Antibiotic-Induced, Increased Conjugative Transfer Is Common to Diverse Naturally Occurring ESBL Plasmids in Escherichia coli
Liu, Gang; Bogaj, Karolina; Bortolaia, Valeria; Olsen, John Elmerdahl; Thomsen, Line Elnif

Published in:
Frontiers in Microbiology

DOI:
10.3389/fmicb.2019.02119

Publication date:
2019

Document version
Publisher's PDF, also known as Version of record

Document license:
CC BY

Citation for published version (APA):
Liu, G., Bogaj, K., Bortolaia, V., Olsen, J. E., & Thomsen, L. E. (2019). Antibiotic-Induced, Increased Conjugative Transfer Is Common to Diverse Naturally Occurring ESBL Plasmids in Escherichia coli. Frontiers in Microbiology, 10, [2119]. https://doi.org/10.3389/fmicb.2019.02119
Antibiotic-Induced, Increased Conjugative Transfer Is Common to Diverse Naturally Occurring ESBL Plasmids in *Escherichia coli*

Gang Liu¹, Karolina Bogaj¹, Valeria Bortolaia², John Elmerdahl Olsen¹ and Line Elnif Thomsen¹*

¹ Department of Veterinary and Animal Sciences, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark; ² National Food Institute, Technical University of Denmark, Lyngby, Denmark

Previously, we showed that cefotaxime (CTX) exposure increases conjugative transfer of a *blaCTX-M-1* encoding IncI1 plasmid (IncI1/pST49/CTX-M-1) in *Escherichia coli* in a SOS-independent manner. This study aimed at investigating whether the observation was unique for that plasmid/strain/antibiotic combination or whether antibiotic-induced plasmid transfer (PT) is a more general phenomenon among plasmids in *E. coli*. Whole genome sequences of 25 *E. coli* strains were analyzed to identify different extended spectrum beta-lactamases (ESBL) plasmids enabling selection of a diverse collection of plasmids. Experiments were performed following exposure of these strains to 1/2 minimal inhibitory concentration (MIC) of CTX, ampicillin (AMP), or ciprofloxacin (CIP) before conjugation experiments. The frequency of PT was measured and compared to that of donors not exposed to antibiotics. Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was used to measure mRNA levels of five PT genes and two SOS response genes in donors exposed to antibiotics. The frequency of PT was measured and compared to that of donors not exposed to antibiotics. Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was used to measure mRNA levels of five PT genes and two SOS response genes in donors exposed to antibiotics. The PT of eight strains (30.8% of strains tested) with IncI1/pST7/CTX-M-1, IncI1/pST49/CTX-M-1, IncI1/pST3/CTX-M-1, IncI1/pST293/CTX-M-1, IncI1/pST295/CTX-M-1, IncI1/pST16/CTX-M-55, and IncFII/CTX-M-14 (*n* = 2) plasmids was significantly increased following antibiotic exposure. CTX increased PT in all of these eight strain/plasmid combinations, AMP and CIP increased the PT in six and three strains, respectively. RT-qPCR showed that PT genes were up-regulated in the presence of the three antibiotics, whereas SOS response genes were up-regulated only following CIP exposure. Our findings reveal that antibiotics can increase PT in *E. coli* strains with various ESBL plasmids. Thus, antibiotic-induced conjugative transfer of ESBL plasmids appears to be a common phenomenon in *E. coli*, having important implications for assessing the risks of antibiotic use.

**Keywords:** antibiotic resistance, extended spectrum beta-lactamases, plasmid transfer, antibiotic induced conjugation, *Escherichia coli*
INTRODUCTION

Conjugation is one of the most important mechanisms for intra- and inter-species horizontal gene transfer (HGT), and it plays a significant role to accelerate the dispersal of antibiotic resistance genes (Mazel and Davies, 1999; Barlow, 2009; Halary et al., 2010). Integrative and conjugative elements (ICEs) and conjugative plasmids are both mobile genetic elements that move between cells using conjugation machinery, and the major difference lies in their respective maintenance mechanism within a bacteria cell, as plasmids are extrachromosomal autonomously replicating elements, while ICEs must integrate into chromosomes for stable inheritance (Johnson and Grossman, 2015). Bacterial plasmid conjugation is primarily mediated by type IV secretion systems (T4SSs) and involves direct cell-to-cell contact, mating-pair formation (Mpf), relaxosome formation, and DNA exchange (de la Cruz et al., 2010).

