

## NAR Breakthrough Article

# IncC conjugative plasmids and SXT/R391 elements repair double-strand breaks caused by CRISPR–Cas during conjugation

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### ABSTRACT

**Bacteria have evolved defence mechanisms against bacteriophages. Restriction-modification systems provide innate immunity by degrading invading DNAs that lack proper methylation. CRISPR–Cas systems provide adaptive immunity by sampling the genome of past invaders and cutting the DNA of closely related DNA molecules. These barriers also restrict horizontal gene transfer mediated by conjugative plasmids. IncC conjugative plasmids are important contributors to the global dissemination of multidrug resistance among pathogenic bacteria infecting animals and humans. Here, we show that IncC conjugative plasmids are highly resilient to host defence systems during entry into a new host by conjugation. Using a TnSeq strategy, we uncover a conserved operon containing five genes (*vcrx089–vcrx093*) that confer a novel host defence evasion (*hde*) phenotype. We show that *vcrx089–vcrx090* promote resistance against type I restriction-modification, whereas *vcrx091–vcrx093* promote CRISPR–Cas evasion by repairing double-strand DNA breaks via recombination between short sequence repeats. *vcrx091*, *vcrx092* and *vcrx093* encode a single-strand binding protein, and a single-strand annealing recombinase and double-strand exonuclease related to Red $\beta$  and  $\lambda$ Exo of bacteriophage  $\lambda$ , respectively. Homologous genes of the integrative and conjugative element R391 also provide CRISPR–Cas evasion. Hence, the conserved *hde* operon considerably broadens the host range of large families of mobile elements spreading multidrug resistance.**

### INTRODUCTION

The global rise of multidrug-resistant bacterial pathogens is a serious public health issue and a growing economic burden. This ongoing antibiotic resistance crisis is mostly due to improper use and overuse in clinical settings and animal husbandry that is exacerbated by a dried-out antibiotic development pipeline (1–3). While a minority of bacterial pathogens have evolved multidrug resistance exclusively by chromosomal mutations, most have acquired resistance by horizontal gene transfer. Resistance genes are frequently associated with mobile genetic elements (MGEs), such as conjugative plasmids, integrative and conjugative elements (ICEs), and bacteriophages that act as vehicles for dissemination (4,5). Conjugative plasmids of the IncC group are large, broad-host range plasmids found in pathogenic species of *Enterobacteriaceae*, *Vibrionaceae* and *Morganeliaceae* (6). IncC plasmids are commonly associated with multidrug resistance and drive the global dissemination of New Delhi metallo- $\beta$ -lactamase genes (*bla<sub>NDM</sub>*) that confer resistance against most  $\beta$ -lactams including carbapenems (7,8). Most of the conserved core genes of IncC conjugative plasmids are also present and syntenic in ICEs of the SXT/R391 family that have played a key role in the emergence of multidrug resistance in seventh-pandemic *Vibrio cholerae* O1, the infectious agent of the diarrhoeal disease cholera (9).

Upon entry into a new host, these MGEs face multiple host-defence systems that protect the cell against invading DNA molecules (10). Restriction-modification systems (R-M) provide innate immunity against foreign DNA that lacks the proper modification at specific recognition sites, while the host genome is protected from restriction endonuclease activity by methylation of the same recognition sites (11). In contrast, Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR) and CRISPR-associated proteins (Cas) provide adaptive immunity based on an RNA-

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guided endonuclease that seeks for sequence complementarity in nucleic acids and cuts the targeted sequence (12). Guide RNAs are produced from spacer sequences in the CRISPR array that are acquired from foreign invading DNA molecules during previous infections by MGEs. Type I CRISPR–Cas systems use the Cas3 endonuclease to target double-strand DNA (dsDNA) and are frequently found in *Enterobacteriaceae* and *Vibrionaceae* (13–15). Given the vast distribution of host-defence systems and their important role as a barrier against horizontal genes transfer, many MGEs have evolved strategies to evade these defences. Bacteriophages and conjugative elements can produce anti-CRISPR proteins, restriction endonuclease inhibiting proteins, or subvert R–M activities by producing their own modification enzymes (16–20).

Here, we investigated the resilience of the multidrug resistance-conferring IncC conjugative plasmid pVCR94 against CRISPR–Cas immunity. We carried out a high-throughput, transposon insertion site sequencing analysis (TnSeq) on pVCR94 to identify genes that allowed resistance to Cas3 cleavage during conjugation in *V. cholerae*. Our analysis revealed a conserved cluster of five genes that provide resistance to DNA restriction and repair Cas3-mediated double-strand DNA breaks (DSB). Two proteins encoded by these genes (*verx092* and *verx093*) are related to the single-strand annealing protein Red $\beta$  (synaptase) and exonuclease  $\lambda$ Exo of bacteriophage  $\lambda$ 's Red recombination system (21,22). Further investigation using the ICE R391 confirmed the role of its synaptase and exonuclease homologues in the repair of Cas3-mediated DSB during conjugation. Altogether, our work demonstrates that the widespread synaptase–exonuclease recombination systems allow conjugative MGEs to evade a new host's CRISPR–Cas immunity, and suggests a similar role for lambdoid bacteriophages during infection.

## MATERIALS AND METHODS

### Bacterial strains, plasmids and media

The bacterial strains and plasmids used in this study are described in Supplemental Table S1. Bacterial strains were grown at 37°C on Luria-Bertani (LB) agar or with agitation in LB broth. Bacterial strains were maintained at –75°C in LB broth containing 20% (vol/vol) glycerol. Antibiotics were used at the following concentrations: ampicillin (Ap), 100  $\mu$ g/ml; kanamycin (Kn), 50  $\mu$ g/ml; spectinomycin (Sp) 50  $\mu$ g/ml; rifampicin (Rf), 50  $\mu$ g/ml for *Escherichia coli* and 100  $\mu$ g/ml for *V. cholerae*; chloramphenicol (Cm), 20  $\mu$ g/ml for *E. coli* and 2  $\mu$ g/ml for *V. cholerae* and nalidixic acid (Nx), 40  $\mu$ g/ml. For complementation assays, bacterial cultures were supplemented with L-arabinose (0.02%, wt/vol).

### Conjugation assays

Bacteria were grown for 16 h in LB broth with the appropriate antibiotics. Mating assays were carried out by mixing 100  $\mu$ l of the donor (containing conjugative plasmid pVCR94 or ICE R391) and the recipient cells. Cells were pelleted by centrifugation, then washed once in 1 volume of LB broth and resuspended in 1/20 volume of LB broth. Bacterial mixtures were incubated for 2.5 h on LB agar

plates at 37°C to allow conjugation. For ICE R391, excision and conjugative transfer were enhanced by overexpressing *setCD* from pGG2B in the presence of 0.02% arabinose. Serial dilutions were then plated on selective LB-agar plates with appropriate antibiotics to discriminate between donor, recipient and transconjugant CFUs. The frequency of transfer was calculated by dividing the number of transconjugant CFUs by the number of donor CFUs. For each assay, 8 transconjugant colonies were randomly picked and stored at –75°C for whole-genome Illumina sequencing.

### Statistical analyses for conjugation assays

One-way analysis of variance (ANOVA) or unpaired Student *t*-test was used to assess statistical differences for frequencies of transconjugant formation in conjugation assays. A *P* value < 0.05 was considered a statistically significant difference. Each conjugation assay was repeated at least three times.

### Plasmids and strains construction

Plasmid DNA was prepared using the QIAprep Spin Miniprep kit (Qiagen), and genomic DNA was isolated with the QIAamp DNA mini kit (Qiagen) as recommended by the manufacturer. PCR products were purified using the PCR purification kit (Qiagen). All molecular biology manipulations were carried out by standard procedures in accordance with the Current Protocols in Molecular Biology (23). Electroporations were done using a Biorad Gene Pulser Xcell apparatus (BioRad) in 1 mm cuvettes under the following conditions: 200  $\Omega$ , 25  $\mu$ F and either 18 kV/cm for *E. coli* or 10 kV/cm for *V. cholerae*.

