3). This result indicates that the functional protein is expressed

from the first AUG codon. Therefore, *E. coli* strain BL21(DE3)/pUIV239 was used to overproduce MbeC-His<sub>6</sub>. SDS-PAGE of crude extracts of overproducing cells showed a major band, with electrophoretic mobility corresponding to a molecular mass of ~15 kDa (close to the 14.9 kDa predicted by sequence). MbeC-His<sub>6</sub> was purified by a three-step procedure, using affinity (His-Trap SPHP column), ion-exchange (HiTrap SPHP column), and a final gel filtration chromatography step (Superdex 75 HR 10/30 column) (Fig. 2A). As shown by gel filtration chromatography, MbeC-His<sub>6</sub> behaved as a dimer of ~30 kDa in these solution conditions.

**Determination of a minimal ColE1 *oriT*.** To determine the minimal length of ColE1 *oriT*, we constructed pUIV245 by cloning a 89-bp segment of ColE1 DNA in pSU18. The 89 bp (coordinates 1439 to 1527 of ColE1, accession no. NC\_001371) contain the *nic* site and the adjacent IR of ColE1 (Fig. 1C). We tested mobilization of pUIV245 and compared it with the mobilization capacity of pUIV201, known to be functional from previous work (38). The results showed that pUIV245 has about the same mobilization frequency as pUIV201 (Table 4),

TABLE 2. Oligonucleotides

Oligonucleotide <sup>a</sup>	Oligonucleotide sequence (5'→3') <sup>b</sup>	Location <sup>c</sup>
mobC-plus	CGC GAG <u>CAT ATG</u> CTC ACA ATA CGG	1858–1881
mobC-minus	TGA AAT TTA ACT <u>AGG ATC</u> CCG CGC C	2196–2172
mbeC-P1-plus	CTG TTC ATG GGC <u>ATA TGA</u> TAC CGA TGA	1829–1855
plus-oriT1	TAC TTT TCA TAG <u>AAT TCG</u> CAC TGA TGC	1427–1453
minus-oriT1	GGC GCG TCA <u>GAA GCT</u> TTT AGC	1542–1522
plus-ΔmbeC	CTT ACG CAT <u>CTA GAC</u> GGC ATT TCA CAC	1389–1412
minus-ΔmbeC	TCA TCG GTA <u>TCA TAT</u> GCC CAT GAA CAG	1855–1829
mbeC-mutR	GTT CAT GGG <u>CAT ATG</u> ATA CCG ATG AAA CGC GAG AGG ATG CTC ACA ATA GCG GTT ACT G	1831–1888
oligo-plus*	GCA CTG ATG CCG CAT GGT TAA GCC AGT ATA CAC TCC GCT ATC GCT ACG TGA CTG GGT CAG GGC TGC GCC CCG ACA CCC GCT AAA	1444–1527
oligo-minus*	TTT AGC GGG TGT CGG GGC GCA GCC CTG ACC CAG TCA CGT AGC GAT AGC GGA GTG TAT ACT GG/C TTA ACC ATG CGG CAT CAG TGC	1527–1444
minus-ab	CCA CTG GAT CCA CTG AAG CTG C	3769–3748
ΔIR-plus*	<u>GAA TTC</u> TTA AGC CAG TAT ACA CTC CGC TAT CGC TAC GTG ACT <u>AAG CTT</u>	1454–1502
ΔIR-minus*	<u>AAG CTT</u> AGT CAC GTA GCG ATA GCG GAG TGT ATA CTG G/CT TAA <u>GAA</u> <u>TTC</u>	1502–1454
nic-plus	CAT GGT TAA GCC AGT ATA CAC TCC GCT ATC GCT ACG TGA CT	1456–1496
nic-minus	AGT CAC GTA GCG ATA GCG GAG TGT ATA CTG G/CT TAA CCA TG	1496–1456
IR-plus	CTG GGT CAG GGC TGC GCC CCG ACA CCC GC	1495–1523
IR-minus	GCG GGT GTC GGG GCG CAG CCC TGA CCC AG	1523–1495

<sup>a</sup> \*, The slash shows the position of MbeA-dependent cleavage.

<sup>b</sup> Oligonucleotides used for site-directed mutagenesis and mutant codons are in boldface. Nucleotides introducing a restriction site are underlined.

<sup>c</sup> The numbers refer to coordinates in the ColE1 DNA sequence (GenBank accession no. NC\_001371). Base position 1 corresponds to the first T in the single EcoRI site.

indicating that 89-bp *oriT* is sufficient for ColE1 mobilization. In order to further reduce the size of the *oriT*, plasmid pUIV267 was constructed by cloning 41 bp of ColE1-*oriT* in pCR-blunt. The 41 bp (coordinates 1456 to 1496) correspond to the region including the *nic* site but not the IR (Fig. 1C). As above, the mobilization of the pUIV267 was tested. Interestingly, results showed that pUIV267 was only about 100 times less efficient than pUIV245, indicating that there is, to some degree, conjugal mobilization even with a 41-bp *oriT* containing the *nic* site but not the IR.