Extended-spectrum beta-lactamase (ESBL) production is a common resistance mechanism against beta-lactam antibiotics in Gram-negative bacteria (Al-Bayssari et al., 2015), and the rapid dissemination of novel genes reflects their evolution under the selective pressure of antibiotic usage. ESBL genes are generally acquired by HGT, and they are able to hydrolyze cephalosporins with an oximino side chain (Malloy and Campos, 2011). Some ESBL genes are mutant derivatives of established plasmid-mediated beta-lactamases (e.g., blatem/SHV) or mobilized from environmental bacteria (e.g., blactx-M) (Overdevest et al., 2011). Many Enterobacteriaceae now carry ESBL genes such as blactx-M, with particular genes associated with different geographical regions (Hawkey and Jones, 2009).

It is generally assumed that antibiotics can enhance HGT, and also a substantial amount of data suggest that sub-inhibitory concentrations of antibiotics may significantly increase the conjugation transfer frequency both in vitro and in the animal gut (Aminov, 2011). For instance, sub-inhibitory concentrations of beta-lactams and tetracycline enhanced the conjugative transfer of a tetracycline resistance plasmid in Staphylococcus aureus or the conjugative transposon CTnDOT in Bacteroides cells, respectively (Barr et al., 1986; Stevens et al., 1993; Whittle et al., 2002). The presence of tetracycline or erythromycin increased the transfer frequency of conjugative transposon Tn916 or plasmid pLFE1 in the guts of gnotobiotic rats, respectively (Bahl et al., 2002).

Furthermore, ciprofloxacin treatment has been shown to enhance the transfer of the ICE SXT, which confer some antibiotic resistance genes (Mazel and Davies, 1999; Barlow, 2009; Halary et al., 2010). Integrative and conjugative elements (ICEs) and conjugative plasmids are both mobile genetic elements that move between cells using conjugation machinery, and the major difference lies in their respective maintenance mechanism within a bacteria cell, as plasmids are extrachromosomal autonomously replicating elements, while ICEs must integrate into chromosomes for stable inheritance (Johnson and Grossman, 2015). Bacterial plasmid conjugation is primarily mediated by type IV secretion systems (T4SSs) and involves direct cell-to-cell contact, mating-pair formation (Mpf), relaxosome formation, and DNA exchange (de la Cruz et al., 2010).

Materials and Methods

Bacterial Strains

Twenty-five ESBL-producing E. coli strains collected in Denmark in 2015 were included in the study. These isolates were collected and whole genome sequenced (WGS) within the Danish programme for surveillance of antimicrobial resistance (DANMAP) 2015 (see Supplementary Table 1 for ENA accession information: PRJEB14641 and PRJEB14086), and originated from pigs’ cecal samples (n = 13), calves’ cecal samples (n = 6), and chicken meat (n = 6) (Table 1). They were selected among the animal- and food-origin ESBL-producing E. coli from DANMAP 2015 to include all detected ESBL genes and all possible plasmids harboring such genes based on a rapid preliminary analysis of beta-lactamase-encoding genes and plasmid replicons detected by using the Center for Genomic Epidemiology (CGE)1 tools. E. coli K-12 MG1655/pTF2 (Kjeldsen et al., 2015) with the IncI1 plasmid pTF2 carrying blactx-M (IncI1/pST49/CTX-M-1) was also used in this study. E. coli J53-1 (NalR) (Taylor and Levine, 1980) and E. coli J53-2 (RifR) (Appelbaum et al., 1972) were used as recipients in conjugation experiments.

Bioinformatics Analysis

Sequencing reads were quality trimmed using Trimmomatic-v0.32 (Bolger et al., 2014) and de novo assembled using SPAdes v.3.5.0 (Bankevich et al., 2012). The assembled genomes were analyzed via the CGE (see text footnote 1) webserver by using a bacterial analysis pipeline (BAP) (Thomsen et al., 2016). Multi-locus sequence types were determined using MLST 2.0 with the allele sequence and profile data from PubMLST2. Other genotypic features such as plasmid replicons, antimicrobial resistance genes, and virulence genes were also identified with a selected threshold equal to 98% identity using the pipelines, PlasmidFinder 2.0, VirulenceFinder 2.0, and ResFinder 3.2 at CGE. The sequences were analyzed for the presence or absence of tra genes, using BLAST in CLR Main Workbench.