The oligonucleotides used in this study are described in Supplemental Table S2. Gene deletions were achieved by using the one-step chromosomal gene inactivation technique ( $\lambda$  Red-mediated mutagenesis) with pKD3 and pKD4 as templates for antibiotic resistance cassettes (24). All deletions were confirmed by Sanger sequencing. Slight modifications were made to adapt  $\lambda$  Red-mediated mutagenesis in *V. cholerae* to obtain the  $\Delta cas$ ,  $\Delta hsdR$  and  $\Delta recA$  mutants based on previously reported procedures for *V. cholerae* (25). Briefly, ~1000-bp fragments upstream and downstream of *casA–E*, *hsdR* or *recA* were PCR-amplified using Q5 high-fidelity polymerase (NEB) and cloned into pCR2.1, flanking the chloramphenicol resistance cassette of pKD3 to yield pCRcrispr, pCRhsdR and pCRrecA, respectively. The resulting plasmids were used as PCR templates for  $\lambda$  Red-mediated mutagenesis to obtain the three deletion mutants.

To introduce protospacers into pVCR94<sup>Kn</sup>, pVCR94<sup>Sp</sup> and R391 and its  $\Delta(bet-exo)$  mutant, the PCR template for  $\lambda$  Red-mediated mutagenesis was constructed as follows. The *cat* chloramphenicol-resistance cassette flanked by FRT sites of pKD3 was digested by HindIII and ligated into pCR2.1. Protospacers V1, V1R, V1NT and V3 were synthesised by oligonucleotide annealing as follows. Complementary pairs of oligonucleotides were designed to include the specific protospacer adjacent motif (PAM) sequence AA in 3' (or 5' for V1NT)

of spacer 1 (V1 and V1R, position 343 477–343 509) or spacer 20 (V3, position 344 636–344 668) of the CRISPR array of *V. cholerae* O395 (Genbank CP000627.1.1) and BamHI restriction sites at both extremities (26) (Figure 1a). *V. cholerae* O395 CRISPR spacer sequences length are typically 32–33 bp and are available in the CRISPRCasdb database at <https://crisprcas.i2bc.paris-saclay.fr/>. For annealing, oligonucleotides were resuspended at 1 mg/ml in annealing buffer (10 mM Tris, pH 7.5–8.0, 50 mM NaCl, 1 mM EDTA). 2 µg of each complementary oligonucleotides were mixed in 50 µl of nuclease-free water and heated at 95°C for 5 min in an aluminium heat block. The heat block was then removed from the heater unit and let to cool down for 60 min. 300 pmol of the mixture was then incubated with 10 units of T4 polynucleotides kinase (New England Biolabs) for 1 h at 37°C, and the enzyme was then inactivated at 65°C for 20 min. Annealed oligonucleotides were digested by BamHI and cloned into pCR2.1::cat to yield pCRV1, pCRV1R, pCRV1NT and pCRV3 that were used as PCR templates for λ Red-mediated mutagenesis after validation of each construction by Sanger sequencing.

The rifampicin-resistant mutants of *V. cholerae* O395 (O395R) was obtained by gradual selection of spontaneous resistant clones grown in LB broth supplemented with increasing concentration of rifampicin (0.1–100 µg/ml). The culture was serially diluted and plated onto LB agar to isolate Rf-resistant clones. Colonies were reisolated and sequenced to validate mutations in *rpoB* (27).

### RNA extraction and cDNA synthesis

Briefly, RNA extractions were performed as follows. *E. coli* VB111 containing pVCR94<sup>Kn-V1</sup> and *V. cholerae* O395R were grown at 37°C for 16 h in LB broth containing the appropriate antibiotics and a conjugation assay was performed as described above. After incubation, the conjugation mixture was resuspended in 1 ml of LB broth and this bacterial suspension was used for total RNA extraction using Direct-zol RNA extraction kit (Zymo Research) and TRI Reagent (Sigma-Aldrich) according to the manufacturer's instructions. Once purified, the RNA samples were treated using 2 units of DNase I (New England Biolabs) according to the manufacturer's instructions to eliminate any residual gDNA. cDNA was synthesised from 0.5 µg of RNA and 2 pmol of gene-specific primer vcrx094RT (Integrated DNA Technologies), using the reverse transcriptase SuperScript III (Invitrogen), according to the manufacturer's instructions. Control reactions without reverse transcriptase treatment ('noRT') were performed for each sample.

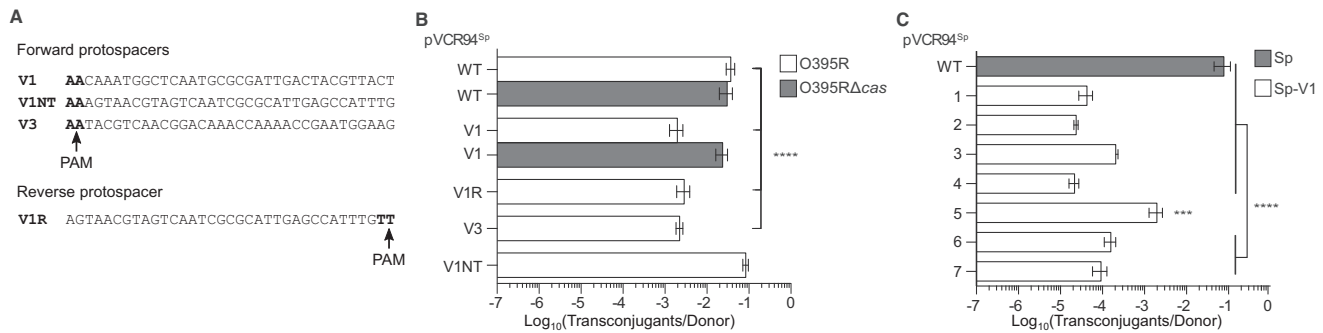
### High-density Tn5 transposon mutagenesis

A conjugation-assisted random transposon mutagenesis experiment was performed on *E. coli* VB111 bearing pVCR94<sup>Kn-V1</sup>. The transposition system was composed of *E. coli* MFD<sup>pir+</sup> carrying pFG036 (a plasmid coding for a thermosensitive cI transcriptional repressor) and pFG051 (a Pi-dependent RP4-mobilizable plasmid coding for the Tn5 transposition machinery repressed by cI, and carrying a mini-Tn5 (Sp) transposon). This diaminopimelate

(DAP)-auxotrophic strain contains a chromosomal RP4 conjugative machinery and expresses the Pi protein required for pFG051 replication. The TnSeq experiment was performed in several successive steps. First, pFG051 was transferred by conjugation from MFD<sup>pir+</sup> to VB111 bearing pVCR94<sup>Kn-V1</sup> in a 2-h mating experiment at 30°C on LB agar plates supplemented with DAP in duplicates. Once in the recipient strain that lacks cI, the Tn5 machinery of pFG051 was constitutively expressed to mediate random mini-Tn5 (Sp) insertions in the genome. The mating mixture was then entirely spread onto 40 large LB agar plates (150 mm) supplemented with Cm, Kn, Nx and Sp. Plates were incubated until near confluence (10k to 40k CFUs) to select clones carrying mini-Tn5 (Sp) insertions. After overnight incubation at 37°C, Cm Nx Kn Sp-resistant colonies were collected using a cell scraper and resuspended in LB broth. The collected sample, designated as the 'input library' was washed, then resuspended in 4.5 ml of LB broth and cryopreserved. Total DNA of a 1.5 ml aliquot of the input library was extracted for sequencing. Another 1 ml aliquot of input library was used to inoculate 50 ml of LB broth supplemented with Cm, Kn, Nx and Sp, which was then grown overnight at 37°C. The resulting culture was used as the donor in a mating, where it was mixed in equal volumes with either *V. cholerae* O395R or O395RΔ*cas*. After a 5 h incubation at 37°C, mating mixtures were spread onto 30 large LB agar plates (150 mm) supplemented with Rf, Kn and Sp to select for mini-Tn5 (Sp) insertions in pVCR94<sup>Kn-V1</sup> that allowed transmission and replication of the plasmid into the recipients. After overnight incubation at 37°C, Rf, Kn and Sp-resistant colonies were collected and subsequently resuspended in LB broth, washed and resuspended in 4.5 ml of LB broth and cryopreserved. These samples were designated as 'output libraries'. Total DNA of a 1.5 ml aliquot of the output libraries was extracted and used for sequencing.