**MbeC binds specifically ColE1 *oriT* but does not recognize the IR.** The in vitro binding of MbeC-His<sub>6</sub> to ssDNA and dsDNA was tested. The dsDNA and ssDNA were prepared as described in Materials and Methods. Experiments revealed that MbeC-His<sub>6</sub> binds specifically to ds-*oriT* (Fig. 3A). Binding specificity was tested by adding into the reactions 38 nM un-

labeled salmon sperm DNA as a nonspecific competitor or various concentrations of unlabeled ds-*oriT* as a specific competitor (Fig. 3B). When unlabeled salmon sperm DNA was added, the MbeC-DNA complex did not dissociate, indicating that MbeC-His<sub>6</sub> binds specifically to ColE1 *oriT*. This finding is strengthened by the fact that when increasing amounts of unlabeled ds-*oriT* were added to the binding reactions, the MbeC-DNA complex dissociated and finally disappeared (Fig. 3B). No MbeC-DNA complexes were formed with ssDNA, using either oligo-plus or oligo-minus (data not shown). Given that IR elimination only reduced mobilization 100 times (Table 4), we thought that MbeC binding site could be located within the *oriT* 1456–1496 segment not containing the IR (Fig. 1C). In order to confirm this possibility, EMSA experiments were performed using the *nic*s (*nic* site fragment) and IRbs (IR-containing fragment) separately (Fig. 1C). As shown in Fig. 3C, MbeC is able to bind the *nic* site containing fragment but not the IR containing fragment, thus confirming our hypothesis.

**MbeC-like proteins belong to the family of accessory proteins for DNA processing in bacterial conjugation.** A PSI-BLAST search using MbeC\_ColE1 as a query, setting a score value of 0.00001, converged in the eighth iteration. A total of 103 homologues were recovered. When the most distantly related hits were used as queries in new rounds of BLAST search in order to find more remote homologues to MbeC, the relaxosomal proteins TraJ\_RP4 and NikA\_R64 were hit, among many of their homologues. MbeC-like proteins are generally smaller than 200 amino acids and share some overall similarity, allowing the construction of a neighbor-joining condensed tree (Fig. 4). It shows clusters of highly related homologues, while the unreliable relationship between clades is not estimated. MbeC homologues were not constrained to mobilizable plasmids but also included conjugative plasmids. Standard PSI-

TABLE 3. Complementation analysis of the ΔmbeC plasmid pUIV248<sup>a</sup>

Complementing plasmid	Mobilization frequency (SD)
pET29c (negative control)	<10 <sup>-7</sup>
pUIV248 (ΔmbeC, negative control)	<10 <sup>-7</sup>
pUIV239 (pET29c::mbeC cloned from the first ATG)	3 × 10 <sup>-2</sup> (1 × 10 <sup>-2</sup> )
pUIV236 (pET29c::mbeC cloned from the second ATG)	3 × 10 <sup>-6</sup> (1 × 10 <sup>-6</sup> )
pUS4601 (ColE1::Km, positive control)	4 × 10 <sup>-2</sup> (2 × 10 <sup>-2</sup> )
pUIV262 (R13A mutant)	1 × 10 <sup>-5</sup> (0.5 × 10 <sup>-5</sup> )

<sup>a</sup> Derivatives of *E. coli* strain BL21(DE3) carrying pUIV248 and each of the plasmids shown in the first column were separately used as donors in triparental matings using DH5α/R64drl-11 as the helper and HMS174 as the final recipient. After filter mating, bacteria were plated on selective media containing Rif to counterselect donors and Cm to select for pUIV248 mobilization. Transfer frequencies are expressed as the number of transconjugants per recipient cell and are the average of at least three separate experiments.

