

Demonstration of Conjugative Transposon (Tn5397)-Mediated Horizontal Gene Transfer between *Clostridium difficile* and *Enterococcus faecalis*[∇]

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Antibiotic-resistant *Enterococcus faecalis* and *Clostridium difficile* are responsible for nosocomial infections in humans, in which they inhabit the same niche. Here, we demonstrate transfer of the conjugative transposon Tn5397 from *C. difficile* 630 to *E. faecalis* JH2-2, the first reported gene transfer between these two bacteria. Furthermore, transfer from the *E. faecalis* EF20A transconjugant to the epidemic ribotype 027 *C. difficile* strain R20291 was also demonstrated. Tn5397 was shown to use a single specific target site in *E. faecalis*; it also has specific target sites in *C. difficile*. These experiments highlight the importance of continual monitoring for emerging resistances in these bacteria.

Clostridium difficile is an enteric pathogen and the leading cause of antibiotic-associated diseases ranging from self-limiting diarrhea to severe pseudomembranous colitis (7, 10, 19). Treatment is usually by oral metronidazole or vancomycin (4, 13, 15). *Enterococcus* spp. are opportunistic pathogens responsible for a wide variety of infections (2, 12, 24). Vancomycin resistance in enterococci is now widespread (3). In *E. faecalis*, the genes responsible have been shown to be carried on conjugative transposon Tn1549 (6, 20). *C. difficile* and *Enterococcus* spp. often coexist in patients, causing increased severity of illness and prolonged duration of hospitalization (1, 8, 5). Both bacteria can acquire resistance to multiple antibiotics via mobile elements (16).

Conjugative transposons (17) are capable of intracellular transposition and intercellular conjugation, often across large phylogenetic distances (16). Conjugative transposon Tn5397 confers resistance to tetracycline (Tc) and was originally identified in *C. difficile* strain 630 (14). This element contains *tet*(M) and has been shown to be transferable from *C. difficile* to *Bacillus subtilis* and vice versa and between *C. difficile* strains (14). In this study, we wanted to assess the potential for gene transfer of Tn5397 between *C. difficile* and *E. faecalis*.

All bacterial strains used in this study are shown in Table 1. *C. difficile* were grown anaerobically (80% N₂, 10% CO₂, and 10% H₂) at 37°C on brain heart infusion (BHI) agar supplemented with 5% defibrinated horse blood (E & O Laboratories, United Kingdom) and *C. difficile*-selective supplement (Oxoid, United Kingdom) or in prerduced BHI broth. *E. faecalis* and *Escherichia coli* were grown aerobically on or in BHI or LB agar or broth at 37°C. Antibiotics were used at concentrations of 10 µg/ml (tetracycline and erythromycin [Erm]), 25 µg/ml (rifampin [Rif]), 5 µg/ml (fusidic acid [Fus]), and 100 µg/ml (ampicillin [Amp]) (Sigma). Filter mating was

carried out as previously described (18). *E. faecalis* transconjugants were selected on tetracycline-, rifampin-, and fusidic acid-containing plates with aerobic growth. *C. difficile* R20291 transconjugants were selected on tetracycline- and erythromycin-containing plates with anaerobic growth. PCR was carried out with primers (Table 2) by using the GeneAmp 2400 PCR system (Perkin-Elmer Cetus) under the following conditions: 94°C for 4 min, followed by 30 cycles of 94°C for 1 min, 50 to 60°C for 1 min (dependent on primer annealing temperature), 72°C for 1 to 2 min (dependent on expected amplicon size), and 72°C for 4 min. Amplicons were sequenced at the Department of Biochemistry, University of Cambridge, United Kingdom. For Southern hybridization, genomic DNAs from the *E. faecalis* JH2-2 (recipient) and transconjugants were digested overnight with XmnI at 37°C, followed by separation on a 1% agarose gel. Fragments were then transferred to a Hybond N+ nylon membrane (Amersham Biosciences, United Kingdom) and hybridized to a 300-bp probe derived from the *tnaX* gene. Analysis of the Southern blot was performed using ECL direct nucleic acid labeling and detection systems (Amersham Biosciences, United Kingdom) according to the manufacturer's instructions.

To determine the insertion site in *E. faecalis*, single-specific-primer PCR (sspPCR) was used. Genomic DNAs (3 µg) of the *E. faecalis* EF20A transconjugant (Table 1) and pUC19 (0.1

TABLE 1. Bacterial strains

Bacterium	Property(ies)	Reference or source
<i>C. difficile</i> 630	Tc ^r , Erm ^r	25
<i>C. difficile</i> R20291 ^a	Erm ^r	Anaerobe Reference Laboratory, Cardiff, UK
<i>E. faecalis</i> JH2-2	Rif ^r , Fus ^r	11
<i>E. faecalis</i> EF20A	Tc ^r , Rif ^r , Fus ^r	This work
<i>E. coli</i> (α-select, bronze efficiency) competent cells	Amp ^r	Biolone, UK

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^a This strain was responsible for an epidemic outbreak at Stoke Mandeville Hospital, United Kingdom (9).