### Preparation of TnSeq sequencing libraries

For each input or output library, a 1.5 ml frozen stock aliquot was thawed on ice for 15 minutes. The aliquot was centrifuged, and cells were resuspended in 300 µl of cell lysis buffer from the Quick gDNA Miniprep kit (ZymoResearch). Total genomic DNA was then purified with AMPure DNA XP magnetic beads (Agencourt). Purified DNA was then fragmented and prepared for Nextera sequencing with the NEBNext Ultra II FS DNA Library Prep Kit (New England Biolabs) for Illumina. At the step of adapter ligation, we used our adapter for Nextera sequencing. Nextera adapter B was generated by annealing two oligonucleotides: 5'-PO<sub>4</sub>-CTGTCTCTTATACACATCTCCGAGCCCACGAGAC-InvdT-3' and 5'-CAAGCAGAAGACGGCATAACGAGATTCGCCTTAGTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGT-3' together. Annealing was performed by heating of each oligonucleotide (40 µM final concentration) in annealing buffer (10 mM Tris NaCl pH 7.5, 50 mM NaCl) to 98°C and then slowly decreasing the temperature by 0.1°C every 10 s down to 4°C. DNA was purified again using DNA Ampure XP beads (Agencourt), and barcoding was performed in a qPCR machine using Veraseq 2.0 High-Fidelity DNA



**Figure 1.** Insertion of O395 protospacers in pVCR94 reduces its transfer to Cas<sup>+</sup> *V. cholerae* O395. (A) Sequence of the protospacers introduced into pVCR94<sup>Sp</sup>. (B) Conjugative transfer of pVCR94<sup>Sp</sup> derivatives bearing O395 protospacers from *E. coli* VB111 to *V. cholerae* O395R (Cas<sup>+</sup>) or O395RΔcas (Cas<sup>-</sup>). (C) Impact of protospacer insertion site on the transfer of pVCR94<sup>Sp</sup> to O395R. Positions of V1 protospacer insertions are indicated by orange arrowheads in Figure 2B. Bars are the mean of three independent experiments. Error bars are the standard error of the mean. Statistical significance: \*\*\*\**P* < 0.0001 and \*\*\**P* < 0.001.

polymerase (Enzymatics). The amplification reaction was stopped at the end of the exponential phase. DNA was purified again and quantified using Quant-it PicoGreen dsDNA assay (Thermo Fisher). The quality and size distribution of the amplified mutant library was assessed on an Agilent 2100 Bioanalyzer instrument using a High Sensitivity DNA Chip. Mutant libraries were then pooled and sequenced by Illumina using the NextSeq<sup>®</sup> 500/550 High Output Kit v2 at the RNomics platform of the Laboratoire de Génomique Fonctionnelle de l'Université de Sherbrooke (<https://rnomics.med.usherbrooke.ca>) (Sherbrooke, QC, Canada). The transposon data analysis began by trimming the reads based on their quality and the presence of the Nextera Illumina adapters using Trimmomatic version 0.36 with the parameters SLIDINGWINDOW:4:20 MINLEN:36 ILLUMINACLIP:<adapters.file.fa>:2:30:15 (28). The quality of the reads was assessed with FastQC version 0.11.5 before and after trimming. The reads were aligned on the genome present in the respective strains with BWA MEM version 0.7.15 using default parameters. SAMtools version 1.3.1 was used to generate alignments statistics and to discard low quality alignments (MAPQ < 10) or multiple alignments (29). In order to get the exact positions of the transposon insertions, we made sam2sites.py, a python script that parses the alignments data and output all the insertions positions and read counts. We used the parameters—read\_len\_threshold 30 and—score\_threshold 0. Note that we chose the middle base pair of the 9-bp Tn5 insertion site duplication to represent the insertion sites (30). The insertions sites files were then encoded in bigWig format using the Kent utilities (31). Visual inspection of the bigWig files was performed using an AssemblyHub on the UCSC Genome Browser (32). To get gene level insertion statistics, we used BEDTools intersect version 2.26.0 together with our python script sites2genes.py (33). The python scripts sam2sites.py and sites2genes.py are available at [https://github.com/fredericQC/HDTM\\_tools](https://github.com/fredericQC/HDTM_tools).

### Preparation of whole-genome sequencing libraries

Whole-genome DNA library preparation was prepared as described for TnSeq libraries with the following modifica-

tions. For each sample, total DNA was extracted from 500 μl of overnight culture, purified and fragmented as specified for TnSeq procedures. For the adaptor ligation step, NEBnext Adaptor for Illumina (New England Biolabs) was used. All the subsequent steps were done as described for TnSeq library preparation above. Different variants of rescued pVCR94<sup>Sp</sup> were sequenced to characterise scars. The assembly workflow started by trimming the reads based on their quality and the presence of the TruSeq Illumina adapters using Trimmomatic version 0.36 with the parameters SLIDINGWINDOW:4:20 MINLEN:36 ILLUMINACLIP:<adapters.file.fa>:2:30:15 (28). The quality of the reads was assessed with FastQC version 0.11.5 before and after trimming. *De novo* assemblies were then generated using SPAdes version 3.11.1 with the parameters -k 21 33 55 77 -careful -only-assembler (34). QUAST version 5.0.2 was used to evaluate the quality of the assemblies (35). Full description of sequenced strains and assembly statistics are available in Supplemental Table S3.

### Bioinformatic analyses

The sequences corresponding to each identified mobile element were downloaded from Genbank Protein homologues of Vcrx089, Bet and Exo were aligned using MUSCLE (36). Pairwise similarity matrices were calculated using UGene 1.32 (37) and plotted as heatmaps using the Heatmapper server (<http://www.heatmapper.ca/pairwise>). Repeat sequences in Supplemental Figure S1 were aligned using ClustalW alignment software (38). mRNA structure predictions were performed using RibEx (39). mRNA structure folding was carried out using the RNAfold web server at <http://rna.tbi.univie.ac.at/cgi-bin/RNAWebSuite/RNAfold.cgi>.

## RESULTS

### CRISPR–Cas3 of *V. cholerae* O395 impedes the transfer of a targeted pVCR94 mutant

To investigate the sensitivity of pVCR94 to CRISPR–Cas activity and identify new anti-CRISPR proteins encoded by IncC plasmids, we took advantage of *V. cholerae*

O395, a strain that naturally expresses CRISPR–Cas3 immunity (26). Since the CRISPR array of O395 is devoid of spacers targeting pVCR94, protospacers corresponding to spacers 1 (V1) and 20 (V3) of O395 CRISPR array were inserted downstream of *verx093* in the spectinomycin-resistant derivative pVCR94<sup>Sp</sup> (Figures 1A and 2B, orange arrowhead 5 on pVCR94<sup>Sp</sup> map).

The resulting plasmids were donated by conjugation in mating assays using *E. coli* VB111 (Nx) as the donor strain, and a rifampicin-resistant derivative of *V. cholerae* O395 (O395R) or a mutant lacking *casA-E* (O395R $\Delta$ *cas*) as the recipient strains. Protospacer V1 reduced plasmid transfer to the Cas<sup>+</sup> strain by 19-fold compared to the Cas<sup>-</sup> strain (Figure 1B). V1 in reverse orientation (V1R) and V3 produced a similar effect, whereas Non-Target V1 (identified as V1NT) with the protospacer adjacent motif (PAM) located at the 5' end produced no interference, thereby confirming the specific yet partial immunity of O395R against pVCR94<sup>Sp-V1</sup> (Figure 1A and B).

Next, to evaluate whether any positional effect resulted from the insertion site, protospacer V1 was also inserted at six alternative intergenic loci across pVCR94<sup>Sp</sup> (Figure 2B, orange arrowheads 1–7 on pVCR94 map) to compare with the phenotype obtained with V1 inserted downstream of *verx093* at position 5. Again, we observed a partial immunity of O395R for all positions. However, the frequency of transfer of the resulting plasmids was differentially impacted by the position of V1, spanning an 80-fold difference between position 5 and positions 2 or 4 (Figure 1C). Notably, pVCR94<sup>Sp-V1</sup> rescue was optimal with V1 inserted at position 5. Although reasons for the observed positional effect will be explored below, insertion of V1 at position 5 was used for all further experiments.