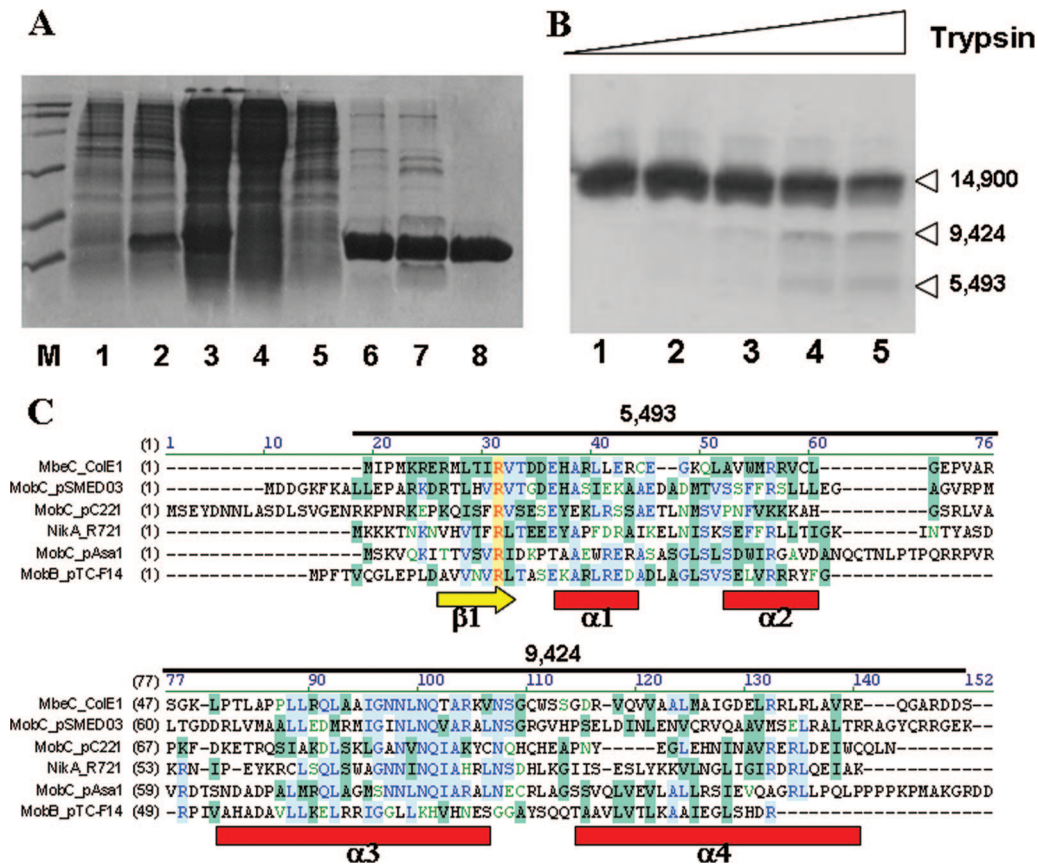


FIG. 2. (A) Purification of MbeC-His<sub>6</sub>. Fractions from the different purification steps were analyzed by SDS-12% PAGE. Lane M, molecular weight marker; lane 1, BL21(DE3)/pUIV239 before induction; lane 2, BL21(DE3)/pUIV239 after 4 h induction; lane 3, clear lysate; lane 4, flowthrough from affinity chromatography; lane 5, wash from affinity chromatography; lane 6, 6  $\mu$ g after affinity chromatography; lane 7, 5  $\mu$ g of sample after ion-exchange chromatography; lane 8, 3  $\mu$ g of sample after gel filtration chromatography. (B) Limited trypsin digestion of MbeC-His<sub>6</sub>. The results are visualized by SDS-15% PAGE. Lane 1, MbeC-His<sub>6</sub> without trypsin. Lanes 2, 3, 4, and 5 show results for 0.5, 1.5, 5, and 15  $\mu$ M trypsin, respectively. Molecular masses (in daltons) of the obtained fragments as determined by MALDI-TOF mass spectrometry are indicated on the right. (C) CLUSTAL W alignment of full-length ColE1 MbeC with other representative MOB<sub>HEN</sub> and MOB<sub>P</sub> accessory proteins. Secondary structure prediction (Jpred, PSIPRED, and GOR4) is shown below the alignment, and the length and mass of trypsin-produced fragments is shown above the alignment.

TABLE 4. Complementation analysis of the  $\Delta oriT$  plasmids pUIV265 and pUIV266<sup>a</sup>

Complementing plasmid	Mobilization frequency (SD)
Using pUIV265	
pUC19 (negative control).....	<10 <sup>-7</sup>
pUIV201 [pUC19::150 bp (ColE1-oriT)].....	5 × 10 <sup>-2</sup> (1 × 10 <sup>-2</sup> )
pUIV247 (pCR-blunt:: $\Delta$ IR).....	5 × 10 <sup>-4</sup> (2 × 10 <sup>-4</sup> )
pUS4601 (ColE1::Km, positive control).....	4 × 10 <sup>-2</sup> (2 × 10 <sup>-2</sup> )
Using pUIV266	
pSU18 (negative control).....	<10 <sup>-7</sup>
pUIV245 [pSU18::89 bp (ColE1-oriT)].....	2 × 10 <sup>-2</sup> (1 × 10 <sup>-2</sup> )
pUIV230 [pSU18::mob(ColE1), positive control].....	2 × 10 <sup>-2</sup> (1 × 10 <sup>-2</sup> )

<sup>a</sup> Derivatives of *E. coli* strain BL21(DE3) carrying pUIV265 and each of the plasmids shown under "using pUIV265" or pUIV266 and each of the plasmids shown under "using pUIV266" were separately used as donors in triparental matings using DH5 $\alpha$ /R64drl-11 as the helper and HMS174 as the final recipient. After filter mating, bacteria were plated on selective media containing Rif to counterselect donors and Cm to select for pUIV265 or pUIV266 mobilization. Transfer frequencies are expressed as the number of transconjugants per recipient cell and are the average of at least three separate experiments.

BLAST analysis initiated from NikA\_R64 hit MbeC\_ColE1 in the third round. The two sequences share 38% homology across 106 residues.