1https://cge.cbs.dtu.dk/services/
2https://pubmlst.org
| Isolate name<sup>a</sup> | β-Lactamase (bla-type) | MLST type | Plasmid replicons<sup>b</sup> | pMLST<sup>d</sup> | Antimicrobial resistance genes<sup>k</sup> | Virulence genes |
|----------------------|------------------------|-----------|-------------------------------|-----------------|------------------------------------------|----------------|
| ESBL 1               | CTX-M-1                | ST101     | IncI1                         | 3               | dfrA17, aadA1, aadA5, bl<sub>TEM</sub>-1, sul<sub>1</sub>, sul<sub>2</sub>, tet(A),  catA<sup>1</sup> | gad, iroN, iss, lipA, mchB, mchC, mchF, mcmA |
| ESBL 2               | CTX-M-1                | ST4373    | IncFII, IncFIB(AP001918)       | –               | dfrA14, strA<sup>1</sup>, strB, bl<sub>TEM</sub>-1, bl<sub>STR</sub>-1A, sul<sub>2</sub>, tet(B) | air, eilA, gad, iroN, lipA |
| ESBL 3               | CTX-M-1                | ST4580    | IncI1, ColpVC, pO111           | 7               | dfrA1, aadA1, bl<sub>CTX</sub>-M-1, sul<sub>1</sub>, sul<sub>2</sub> | ctnf<sup>1</sup>, gad, iroN, iss, lipA, mchB, mchC, mchF, mcmA |
| ESBL 4               | CTX-M-1                | ST58      | IncI1                         | 7               | dfrA1, aadA1, bl<sub>CTX</sub>-M-1, sul<sub>1</sub> | gad, iroN, lipA |
| ESBL 5               | CTX-M-1                | ST165     | IncFII(pRSB107), pO111         | –               | bl<sub>CTX</sub>-M-1, tet(A), erm(B), mph(A) | gad |
| ESBL 6               | CTX-M-1                | ST295     | IncFII(Fil), IncFIB(AP001918)  | –               | dfrA8, aadA1, bl<sub>CTX</sub>-M-1, mph(A) | iroN, iss, lipA, mchF, tsh |
| ESBL 7               | CTX-M-1                | ST362     | IncFIL, IncI1, IncQ1<sup>f</sup> | NT<sup>★</sup> | dfrA<sup>1</sup>, dfrA<sup>17</sup>, aph(3')-la, str<sup>B</sup>, aprA<sup>4</sup>-la, aac(3)-Ia, aac(3)-Ia, aadA<sup>1</sup>, strA, aadA<sup>5</sup>, bl<sub>CTX</sub>-M-1, bl<sub>TEM</sub>-1B, sul<sub>2</sub>, tet(A), catA<sup>1</sup> | ireA, iss, mcmA |
| ESBL 8               | CTX-M-1                | ST345     | IncI1, IncFIB(AP001918), Col156 | 3               | dfrA<sup>17</sup>, strB<sup>1</sup>, aadA<sup>5</sup>, bl<sub>CTX</sub>-M-1, bl<sub>TEM</sub>-1C, sul<sub>1</sub>, sul<sub>2</sub>, tet(A) | ctnf<sup>1</sup>, gad, iroN, iss, lipA, mchF |
| ESBL 9               | CTX-M-1                | ST101     | IncFII(29), IncI1              | NT<sup>★</sup> | dfrA<sup>17</sup>, aadA<sup>1</sup>, aadA<sup>5</sup>, bl<sub>CTX</sub>-M-1, sul<sub>1</sub>, sul<sub>2</sub>, tet(A), catA<sup>1</sup> | cba, cma, gad, iroN, iss, lipA, mchB, mchC, mchF, mcmA |
| ESBL 10              | CTX-M-1                | ST647     | IncFII(pSE111), IncI1, IncFIB(AP001918) | 29<sup>3</sup>f | dfrA<sup>17</sup>, strB<sup>1</sup>, aadA<sup>5</sup>, bl<sub>CTX</sub>-M-1, sul<sub>1</sub> | cba, cdtB, cma, gad, iroN, iss, sfa<sup>5</sup>, vat |
| ESBL 11              | CTX-M-1                | ST48      | IncFII(29), IncI1, IncY, IncFIB(AP001918), Col(MP18) | 29<sup>4</sup>g | aadA<sup>1</sup>, bl<sub>CTX</sub>-M-1, sul<sub>2</sub> | ND<sup>n</sup> |
| ESBL 12              | CTX-M-1                | ST1640    | IncFII, IncI1, IncN, IncFIB(AP001918), IncQ1<sup>e</sup>, ColpVC | 29<sup>5</sup>a | dfrA<sup>14</sup>, dfrA<sup>17</sup>, gyr<sup>S</sup>1, gyrA<sup>B</sup>83<sup>L</sup>, str<sup>B</sup>, strA<sup>1</sup>, aadA<sup>5</sup>, bl<sub>CTX</sub>-M-1, bl<sub>TEM</sub>-1B, sul<sub>2</sub>, tet(A), catA<sup>1</sup> | astA, cba, cdtB, cma, gad, hlyE, iha, ireA, iss |
| ESBL 13              | CTX-M-1                | ST4243    | IncI<sup>e</sup>, ColpVC, IncB/O/K/Z | NT<sup>★</sup> | dfrA1, bl<sub>CTX</sub>-M-1, bl<sub>TEM</sub>-1B, sul<sub>2</sub> | air, astA, cma, eilA, gad, ighA |
| ESBL 14              | CTX-M-1                | ST117     | IncFII, IncI1, IncFIB(AP001918), ColpVC | 29<sup>6</sup>f | dfrA<sup>17</sup>, aadA<sup>5</sup>, sul<sub>2</sub>, bl<sub>CTX</sub>-M-1 | gad, hlyE, ireA, iss, lipA, vat, mchB, mchC, mchF, mcmA |
| ESBL 15              | CTX-M-1                | ST752     | IncFII(Fil), IncI1, IncFIB(AP001918), pO111 | 7 | strA, strB, bl<sub>CTX</sub>-M-1 | astA, cfr, eae, espA, espF, espP, gad, iroN, iss, rieB, tcp<sup>P</sup>, tv, tsh |
| ESBL 16              | CTX-M-1                | ST88      | IncFII, IncI1, Col(MG8280), IncFIB(AP001918) | 7 | dfrA1, strA, strB, aadA1, bl<sub>CTX</sub>-M-1, bl<sub>TEM</sub>-1C, sul<sub>1</sub>, sul<sub>2</sub>, tet(A) | gad, iroN, iss, lipA, mchF |
| ESBL 17              | CTX-M-1                | ST101     | IncFII, IncI1, IncFIB(AP001918) | 3 | dfrA5, dfrA17, aadA5, bl<sub>CTX</sub>-M-1, sul<sub>2</sub> | gad, iroN, iss, lipA, mchF |
| ESBL 18              | TEM-52                 | ST10      | IncRA, IncFII, IncFIB(AP001918), IncX1 | – | bl<sub>TEM</sub>-52B | astA, gad, iss |
| ESBL 19              | TEM-52                 | ST10      | IncRA, IncFII, IncX1, IncFIB(AP001918) | – | bl<sub>TEM</sub>-52B | astA, gad, iss |
| ESBL 20              | TEM-52                 | ST2607    | IncFII(Fil), IncI1, IncN, IncFIB(AP001918) | – | aadA<sup>17</sup>, bl<sub>TEM</sub>-52B | gad, iss, lipA |
| ESBL 21              | CTX-M-55               | ST2952    | IncI<sup>e</sup>, IncX1         | 16 | bl<sub>CTX</sub>-M-55 | gad, lipA |