### Identification by TnSeq of a conserved host defence evasion (*hde*) locus

Since the strain O395R failed to abolish the entry and persistence to pVCR94<sup>Sp-V1</sup> (partial immunity), we hypothesized that this plasmid encodes a CRISPR–Cas evasion system. To identify the gene(s) responsible for this phenotype, we devised a TnSeq strategy aimed at targeting critical genes that allow evasion of the O395R CRISPR–Cas system during conjugation (Figure 2A). We constructed a high-density Tn5 insertion library in *E. coli* VB111 bearing the kanamycin-resistant variant pVCR94<sup>Kn-V1</sup>. This pVCR94 variant lacks *acr2*, the main repressor of IncC plasmid transfer (40,41), to increase conjugative transfer and prevent an insertion bias in this negative regulator gene. This set of pVCR94<sup>Kn-V1</sup> mutants represents the ‘input library’ enriched in Tn5 insertions enabling plasmid replication and maintenance. The input library was used in mating assays to donate the set of mutated plasmids to *V. cholerae* O395R and O395R $\Delta$ *cas*. The two sets of transconjugant colonies correspond to ‘output libraries’. Insertions in pVCR94<sup>Kn-V1</sup> that are depleted in the output libraries compared to the input library were expected to reveal plasmid genes or sequences that are important for conjugation (with O395R $\Delta$ *cas*) and involved in CRISPR–Cas evasion or DNA repair (with O395R).

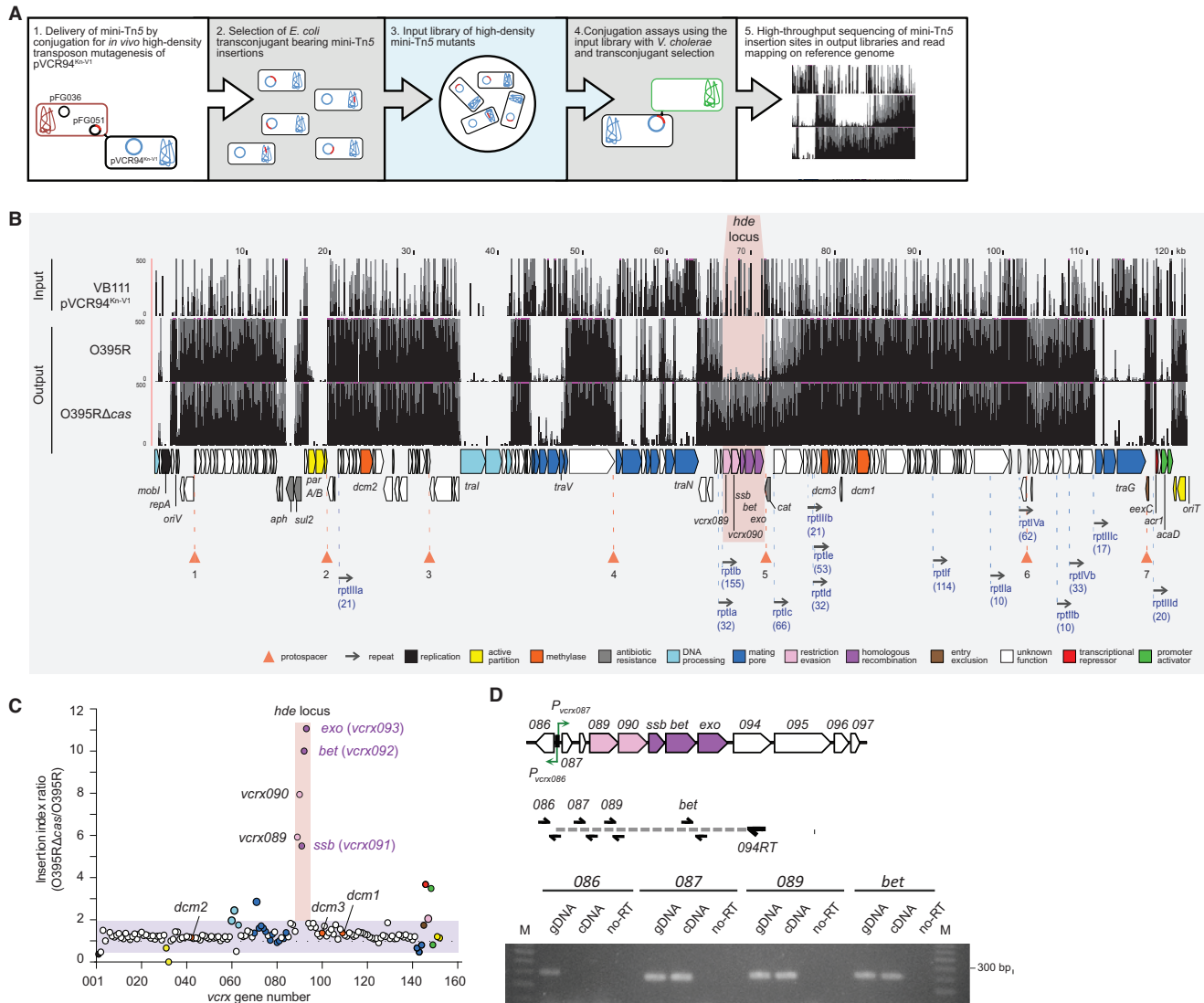
Mean coverages of 289 $\times$ , 3608 $\times$  and 3537 $\times$  were obtained for the input library, and the O395R and O395R $\Delta$ *cas* output libraries, respectively. These results suggested sufficient saturation of the libraries to enable identification of genes essential for replication, maintenance, transfer, and repair. O395R and O395R $\Delta$ *cas* outputs exhibited strikingly similar Tn5 insertion patterns (Figure 2B). Transfer-associated genes (e.g. *traN* and *traG*), plasmid replication and maintenance genes (e.g. *repA* and *parAB*) had virtually identical Tn5 insertion depletions in both conditions (Supplemental Table S4), except for a discrete  $\sim$ 4.7-kb region, hereafter referred to as *hde* for Host Defence Evasion, exhibiting rare Tn5 insertions in O395R. Indeed, the five genes *verx089*, *verx090*, *verx091*, *verx092* and *verx093* located in the *hde* locus underwent a 5- to 12-fold depletion of Tn5 insertions in the Cas<sup>+</sup> strain (Figure 2C and Supplemental Table S4).

While no Pfam domain was found for the translation product of *verx090*, *verx089* is predicted to code for a protein containing a CbbQ\_C domain (Pfam PF08406) and an ATPase AAA\_5 domain (Pfam PF07728). *verx091*, *verx092* and *verx093* code for a single-strand binding protein (Pfam PF00436: SSB), a single-strand annealing protein (Pfam PF03837: RecT) and a double-strand specific 5' to 3' exonuclease (Pfam PF09588: YqaJ). The translation products of *verx092*, and *verx093*, hereafter named Bet and Exo, share 25% and 24% identity with Red $\beta$  and  $\lambda$ Exo encoded by bacteriophage  $\lambda$ . Together with *gam*, *bet* and *exo* form the  $\lambda$  Red system that is widely used as a chromosomal gene deletion and replacement tool due to its ability to facilitate RecA-independent recombination between identical sequences as short as 36 bp (24).

Based on their location, we hypothesized that the five genes of the *hde* locus are part of a polycistronic transcript. A reverse transcription experiment was performed using a primer located at the 3' end of *verx094* and total RNA extracted from VB111 cells bearing pVCR94<sup>Sp</sup>. PCR amplification using the generated cDNA revealed that *verx087*, *verx089* and *bet* but not *verx086* are part of the same mRNA transcript, likely initiated at *P*<sub>*verx087*</sub> (Figure 2D). Therefore, all five *hde* genes are part of the same operon.