**MbeC-homologue NikA structure shows an RHH fold.** The nuclear magnetic resonance (NMR) structure of R64 plasmid NikA N-terminal fragment (Nika-N, residues 1 to 51) is deposited in the PDB databank under the accession number 2ba3. The final set contains 20 structures with two NikA-N chains. Each NikA-N chain exhibits the RHH fold, consisting of a  $\beta$ -strand formed by residues 17 to 23 ( $\beta 1$ ), followed by two  $\alpha$ -helices formed by residues 24 to 38 ( $\alpha A$ ) and 42 to 52 ( $\alpha B$ ) (Fig. 5B). Residues 1 to 16 show a different orientation on each of the 20 models, and thus we consider these residues as disorder residues in solution. A DALI structural similarity search using chain A of the NikA\_R64 NMR structure (residues 15 to 51) found the Arc repressor as the first structural homologue ( $z$ -score = 5.7, root mean square deviation over 36 residues = 1.2, and 25% sequence identity). Arc is the paradigm of the RHH-containing Arc/MetJ family of transcriptional repressors. Two Arc dimers contact each side of the

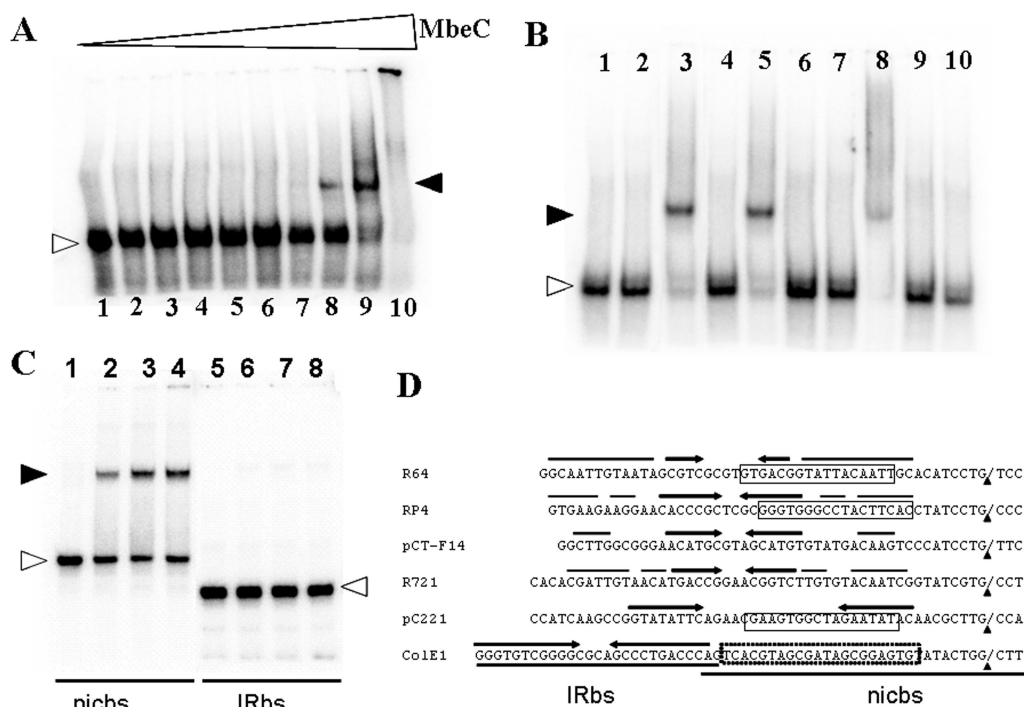


FIG. 3. (A) Gel shift of *ds-oriT* by MbeC-His<sub>6</sub> in the presence of 1 nM BSA. Lane 1, no MbeC-His<sub>6</sub> and no BSA; lane 2, no MbeC-His<sub>6</sub>. Lanes 3, 4, 5, 6, 7, 8, 9, and 10 show results for 10 nM, 30 nM, 0.1 μM, 0.2 μM, 0.4 μM, 0.6 μM, 1 μM, and 3 μM MbeC-His<sub>6</sub>, respectively. (B) Gel shift of *ds-oriT* by 1 μM MbeC-His<sub>6</sub> in the presence of 1 nM BSA, after the addition of various amounts of specific and nonspecific dsDNA. Lane 1, 10 nM *ds-oriT*; lane 2, 10 nM *ds-oriT* and 1 nM BSA; lane 3, 10 nM *ds-oriT* and 1 μM MbeC-His<sub>6</sub>, no unlabeled salmon sperm DNA; lane 4, 10 nM *ds-oriT* and 38 nM unlabeled salmon sperm DNA; lane 5, 10 nM *ds-oriT*, 38 nM unlabeled salmon sperm DNA, 1 μM MbeC-His<sub>6</sub>. Lanes 6, 7, 8, 9, and 10 show results for 10, 50, 5, 100, and 200 nM unlabeled *ds-oriT*, respectively, in 10 nM *ds-oriT* and 1 μM MbeC-His<sub>6</sub>. (C) Gel shift of IRbs and nicbs by MbeC-His<sub>6</sub>. Lane 1, nicbs without MbeC-His<sub>6</sub>; lanes 2, 3, and 4, nicbs and 0.6, 0.8, and 1 μM MbeC-His<sub>6</sub>, respectively; lane 5, IRbs without MbeC-His<sub>6</sub>; lanes 6, 7, and 8, IRbs and 0.6, 0.8, and 1 μM MbeC-His<sub>6</sub>, respectively. In all gel shifts, free DNA bands are indicated by open arrowheads, while DNA-MbeC complexes are indicated by filled arrowheads. (D) Comparison of different MOB<sub>p</sub> *oriT*s around the *nic* site. Black arrowheads indicate the position of the *nic* sites. Binding sites of Nika\_R64, TraJ\_RP4 and MobC\_pC221 are squared, and IRs are indicated by arrows. The putative MbeC binding region is indicated by a dotted-line square. The IRbs and nicbs fragments used for EMSAs in panel C are also shown.