<sup>a</sup> Isolate name: E. coli (bla-type) | β-Lactamase (bla-type) | MLST type | Plasmid replicons | pMLST | Antimicrobial resistance genes | Virulence genes |

*(Continued)*
TABLE 1 | Continued

| Isolate name<sup>a</sup> | β-Lactamase (bla-type) | MLST type | Plasmid replicons<sup>b</sup> | pMLST<sup>d</sup> | Antimicrobial resistance genes<sup>k</sup> | Virulence genes |
|-------------------------|------------------------|-----------|---------------------------|----------------|---------------------------------|----------------|
| ESBL 22                 | CTX-M-14               | ST1310    | IncFiI(29), IncFIA, IncFIB(AP001918), IncB/O/K/Z, IncQ1<sup>e</sup> | –              | bla<sub>CTX</sub>-M-14           | gad, iss, kpfA |
| ESBL 23                 | CTX-M-14               | ST88      | IncFl, IncFIB(AP001918), IncB/O/K/Z, IncQ1<sup>e</sup> | –              | strA, strB, bla<sub>CTX</sub>-M-14, bla<sub>TEM</sub>-1B, sul2, tet(A) | cnf1, gad, icaN, iss, kpfA, mchF |
| ESBL 24                 | CTX-M-14               | ST34      | IncI1, IncQ1<sup>e</sup>, ColM(G328B), ColIPC | 259            | gnrB19<sup>f</sup>, bla<sub>CTX</sub>-M-14, bla<sub>TEM</sub>-1B, sul2, tet(A), floR<sup>f</sup> | ND<sup>m</sup> |
| ESBL 25                 | SHV-12                 | ST115     | IncFl, IncFIB(AP001918), pO111 | –              | gnrB19, bla<sub>SHV</sub>-12 | air, ceb, elA, ica, iss |