### *verx089-090* is involved in restriction evasion

To investigate the role of the *hde* genes in the CRISPR–Cas evasion phenotype observed above, we constructed single and combined deletion mutants both in the presence and absence of protospacer V1 at position 5. In the absence of V1, deletion of both *verx089* and *verx090* led to a 25-fold reduction of conjugative transfer of pVCR94<sup>Sp</sup> to O395R (Figure 3A). Transfer was restored to wild-type level upon ectopic expression of both genes in the recipient, but not in the donor cells, suggesting that *verx089-090* acts upon entry into the new hosts. The same experiment conducted using *E. coli* VB112 as the recipient strain showed that pVCR94<sup>Sp</sup> and its  $\Delta$ (*verx089-90*) mutant transferred at the same rate (Figure 3B). *E. coli* VB111 and VB112 are virtually isogenic strains with identical EcoKI (AACN<sub>6</sub>GTGC) methylation patterns. In contrast *V. cholerae* O395 codes for a type I R-M enzyme predicted by REBASE (42) to recognizes the pattern AAGN<sub>6</sub>CATC. To test whether the trans-

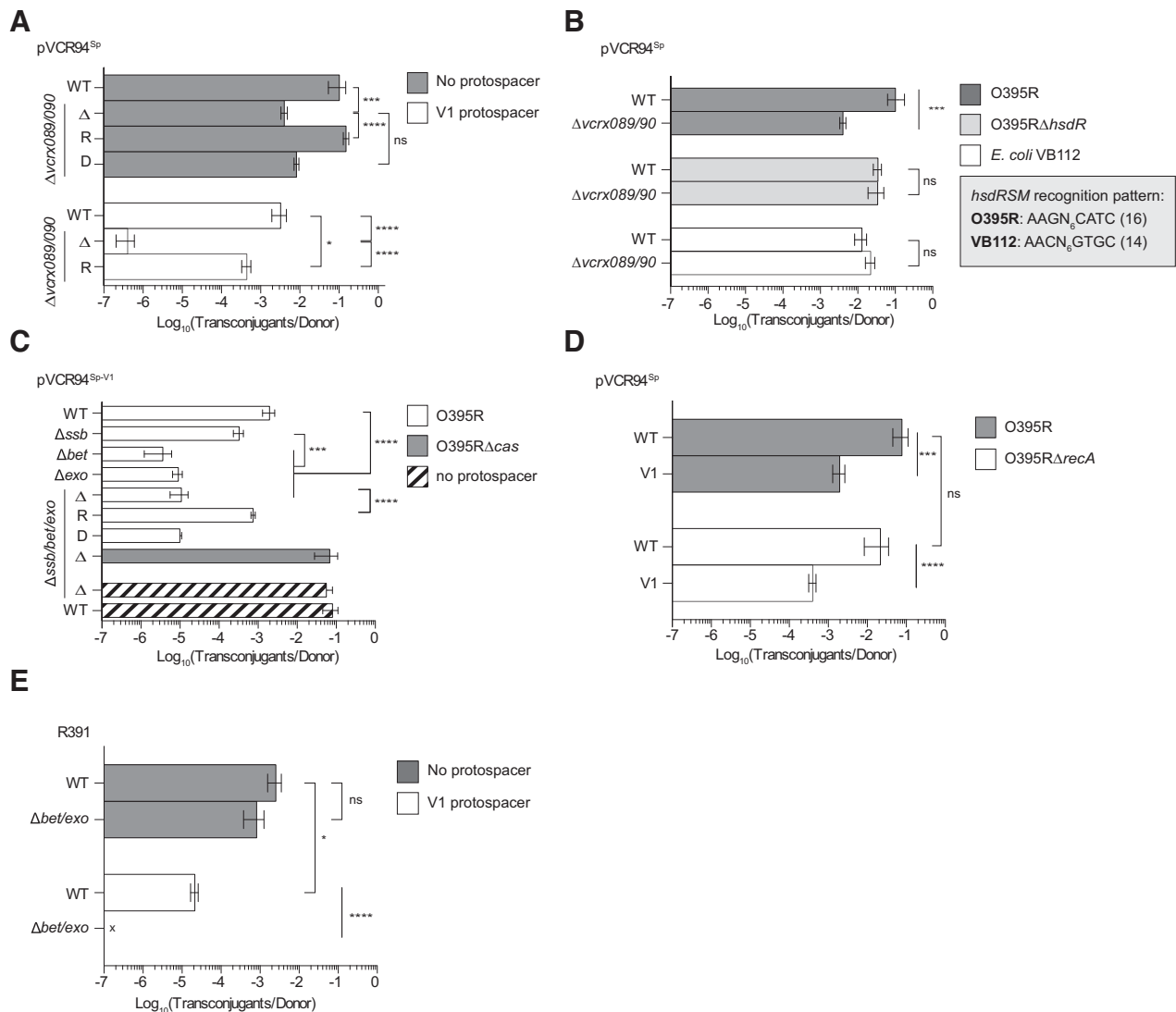


**Figure 2.** Identification of CRISPR and restriction evasion functions by TnSeq. (A) Overview of the TnSeq workflow for the identification of anti-CRISPR functions in the IncC plasmid pVCR94. (B) A cluster of five genes is essential for the transfer of pVCR94<sup>Kn-V1</sup> from *E. coli* VB111 to *V. cholerae* O395R (Cas<sup>+</sup>) but not O395RΔ*cas* (Cas<sup>-</sup>). The three first tracks plot the number of mini-Tn5 reads mapped as a function of the position in pVCR94<sup>Kn-V1</sup> (black bars). (C) Plot of mini-Tn5 insertion index for each gene (*vcrx*) of pVCR94<sup>Kn-V1</sup>. Insertion indexes were calculated as the ratios of insertion counts between O395RΔ*cas* and O395R. Genes are color-coded as in panel b. (D) *hde* genes are part of a single operon. A 2% agarose gel from an assay to amplify genes of the *hde* locus on the *vcrx094*-derived reverse transcribed RNA. Genomic DNA of VB111 carrying pVCR94<sup>Sp</sup> (gDNA) and reverse transcription samples in the absence of reverse transcriptase ('noRT') were used as positive and negative PCR controls, respectively. M, 100 bp Plus II DNA ladder (Civic Bioscience).

fer deficiency of pVCR94<sup>Sp</sup> Δ(*vcrx089-90*) resulted from restriction by the recipient strain, mating assays were carried out using a Δ*hdsR* mutant of O395R that is unable to produce the endonuclease subunit of the type I R-M enzyme (locus tag VC395\_1879; Genbank CP001235.1). pVCR94<sup>Sp</sup> and its Δ(*vcrx089-90*) mutant transferred at the same rate to O395RΔ*hdsR*, thereby suggesting that *vcrx089-90* encodes an anti-restriction mechanism (Figure 3B).

In the presence of protospacer V1, we observed a 5000-fold reduction of transfer of the Δ(*vcrx089-90*) mutant to O395R, suggesting a key role of *vcrx089-90* in CRISPR–Cas evasion. However, complementation assays failed to restore the transfer of this mutant to the level of

pVCR94<sup>Sp-V1</sup> (Figure 3A). Hence, the strong transfer deficiency of pVCR94<sup>Sp-V1</sup> Δ(*vcrx089-90*) likely results from both a sensitivity to restriction and a probable polar effect of the mutation on the expression of the downstream genes *ssb*, *bet* and *exo*. While the cause of this phenotype is not clearly understood, deletion of *vcrx089-90* preserved a predicted RNA secondary structure that likely sequesters the ribosome binding site of *vcrx089* (Supplemental Figure S2). The deletion places this structure 212 bp upstream of the start codon of *ssb*, leaving a large untranslated region that could negatively impact the translation initiation of *ssb*, *bet* and *exo* or the stability of the altered *hde* mRNA. Hence, in the context of Cas-mediated DSB, *vcrx089-90* restric-



**Figure 3.** *hde* genes impact the resilience of translocated DNA against DNA cleavage in recipient cells during conjugation. (A) Effect of *vcrx089/090* on transfer of pVCR94<sup>Sp</sup> with or without protospacer V1 using O395R as the recipient. Complementation of the  $\Delta(vcrx089-090)$  mutation was carried out using pVcrx089-90 in either donor (D) or recipient (R) cells. (B) Effect of *vcrx089/090* on transfer of pVCR94<sup>Sp</sup> devoid of protospacer V1 using O395R and its restriction deficient  $\Delta$ *hsdR* mutant or VB112 as the recipient. The pattern recognized by the type I R-M enzyme of each recipient is indicated. The corresponding number of sites in pVCR94<sup>Sp</sup> is shown in parenthesis. (C) Effect of *ssb*, *bet* and *exo* on transfer of pVCR94<sup>Sp</sup> with or without protospacer V1 using O395R or its  $\Delta$ *cas* mutant as the recipient. Complementation of the  $\Delta(ssb-exo)$  mutation was carried out using pSsbBetExo in either donor (D) or recipient (R) cells. (D) Effect of *bet* and *exo* on the transfer of R391 with or without protospacer V1 using O395R as the recipient. (E) Effect of *recA* in recipient cells on transfer pVCR94<sup>Sp</sup> with or without protospacer V1. For all mating assays, *E. coli* VB111 containing pVCR94<sup>Sp</sup> variants was used as the donor strain and transconjugants were selected as Rf Sp-resistant colonies. Bars are the mean of three independent experiments. Errors bars are the standard error of the mean. An 'x' indicates that the frequency of transconjugant formation was below the detection limit ( $<10^{-7}$ ). Statistical significance: \*\*\*\* $P < 0.0001$ ; \*\*\* $P < 0.001$ ; \* $P < 0.05$ ; ns, not statistically significant.

tion evasion function seems to be at least in part dependent upon the downstream *ssb/bet/exo* genes.