operator (Fig. 5C), and the DNA specific contacts are made by using polar amino acids (Gln9, Asn11, and Arg13; Fig. 5D) of their N-terminal β-sheets (31). Sequence alignments showed that Nika\_R64 could be contacting DNA via the residues Val18, Thr20, and Arg22 (Fig. 5D). A polar residue could have been expected instead of Val18, but there are some RHH sequences containing a hydrophobic side chain at this position (34). Following the same line of reasoning, the MbeC\_ColE1 residues involved in DNA contact could be the polar residues Thr11, Arg13, and Thr15. According to this, the mutation R13A in MbeC should impede or reduce DNA binding.

While this manuscript was in preparation, the NMR structure of the R64 plasmid Nika N-terminal 51-residue fragment of Nika was published (41). The characterization of the three-dimensional structure of Nika(1-51) by solution-state NMR has been described, which revealed that Nika is in fact an RHH protein. Residue R22 is oriented toward the outside of the β-sheet and is responsible for binding to the *oriT*, thus confirming our “in silico” analysis.

**Mutation of R13 within the predicted β1 abolished MbeC DNA-binding activity.** In order to test MbeC(R13A)-His<sub>6</sub> functionality, the Δ*mbeC* ColE1-derived nonmobilizable plas-

mid pUIV248 was used. This plasmid can be rendered mobilizable in the presence of plasmids contributing a functional MbeC protein. Mating experiments showed that complementation by MbeC(R13A)-His<sub>6</sub> was 3,000-fold less efficient than complementation by wild-type MbeC. Therefore, the mutant MbeC(R13A) appears to be nonfunctional. For the overproduction of MbeC(R13A)-His<sub>6</sub>, BL21(DE3)/pUIV262 was used, following the same procedure as for the purification of MbeC-His<sub>6</sub>. The mutant protein behaved as a dimer in gel filtration like the wild-type MbeC. The purified MbeC(R13A)-His<sub>6</sub> was used for in vitro DNA-binding reactions, following the same procedure as described for MbeC-His<sub>6</sub>. EMSA showed that the mutant MbeC(R13A)-His<sub>6</sub> was unable to bind to ssDNA or dsDNA (data not shown).

**Identification of two MbeC domains by limited proteolysis.** The predicted tertiary structure of MbeC was probed by limited proteolysis. We observed, after trypsin treatment (Fig. 2B), two discrete bands with lower molecular masses than the full-length MbeC. Matrix-assisted laser desorption ionization-time of flight (MALDI-TOF) spectrometry gave molecular masses corresponding to two C-terminal fragments (5,493 and 9,424 Da), which together add up nicely to the determined