TABLE 2. Primer and probe details

Primer/probe ^a	Sequence (5'-3')	Amplicon and/or purpose ^b
Tndx (F30)	CTTACAATGTTAAAACAGCAAGC	1.6-kb fragment of <i>tndX</i>
Tndx (R1612)	GAGAATGTATCAATGAGACACTG	1.6-kb fragment of <i>tndX</i> 300-bp probe for Southern analysis
TndX probe	CTTTAGGGAAAATAACTGAT	300-bp probe for Southern analysis
LEO	CCACTTGATATGAAAAATCAAATGGCTC	sspPCR, CI, left-end transposon-genome junction
REO	ACGTGTATCAAGCAGAGGGAATCGGTAAA	sspPCR, CI, right-end transposon-genome junction
PTS forward primer	GTGTCAATGACCGCAGAAGA	Chromosomal target site, left-end transposon-genome junction
PTS reverse primer	TCGCTAGAATGACCTGTAGAAGAA	Chromosomal target site, right-end transposon-genome junction
M13 forward primer	GTAAAACGACGGCCAGT	sspPCR
M13 reverse primer	CAGGAAACAGCTATGAC	sspPCR

^a PTS, phosphotransferase system.

^b CI, circular intermediate.

μg) were digested with either EcoRI or BamHI. Digested pUC19 was dephosphorylated with calf intestine alkaline phosphatase (Sigma). Both digested DNAs were cleaned by using a Qiagen Miniprep column. Ligations were carried out overnight with T4 DNA ligase (Promega) at 4°C. One microliter of ligation mixture was used as a template in PCR. Amplification was carried out using the Tn5397-specific primers (LEO or REO) and either M13 forward or M13 reverse primer (Table 2). Sequenced PCR products were analyzed using NCBI tools (<http://www.ncbi.nlm.nih.gov/>) and the Expaty molecular biology server (<http://www.expaty.ch/tools/>). Multiple sequence alignments were carried out using ClustalW (<http://www.ebi.ac.uk/clustalW>).

Tn5397 was transferred from *C. difficile* 630 to *E. faecalis* JH2-2 with a mean transfer frequency (± standard deviation [SD]) of 8.85×10^{-8} (± 2.14×10^{-7}) transconjugants per recipient (average of 5 independent experiments). All 5 transconjugants analyzed were generated independently from different filter-mating experiments to exclude the possibility of analyzing siblings. All tetracycline-resistant transconjugants were positive for the *tndX* determinant of Tn5397 (22). The *tndX* gene is unique to Tn5397, the product of which is required and sufficient for its excision and insertion (22). Southern blot analysis showed that a single copy of Tn5397 had inserted in the same site in all *E. faecalis* transconjugants (results not shown). Determination of the insertion site of Tn5397 in *E. faecalis*, using sspPCR, showed that Tn5397 had inserted into a site within the JH2-2 chromosome which has identity to an open reading frame (ORF) from *E. faecalis* V583 encoding

a IIA component of a mannose/sorbose-specific sugar phosphotransferase system (NCBI accession number NP_814245) involved in the uptake and phosphorylation of carbohydrates. The sequences of the target site and the joint of the circular form of Tn5397 in *E. faecalis* are shown in Fig. 1. PCR analysis for the transposon-genome junction for all 5 transconjugants confirmed that the element had entered this site in all transconjugants analyzed.

One transconjugant, EF20A, was used as a donor with *C. difficile* R20291 as the recipient. Tn5397 transferred at a frequency of 6.15×10^{-8} transconjugants per donor or 4.44×10^{-6} transconjugants per recipient. Tn5397 usually occupies a single specific site in the *C. difficile* 630 genome, whereas in R20291, there are two specific sites (Fig. 1). We confirmed by PCR that Tn5397 insertion in *C. difficile* R20291 transconjugants had occurred at one of these sites (results not shown).

This is the first study to demonstrate reciprocal genetic exchange between *E. faecalis* and *C. difficile*. The fact that Tn5397 has a single preferred target site is similar to the situation in *C. difficile* 630 but different from that in *B. subtilis*, in which insertion occurs at multiple sites (21, 23). Demonstration of two-way genetic transfer between *E. faecalis* and *C. difficile* in the laboratory environment represents an ideal situation for gene transfer, and this setting is different from the usual environment of the bacteria; however, these results highlight the need for continual monitoring of emerging resistances in both *C. difficile* and *E. faecalis*, specifically for resistance to clinically important vancomycin, clindamycin, and fluoroquinolones.

TS in CD630 & R20291	TTGTATATGTTTCATCCTTTTGTAGT	GA TGGTAATGGAAGAACATCAAGAGCC
TS in JH2-2 and EF20A	ATGTCCCACGCTTGTATTAGCT	GA CTGATTTAAAAGGAGGAACACCTTG
TS2 in R20291	TTGAATTGATACATCCATTTCAA	GA TGGTAATGGTCGCATAGGTAGATT
LE in EF20A	ATGTCCCACGCTTGTATTAGCT	GA TGGAATGTACCATCAAGACACCTG
RE in EF20A	AGTGTCTCATTGATACATTCTCT	GA CTGATTTAAAAGGAGGAACACCTTG
CI Joint in EF20A	AGTGTCTCATTGATACATTCTCT	GA TGGAATGTACCATCAAGACACCTG

FIG. 1. Target sites of Tn5397 within *C. difficile* 630 (CD630), *C. difficile* R20291, and *E. faecalis* JH2-2. Tn5397 sequence is shown in boldface. TS, target site in the genome; LE, left end of Tn5397; RE, right end of Tn5397; CI, circular intermediate of Tn5397; TS2, the second target site in *C. difficile* R20291. All sequences are derived from PCR amplicons.

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