### *ssb/bet/exo* allows CRISPR–Cas evasion of IncC plasmid during conjugation

To test this hypothesis, *ssb*, *bet* and *exo* deletion mutants of pVCR94<sup>Sp</sup> or pVCR94<sup>Sp-V1</sup> were used in mating experiments. The  $\Delta(ssb-exo)$  mutation had no impact on the transfer of pVCR94<sup>Sp</sup> to O395R (Figure 3C). In contrast, deletion of any of the three genes resulted in a 6- to >200-fold reduction of pVCR94<sup>Sp-V1</sup> transfer, thereby suggesting an enhanced sensitivity of these mutants to Cas3-mediated

cleavage (Figure 3C). This phenotype was reversed when *ssb-bet-exo* was ectopically expressed in the recipient but not in the donor cells or when O395R $\Delta$ *cas* was used as the recipient (Figure 3C). Altogether, these results confirm that *ssb*, *bet* and *exo*, while not required for transfer, provide an evasion mechanism against CRISPR–Cas immunity during entry by conjugation into a Cas<sup>+</sup> strain.

### Bet-Exo-mediated DNA repair is RecA-independent

$\lambda$  Red mediates homologous recombination by single-strand annealing (SSA) in a RecA-independent fashion (43). Likewise, Bet and Exo of SXT/R391 ICEs have been

**Table 1.** Summary of the scar patterns found in rescued pVCR94<sup>Sp</sup> carrying V1 protospacer at diverse positions

Protospacer position	Number of sequenced clones <sup>a</sup>	Type of scars <sup>b</sup> (distribution)	Conjugation properties	Repeat ID used for repair <sup>c</sup>	Deletion size (bp) <sup>d</sup>
5	73/85	IA (43/85)	Transmissible	rptIb-rptIf	23 702
		IB (1/85)	Transmissible	rptIb-rptIf	23 702
		IC (7/85)	Transmissible	rptIb-rptIe	14 076
		ID (1/85)	Transmissible	rptIb-rptIe	14 076
		IE (1/85)	Transmissible	rptIb-rptId	14 003
		IF (4/85)	Transmissible	rptIb-rptIc	10 310
		IG (16/85)	Transmissible	rptIa-rptIc	9268
1	0/85	No scar		NA	NA
7	1/85	III (1/85)	Deficient	rptIIIc-rptIIIId	6728
3	2/85	IV (2/85)	Deficient	rptIIIa-rptIIIb	59 432
6	9/85	VA (1/85)	Transmissible	rptIIa-rptIIb	5138
		VB (8/85)	Transmissible	rptIVa-rptIVb	4936
2	0/85	No scar		NA	NA
4	0/85	No scar		NA	NA

<sup>a</sup>0 values, no repaired transconjugants was obtained.

<sup>b</sup>Partial sequences of characterized scars are available in Supplemental Figure S2.

<sup>c</sup>As depicted in Figure 2B. Sequences of repeat IDs are shown in Table 2. NA, not applicable.

<sup>d</sup>Exact deletion size from pVCR94<sup>Sp-V1</sup> variants

shown to promote ICE diversity independently of RecA (21). To test whether RecA is involved in IncC plasmid rescue, a  $\Delta recA$  mutant of O395R was constructed and used as the recipient strain in mating assays. Transfer of pVCR94<sup>Sp-V1</sup> remained unaffected by the *recA* mutation (Figure 3D), confirming that the DNA repair process mediated by Bet-Exo recombination occurs independently of RecA.

### Bet/Exo repairs Cas-generated DSBs by annealing conserved short direct repeats

Since the position of protospacer insertions influences the efficiency of DSB repair (Figure 1C), rescued plasmids that were recovered from O395R transconjugants were further investigated by PCR amplification and sequencing of scar sequences. Only a handful of transconjugants were obtained using pVCR94<sup>Sp</sup> with V1 inserted at positions 1, 2 and 4 (Table 1). Analysis of these rescued plasmids revealed an intact protospacer and surrounding region, suggesting Cas3 loss-of-function, a phenotype also reported for Cas9 endonuclease at a similar rate (44). Surprisingly, standard PCR assays failed to amplify regions adjacent to all other V1 insertion loci, suggesting large deletions or insertions at scar sites. Therefore, whole-genome Illumina sequencing was carried out to determine the structure of the repaired plasmids. Since fragments of the O395R recipient genome could have been used as templates for Bet/Exo-mediated repair, the rescued plasmids were first retransferred into *E. coli* VB111 to facilitate the identification of possible insertions of chromosomal DNA originating from O395R. Out of 85 selected transconjugants carrying rescued IncC plasmids, 78 could donate their plasmid to VB111. Seven plasmids partially or totally lost the ability to conjugate. Consequently, these plasmids were sequenced in O395R. *De novo* assemblies yielded a 61x mean coverage (Supplemental Table S3).

Analysis of the scar sites revealed different repair patterns, even when DSBs occurred at the same position.

Eleven different scar patterns were observed for all protospacer positions (Table 1 and Supplemental Table S3). All repairs led to deletions; no insertions were found. With a set of 73 representatives, rescued pVCR94<sup>Sp-V1</sup> plasmids were the most frequently sequenced since the protospacer at position 5 yielded the highest rate of transconjugant formation (Figure 1C). Within this set, seven different deletion patterns spanning from positions 9268 to 23 702 bp were identified. No key transfer genes were deleted within this set, allowing the repaired plasmids to remain self-transmissible (Table 1) while being protected from O395R Cas3 activity. Rescued plasmids with protospacer at positions 3, 6 and 7 lost DNA fragments spanning from positions 4936 to 59 432 bp.

All repairs occurred by homologous recombination between repeated sequences distributed across pVCR94 (all repeats found in pVCR94 are listed in Supplemental Table S5). Four distinct repeated sequences (rptI-IV) were used for recombination depending on DSB location and resulted in different scar patterns (Table 1). Except for rptII that is located within two genes of unknown function, all repeats were found to be located within intergenic sequences. rptId allowed recombination between only seven identical nucleotides. The longest repeat, rptIb, was 155 bp and common to all repairs occurring at position 5 except for type IG scars (Table 1 and Supplemental Figure S3). All other type I repeats (rptIa, rptIc-f) were found to be partial repeats of rptIb (Table 2 and Supplemental Figure S1). The use of the same two repeats could yield slightly different scars depending on where homologous recombination took place within the repeats (Table 1 and Supplemental Figure S3), as observed for IA-IB and IC-ID. rptIb and rptIf shared the longest identity stretches (Supplemental Figure S1) and generated the majority of characterised scars (Table 1) when DSB occurred at position 5.