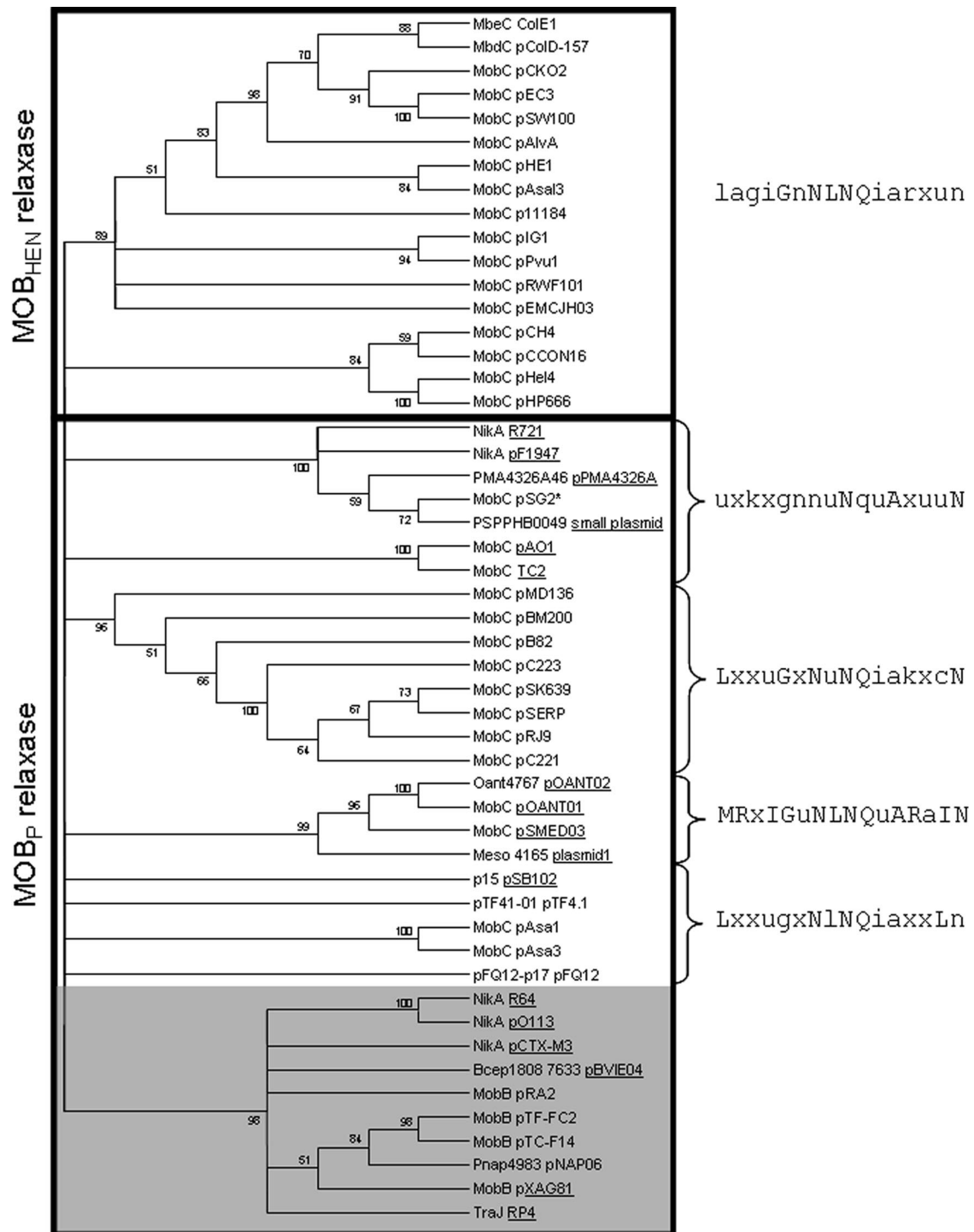


FIG. 4. Neighbor-joining condensed tree of aligned MbeC homologues. MbeC-like proteins linked to MOB<sub>HEN</sub> or MOB<sub>P</sub> relaxases are enclosed by a rectangle. The NLNQ motif associated with each clade is indicated at the right of the figure. The group of elements lacking this motif is gray shaded. When MbeC-like protein is contained in a conjugative plasmid, the element is underlined. An asterisk is placed after plasmids having a truncated relaxase.

molecular mass of the full-length protein (14,900 Da). The fragments perfectly fit the mass of the 46 N-terminal and the 51 C-terminal amino acids of the protein, respectively (Fig. 2C). The existence of this breakpoint possibly signals the interface between two domains as analyzed further in the Discussion.

## DISCUSSION

In the present study we analyzed protein MbeC encoded by plasmid ColE1 in an effort to characterize its role in ColE1 mobilization. We first determined the start codon of *mbeC* by

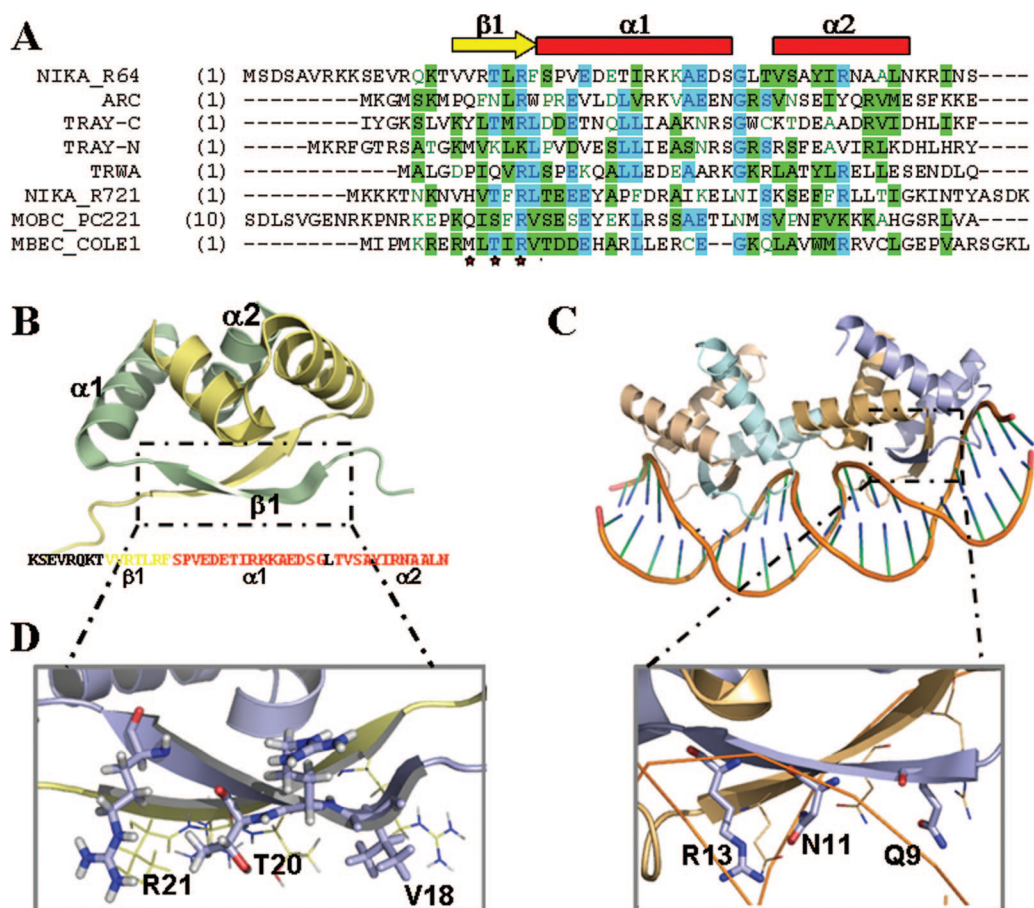


FIG. 5. (A) CLUSTALW alignment of the 50 N-terminal amino acids of MbeC with other RHH accessory proteins (TrwA\_R388, TraY\_F [N-terminal and C-terminal domain], NikA\_R721, NikA\_R64, and MobC\_pC221) and the RHH repressor Arc. The secondary structure of the solved NikA\_R64 NMR structure is shown above the alignment (the  $\beta$ -sheet is represented by a yellow arrow, and  $\alpha$ -helices are indicated by red squares). (B) Ribbon diagram representation of the NMR structure of NikA\_R64 N-terminal domain (PDB accession no. 2ba3). One monomer is shown in blue, and the other is shown in wheat color. The sequence of the solved domain is shown below the structure with a color code secondary structure (yellow  $\beta$ -sheet and red  $\alpha$ -helix forming residues). (C) Ribbon diagram representation of the crystal structure of two Arc dimers bound to its operator (PDB accession no. 1par). (D) Comparison of NikA (left) and Arc (right)  $\beta$ -sheet polar residues.

comparing the complementation properties of two potential starts (GenBank accession no. X15873, coordinates 1843 to 1845 and 1867 to 1869; Fig. 1B). Former work proposed the first AUG as the *mbeC* start codon (8), based on the criterion that a Shine-Dalgarno sequence should be found within 3 to 10 bp upstream. A different work proposed the second AUG as the start codon (4), based on a comparison of *mbeC* with the related genes *mbaC* and *mbkC* of plasmids ColA and ColK, respectively. According to our mobilization results, the conjugation frequency obtained from the MbeC started from the second AUG was 10,000 times lower compared to the MbeC started from the first AUG (Table 3), demonstrating that MbeC starts at the first AUG.

We also localized the ColE1 *oriT* to a minimal functional DNA segment. According to an earlier study (14), ColE1 *oriT* should not exceed 100 bp, given the size of the fragment that contains the *nic* site and the 32-bp upstream IR (Fig. 1C). This assumption was based on the fact that IRs are usually protein-binding sites. In the case of ColE1, the IR was not absolutely required for MbeA\_ColE1 binding to DNA, since a shift occurred using an oligonucleotide not containing the IR (38).

According to our retardation experiments, the IR is not required for MbeC\_ColE1 binding either, since MbeC does not bind to a 29-bp dsDNA fragment containing the IR (Fig. 3C). However, MbeC\_ColE1 binds to a 41-bp dsDNA fragment containing the *nic* site (Fig. 3C). This is in accordance with experiments that showed that ColE1 mobilization frequency only drops 100 times when a 41-bp *oriT* fragment is used (Table 4). When different *oriT*s of the MOB<sub>p</sub> family were compared, we observed that the MobC/NikA binding sites were located in the corresponding coordinates 1456 to 1496 of MbeC\_ColE1 *oriT* (Fig. 3D). Thus, ColE1 *oriT*, with only 41 bp being sufficient for conjugation, is one of the shortest *oriT* characterized thus far. The fact that the 89-bp *oriT* containing the IR was mobilized 100 times better than the minimal 41-bp *oriT* could be due to the role of this IR in termination of DNA transfer, as proposed for plasmid R64 (16). Other systems, such as R388 and F plasmids, are strongly regulated and thus require longer *oriT*s (>300 bp). The presence of additional regulatory elements, such as extra IRs and protein binding sites, demonstrates the degree of complication of these systems compared to the simple ColE1-based *oriT*.