NCBI blastn search of rptI (rptIb), rptIII and rptIV sequences revealed a strong conservation among diverse species of *Gammaproteobacteria*. rptI was found in most IncC conjugative plasmids of *Enterobacteriaceae*



Table 2. Repeated sequence motifs recombined in repaired plasmids

Repeat ID	Nucleotide sequence (5' to 3') <sup>a,b</sup>	Location <sup>c</sup>
rptIa	GTGAGATAGCTGGTCTCCGAAAACAAACAGCA	65,107:65,138
rptIb	AGTCTTCTTACTCATGGTAAAGCCATCAGTAAGCAGATTTTGTCTCCTC CCTGAGGGCGAAAAC <b>GTGCGATAGCTGGTCGCCAAAAACAAACAGCA</b> <u>AATTAACGTTAATTTACTAGCCCAACCGGGCGCATCCGCCCGGTTCCG</u> GGACGTGGTGCGCC	66,087:66,241
rptIc	AGAAAGTGGATTTTGTCTCTCTGAGGGCGAAAAGTGCATAGCTGG TCGCCAAAAACAAACAGCA	75,383:75,448
rptId	GTGAGGTAGCTGGCCTCCCCAAACAAACAGCA	79,816:79,847
rptIe	AATCTACTAACCTAAACCGGTGCATCCGCCCGGTTCCGGGACGTGG TGCGCC	80,265:80,317
rptIf	AAGTCTTCTTACTCATGGTAAAGCCATCAGTAAGCAGATTTTGTCTCCT CCCTGAGGGCGAAAACGTGCATAGCTGGTCGCCAAAAACAAACAGC <u>AAATTAACGTTAATTTATTA</u>	89,788:89,901
rptIIa	CGTCATCGAG	97,397:97,406
rptIIb	CGTCATCGAG	102,535:102,544
rptIIIa	AAA <u>ACTTT</u> CACATGTGAAAGT	20,315:20,335
rptIIIb	<u>ACTTT</u> CACATGTGAAAGT	79,750:79,767
rptIIIc	<u>CTTT</u> CACATGTGAAAGT	109,011:109,027
rptIIId	AAA <u>ACTTT</u> CACATGTGAAAGT	115,736:115,755
rptIVa <sup>d</sup>	AATTACCGTATTACCGTAATTATGATAATTACCGTATTACCGTAATT ATGATAATTACGGTA	101,144:101,205
rptIVb	TAATTATGATAATTACCGTATTATCGTAATTAT	106,096:106,128

<sup>a</sup> Repeat sequence alignments for rptI are shown in Supplemental Fig. 1.

<sup>b</sup> Palindromic sequences are underlined. Sequence in bold represents sequence found in SG11.

<sup>c</sup> Based on pVCR94deltaX reference sequence (Genbank accession number KF551948.1).

<sup>d</sup> Repeats within the repeat sequence are indicated in different colours

and *Vibrionaceae*, such as multidrug resistance plasmid pMG252 from *E. coli* and pVPS129 from *Vibrio parahaemolyticus*, and many other IncC plasmids from *Klebsiella*, *Salmonella*, *Providencia*, *Escherichia*, *Vibrio*, *Citrobacter* and *Shewanella*. Most type 2 IncC plasmids shared repeats highly similar to rptI (>98% identity), whereas type 1a and 1b rptI-like repeats were more divergent (>92% identity). Interestingly, a 32-bp rptIb segment was also found (100% identity) in *Salmonella* genomic island 1 (SG11), a multidrug resistance island that requires IncC plasmids for its own dissemination (45). The rptIII and rptIV repeats were also strictly conserved (100% identity) in most IncC plasmids.

### *ssb/bet/exo* allows CRISPR–Cas evasion of ICE R391 during conjugation

The translation products of *vcx089*, *vcx092*, and *vcx093* share 53%, 59% and 45% with Orf70, Bet and Exo encoded by SXT/R391 ICEs, respectively, and homologues are found in several other mobile genetic elements including IncA, IncT and IncP-7 conjugative plasmids (Supplemental Figure S4a and S4b). Bet and Exo of SXT/R391 ICEs have been shown to promote hybrid ICE formation by RecA-independent homologous recombination between SXT and R391 integrated in a tandem fashion in the chromosome (21). Like IncC plasmids, SXT/R391 ICEs lack  $\lambda$  *gam*, which encodes an inhibitor of *E. coli* SbcCD and

RecBCD exonuclease, but carry a *ssb* gene (46–48). Since SXT/R391 ICEs share homologous *bet* and *exo* genes (46), the role of this system was also investigated using R391. Insertion of protospacer V1 in R391 resulted in CRISPR–Cas interference (76-fold reduction of transfer) (Figure 3e). Deletion of *bet* and *exo* completely abolished the transfer of R391<sup>V1</sup>, confirming the key role of *bet* and *exo* in CRISPR–Cas evasion during ICE transfer.

## DISCUSSION

Although many transfer and maintenance-related genes have been characterised for IncC conjugative plasmids and SXT/R391 ICEs, many genes remain to be characterised (6,8,22,49). Here, we sought genes that could be instrumental for these globally prevalent MGEs to evade CRISPR–Cas immunity deployed by bacteria as a barrier against invading DNA molecules during conjugation, the main mechanism of multidrug resistance dissemination. Several anti-CRISPR proteins are known to directly inhibit type I system Cas3 endonucleases such as AcrIE and AcrIF (16,50). Most of these anti-CRISPR proteins are produced by bacteriophages, and are challenging to predict due to their small size and the lack of conserved sequence motif (16,20,50).

In contrast, our TnSeq approach allowed us to identify host defence evasion (*hde*) functions that are conserved among conjugative plasmids belonging to several incompatibility groups as well as ICEs of the SXT/R391 family. While seeking anti-CRISPR functions, we serendipitously identified the *hde* genes *verx089-090* as genes that protect IncC plasmids against type I R–M systems. *De novo* expression of these genes was required in the recipient but not the donor strain for effective protection against the recipient's type I R–M endonuclease activity. Phages and conjugative elements deploy diverse anti-restriction strategies. Besides evolving genome sequences depleted for restriction sites, these MGEs often code for active protection mechanisms. For instance, IncW plasmids produce ArdC that occludes restriction sites (51). Ral of phage  $\lambda$  stimulates the methylation activity of EcoKI to expedite the protection of its own genome (52). Phage T3 encodes a hydrolase that depletes the pool of S-adenosyl methionine, an essential cofactor for type I R–M endonuclease activity (53). ArdA proteins encoded by diverse conjugative elements such as IncN plasmids and Tn916 ICE mimic the DNA structure to bind and block type I R–M enzymes (54). IncC plasmids code for three conserved cytosine-specific methylation enzymes (*dcm1* [*verx109*, PF00145], *dcm2* [*verx042*, PF00145] and *dcm3* [*verx100*, PF06044]) that were shown to change the host's methylation pattern and help the IncC plasmid pVC211 to cross the R–M barrier of *V. cholerae* O1 El Tor C6706 (17). However, none of the three *dcm* genes came out as essential for escaping restriction in our assays using *V. cholerae* O395R as the recipient (Figure 2B and C), suggesting possible functional redundancy. Neither Vcrx089 nor Vcrx090 share any homology with known anti-restriction factors. CbbQ\_C domain proteins are part of CbbQ/NirQ/NorQ family of proteins which play a role in the post-translational activation of Rubisco. The biological role and origin of such proteins in bacteria is unknown (55). ATPase\_AAA\_5 domain proteins are part of AAA+ super-

family of ATPases that is largely distributed among bacteria and participate in a wide range of cellular processes such as membrane fusion, proteolysis and DNA replication (56). The presence of predicted CbbQ\_C and ATPase AAA\_5 domains in Vcrx089 suggests a novel anti-restriction mechanism that is currently being characterized.

CRISPR–Cas systems can be acquired by horizontal gene transfer and are widely distributed in *Enterobacteriaceae* and *Vibrionaceae* (13). Considering that as adaptive bacterial immune systems, CRISPR arrays collect spacers from past invading DNA molecules, acquisition of IncC plasmid-specific spacer by natural strains is a possible event. For instance, the CRISPR array of *Shewanella putrefaciens* W3-18-1 (Genbank CP000503.1) contains three spacers, 26, 29 and 41, that seem to target the conjugal transfer *traC* gene of IncC plasmids (data not shown). Hence, mitigation of CRISPR–Cas immunity via dsDNA break repair likely broadens the host range of IncC and related plasmids. We have identified Bet (Vcrx092) and Exo (Vcrx093) as key actors in the evasion of IncC plasmids from the host's CRISPR–Cas immunity (Figure 2). Consistent with this function, the deletion of *bet* and *exo* in the distantly related ICE R391 increased its sensitivity to CRISPR–Cas-mediated cleavage. Together with *verx089-090* and *ssb* (*verx091*), *bet* and *exo* are part of a large operon that is conserved in IncC plasmids and SXT/R391 ICEs (Supplemental Figure S4a) (46). The proteins encoded by *bet* and *exo* are distantly related to the Red system of bacteriophage  $\lambda$  (Supplemental Figure S4b) (46,57). While  $\lambda$  Red is widely used for bacterial mutagenesis and synthetic biology (24), its exact biological role remains ambiguous despite its involvement in  $\lambda$  replication (58). Likewise, the products of *bet* and *exo* of SXT/R391 ICEs have been shown to promote recombination between ICEs integrated in the chromosome in a tandem fashion, increasing ICE plasticity and diversity (21). However, ICE tandem arrays were not found in nature, and were shown to be unstable and short-lived in laboratory conditions, suggesting that generation of ICE diversity is likely an accessory consequence of another key function mediated by *bet* and *exo* (46). Moreover, IncC plasmids do not naturally integrate into the chromosome in a site-specific manner, are mutually incompatible, and exclude each other's entry into the same host (59,60), thereby limiting the opportunities for two different IncC plasmids to coexist together within the same cell and recombine. Hence, *bet* and *exo* genes are unlikely dedicated catalysts of plasmid diversity through recombination. Rather, generation of diversity is likely a by-product of the repair function that depends on the availability of compatible repair templates. Indeed, our results strongly support the notion that the products of *bet* and *exo* of IncC plasmids and SXT/R391 ICEs repair DSBs mediated by Cas endonucleases upon entry into an immune host cell. Notably, successful complementation of the *ssb*, *bet* and *exo* null mutants occurred exclusively when the genes were expressed in recipient cells, confirming that CRISPR–Cas evasion does result from pre-existing diversity generated in donor cells prior to conjugation.