BLAST analysis indicated that MbeC-like proteins show two main regions of homology, which correspond to the N- and C-terminal folding domains (Fig. 2C). The most conserved is the C-terminal domain, which contains an NLNQ motif. The sequence motif [LF]xxx[GS]xNxNQxAxxxN was described for MobC proteins from gram-positive and gram-negative bacteria (1). There are several MbeC homologues that lack this motif, and they are clustered in a single monophyletic group that contains members from conjugative and mobilizable plasmids (Fig. 4). A distinctive feature of the plasmids contained in this group is the organization of their mobilization region, having their mobilization genes located at both sides of *oriT*. On the other hand, MbeC-like proteins containing the NLNQ motif belong to plasmids whose mobilization genes are placed on the same position with regard to *oriT*. The NLNQ motif is always present in MOB<sub>HEN</sub> and in most MOB<sub>P</sub> accessory proteins, and it could be involved in the interaction with the conjugative helper apparatus, probably with the coupling protein. In fact, TrwA\_R388 has been shown to interact via its C-terminal domain with the coupling protein TrwB\_R388 (37). This interaction stimulates TrwB ATPase activity, being the energy released from ATP hydrolysis used for DNA pumping. Thus, the interaction of the coupling protein with the C-terminal domain of the accessory protein allows the relaxosome to be recruited by the T4SS for DNA transport. We do not exclude the possibility that the NLNQ motif could interact with the MbeA relaxase or any other proteins, but in any case it seems feasible that this motif plays an important function in conjugation.

The conserved RHH N-terminal domain shows a wide distribution among conjugative accessory proteins. In most MbeC-like proteins, an RHH domain can be recognized at the N terminus by second-structure prediction (Fig. 5A). The RHH motif was not observed up to now probably because of the high conservation of the NxNQ motif in the C-terminal domain and the difficulty in the assignment of an RHH signature, due to the small size of the motif and the similarity to the HTH DNA-binding motif. In a previous study (34), the sequences of characterized and putative RHH proteins were compared and showed that they share alternating hydrophilic and hydrophobic amino acids within the N-terminal  $\beta$ 1-strand. In addition, there is a conserved G-X-S/T/N motif in the loop between helix  $\alpha$ 1 and  $\alpha$ 2 and some conserved hydrophobic residues in  $\alpha$ 1 and  $\alpha$ 2. MbeC predicted  $\beta$ 1 strand contains an alternation of hydrophobic (L10, I12, and V14) and hydrophilic (T11 and R13) residues. The MbeC putative loop between helix  $\alpha$ 1 and  $\alpha$ 2 contains the sequence GXQ, and there are conserved hydrophobic residues in  $\alpha$ 1 (L22) and  $\alpha$ 2 (L31 and L40) (Fig. 5A), which is compatible with the consensus RHH motif (34).

A standard PSI-BLAST analysis initiated from the RHH accessory protein NikA\_R64 hit MbeC\_ColE1 in the third iteration. Our in silico analysis of the accessory protein NikA and the three-dimensional structure of its N-terminal domain (which was published [41] while the present study was in preparation), as well as secondary structure prediction Jpred3, PSIPRED, and GOR4 on MbeC, gave more credence to the predicted RHH DNA-binding domain of MbeC.

The existence of the N- and C-terminal folding domains found by the BLAST analysis is in accordance with our experimental results of limited proteolysis of MbeC. After trypsin

treatment and MALDI-TOF spectrometry, we observed two discrete bands of 5,493 and 9,424 Da. The existence of this breakpoint possibly signals the interface between both domains (Fig. 2C). The N-terminal domain (46 amino acids) contains the predicted RHH domain, while the C-terminal domain (51 amino acids) contains the NxNQ domain.

Comparison with other RHH proteins leads to the prediction that the mutant MbeC(R13A) should be critically affected on DNA binding (Fig. 5A). In fact, the point mutation R13A abrogated MbeC binding to DNA and reduced ColE1 mobilization by 3,000 times (Table 3). A similar effect was reported when the arginine of the DNA binding N-terminal beta sheet (R10) was mutated to alanine for the RHH accessory protein TrwA\_R388 (27). This result again confirms the RHH prediction for MbeC and MbeC-like proteins. Moreover, at the mobilization experiments testing the starting codon of MbeC, the MbeC beginning from the second codon, gave conjugation diminution by 10,000 times (Table 4). The predicted N-terminal  $\beta$ -strand of MbeC contains a pattern of alternating hydrophilic and hydrophobic side chains with positive charged Arg or Lys residues at positions 2 or 6 (Fig. 5A) in most of the proteins. MbeC starting from the second AUG loses a predicted polar residue (R6) of the  $\beta$ -sheet (Fig. 5C), which explains the dramatic drop of the mobilization frequency. This finding strengthens even more our hypothesis of a putative RHH N-terminal domain in MbeC.

In conclusion, our study demonstrates that MbeC\_ColE1 is a DNA-binding protein possibly belonging to the RHH family. By comparison to other accessory proteins, we propose that MbeC induces DNA bending, helping the MbeA relaxase to melt the DNA around the *nic* site and cleave the phosphodiester bond. This role has been suggested for most accessory *nic*-processing proteins such as TrwA\_R388, TraJ\_RP4, TraY\_F, and NikA\_R64, all belonging to the RHH family, as confirmed here by sequence comparison. For this reason, despite the differences among the various conjugative systems, we propose that this potentially universal mechanism for DNA processing in bacterial conjugation exists for all accessory proteins.

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