RecA-mediated homologous recombination and repair of DSBs by double-strand invasion requires long stretches of identical complementary DNA (61). IncC plasmids are large, low-copy number, entities that are likely maintained

in a single copy per cell, providing little DNA substrate for such repair mechanism upon entry into a new host. Scars characterisation of rescued plasmids revealed that Bet/Exo repaired DSBs by recombining identical repeated sequences as short as 7 bp compared to 9 bp for  $\lambda$  Red (62). These repeats are conserved among IncC plasmids. Bet/Exo-mediated repairs resulted in different deletions (Table 1) consistent with the SSA homologous repair mechanism. Bet/Exo failed to repair DSBs occurring at positions 2 and 4 (Table 1). Several factors may affect the efficiency of DSB repair, including the proximity of essential genes at these two positions, such as partition system *parAB*, replication gene *repA*, antibiotic resistance genes *aph* and *sul2* used for selection, or toxin/antitoxin addiction systems (49). The lack of available repeats near the cleavage sites is another factor that could prevent DSB repair. Bet/Exo-mediated repair of DSB induced by CRISPR–Cas is non-conservative since the region containing the protospacer was eliminated, thereby immunizing the rescued plasmids against the new host. As expected, several rescued plasmids lost their conjugative properties due to the deletion of conjugation-related genes (Table 1). Although our experimental design led to isolation of deletion mutant only, we cannot rule out that DSB repair facilitates DNA insertion. With the proper selective pressure and DNA template, SSA-mediated DSB repair could eventually led to the incorporation of antibiotic resistance genes. Such event could explain the existence of hotspots for the insertion of variable DNA in SXT/R391 ICEs (21,22).

The Ssb protein encoded by *vcx091* was important but not essential for DSB repair, suggesting an accessory role that facilitates DSB repair. The lack of an *ssb* homologue in the *hde* loci of IncA plasmid pRA1 and untyped plasmid pAsa4C is consistent with this observation (Supplemental Figure S4a). Red $\beta$  was recently shown to bind *E. coli* Ssb to facilitate annealing of the 3'-overhang to Ssb-coated ssDNA at the replication fork (63). Whether Bet encoded by IncC conjugative plasmids and SXT/R391 ICEs can bind their host-encoded Ssb protein is unclear. However, since DSB repair of the  $\Delta$ *ssb* mutant of pVCR94<sup>Sp-V1</sup> was impaired, we hypothesize that the plasmid-encoded Ssb protein can coat the 3'-overhangs generated by Exo and facilitate the loading of Bet on these 3'-overhangs. Alternatively, Bet could be directly loaded on the 3'-overhangs via an Exo-Bet protein-protein interaction, and then interacts with Ssb-coated ssDNA of the lagging strand at the replication fork of the same or another copy of the replicating plasmid.

Here, we report that the Bet/Exo system encoded by IncC plasmids and SXT/R391 ICEs protects these MGEs against the host-encoded defence mechanisms, by recombining direct repeats in a RecA-independent fashion to repair DSBs caused by CRISPR–Cas (58,64). Bacteriophage  $\lambda$  can mediate recombination by a RecA-independent SSA mechanism between  $\lambda$ Exo-processed complementary DNA sequences. By analogy, we proposed that, following CRISPR–Cas-mediated DSB, Exo resects the 5'-end of duplex DNA ends to generate a 3'-overhang (Supplemental Figure S5). Specific patterns such as palindromic sequences found in the repeats (Table 2) could pause or slow down the degradation mediated by Exo, similar to Chi sequences that regulate RecBCD nuclease activities in the RecA-dependent

DNA break repair mechanism (65). Ssb binds to the resulting 3'-overhang, protects it from host nuclease and helps recruiting Bet to anneal to a complementary ssDNA strand in a RecA-independent manner. The host ligase, with the collaboration of Bet and Exo, likely seals the resulting nick upon cleavage of 3' ssDNA regions excluded from the duplex (58,64,66). Consistent with our results and this SSA repair model, one copy of the repeat and the region within these repeats are lost, giving rise to different scars, and contributing to plasmid plasticity.

This novel CRISPR evasion mechanism differs from anti-CRISPR proteins as it does not prevent DSB but rescues plasmids by SSA repair after Cas-mediated cleavage (16). Hence, synaptase-exonuclease may protect MGEs against several types of CRISPR–Cas system considering the large distribution of these two-component recombination systems in MGEs. Besides bacteriophage  $\lambda$  and related phages, homologues of Bet and Exo are encoded by IncC, IncA, IncT and IncP-7 conjugative plasmids, SXT/R391 ICEs, as well as many other uncharacterized mobile elements in distantly related bacteria such as *Bacillus subtilis* and *Borrelia hermsii* (46,67,68). Interestingly, most IncC plasmids but not SXT/R391 ICEs share repeats I and III, thus suggesting that distinct Bet/Exo systems may have evolved together with specific sets of repeated motifs. The presence of a portion of *rptIb* in SGI variants may help explain the evolution of these genomic islands that are widespread in *S. enterica* and highly reliant on IncC plasmids for their propagation (69). The involvement of Bet (RecT) has been suggested in the elevated recombination rate that occurs in SGI1 in the presence of an IncC helper plasmid and leads to the generation of SGI1 deletion mutants (69). However, the involvement of *rptIb* has not been demonstrated.

Therefore, synaptase-exonuclease recombination systems, together with the Vcrx089-90 anti-restriction system, play a crucial role in the acquisition, persistence, and dissemination of antibiotic resistance genes in bacterial populations by helping the conjugative elements that carry these genes to evade defence systems deployed by bacteria as barriers against invading DNA molecules. By broadening the host range of highly prevalent MGEs, *hde* functions could hinder efforts to develop CRISPR–Cas targeting systems, such as engineered conjugative plasmids or bacteriophages, aimed at controlling the dissemination of multidrug-resistant bacteria (70–72).

## DATA AVAILABILITY

Raw sequencing data were submitted to Genbank under Bioproject accession number PRJNA602297 with the following Biosample accession numbers: for TnSeq with O395R and O395R $\Delta$ *cas* as the recipient strains, SAMN13887834 and SAMN13887835, respectively; for TnSeq with VB111 bearing pVCR94<sup>Kn-V1</sup> as the donor strain (input), SAMN13910572; for whole-genome sequencing of transconjugants bearing repaired IncC plasmids, SAMN13887703 to SAMN13887713. The python scripts *sam2sites.py* and *sites2genes.py* are available at [https://github.com/fredericQC/HDTM\\_tools](https://github.com/fredericQC/HDTM_tools).

Complete data from aligned reads for TnSeq experiments can also be visualized using the UCSC genome

browser at [http://genome.ucsc.edu/cgi-bin/hgTracks?genome=pVCR94dX3dacr2spacer1&hubUrl=https://datahub-101-cw2.p.genap.ca/roy\\_2020/hub.txt](http://genome.ucsc.edu/cgi-bin/hgTracks?genome=pVCR94dX3dacr2spacer1&hubUrl=https://datahub-101-cw2.p.genap.ca/roy_2020/hub.txt).

## SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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