<sup>a</sup>Original ID as recorded in European Nucleotide Archive (ENA) is provided in Supplementary Table 1. Isolates for which ESBL plasmid transfer significantly increased upon antibiotic exposure are underlined. <sup>b</sup>The replicons of the plasmids associated with the ESBL gene are underlined. <sup>c</sup>Only part of the replicon was detected. <sup>d</sup>pMLST was performed on IncI1. <sup>e</sup>No detection of the sogS allele. The remaining allele variants were repI1, 2; ardA, 1; trbA, 4; pilL, 2. <sup>f</sup>New pMLST identified in this study. The allele variants were repI1, 2; ardA, 4; pilL, 2. <sup>g</sup>New pMLST identified in this study. The allele variants were repI1, 2; ardA, 1; trbA, 4; pilL, 2. <sup>h</sup>Antimicrobial resistance phenotype: aminoglycoside [aadA1, aadA5, aadA17, strA, strB, aph(3′)-Ia, aph(4)-Ia, aac(3)-Iva], quinolone (gyrA S83L, qnrS1, qnrB19), trimethoprim (dfrA1, dfrA5, dfrA8, dfrA14, dfrA17), beta-lactam (bla<sub>CTX</sub>-M-1, bla<sub>CTX</sub>-M-14, bla<sub>CTX</sub>-M-55, bla<sub>TEM</sub>-1A, bla<sub>TEM</sub>-1B, bla<sub>TEM</sub>-1C, bla<sub>TEM</sub>-GE, bla<sub>SHV</sub>-12), sulfonamide (sul1, sul2), tetracycline (tetA, tetB), macrolide (mphA, ermB), and phenicol (catA1, floR). <sup>i</sup>Genes with < 100% match with the reference in ResFinder database. <sup>j</sup>ND, not identified.

8.0.0 (CLC Bio, Denmark). To reconstruct ESBL plasmids, a curated dataset of complete plasmid sequences (Orlek et al., 2017) was used to identify the best reference plasmid for each strain using (i) information on replicon, pMLST, and ESBL gene content; and (ii) highest percentage of ID over highest percentage of length as obtained using nucleotide BLAST at NCBI. Using the best reference plasmid, contigs were assigned to a plasmid scaffold within each strain using nucleotide BLAST at NCBI.

Antibiotic Susceptibility Testing

The MIC of CTX and AMP was determined on donors and transconjugants from the first conjugation round (see below) using the broth microdilution method [concentrations ranging from 0 to 2048 mg L<sup>−1</sup> (CTX) or 0 to 4096 mg L<sup>−1</sup> (AMP) by twofold dilution increases] in poly-styrene microtiter plates using the control strain E. coli ATCC® 25922 according to the CLSI guidelines M100-S25 (Wayne et al., 2015). In addition, MIC determination experiments were performed using serial increases of 200 mg L<sup>−1</sup> AMP to obtain a value as close to the real MIC as possible. The MIC of CIP was measured by using ciprofloxacin single pack E-test (Biomerieux, United States) (concentration ranging from 0.002 to 32 mg L<sup>−1</sup>) following the manufacturer’s protocol. MIC values for donors and transconjugants are available in Supplementary Table 2.

Conjugation Experiment

The 25 strains containing different ESBL plasmids and MG1655/pTF2 (Kjeldsen et al., 2015) were used as donor strains and E. coli J53-2 as recipient strain in the first round of conjugation. In the second round, transconjugants from the first round of conjugation were used as donors and E. coli J53-1 as recipient. Strains were exposed to 1/2 MIC of antibiotics. Conjugation was performed as previously described (Moller et al., 2017). Briefly, donor strains were grown either without or with sub-inhibitory concentrations (1/2 MIC) of antibiotics in Luria-Bertani (LB) broth (Sigma, Copenhagen, Denmark) with shaking (125 rpm) at 37°C to exponential phase (OD<sub>600</sub> = 0.5), and the recipient strain was grown without antibiotics to the same OD<sub>600</sub>. Antibiotics were removed by a washing step and conjugation was performed by mixing donor and recipient strain in a 1:1 ratio on filters (0.22 µM, Millipore, Copenhagen, Denmark) on LB agar plates (Becton, Dickinson, Albertslund, Denmark) at 37°C for 30 and 60 min. The bacterial material was washed from the filters using isotonic NaCl and plated on LB agar plates containing (in the first conjugation round): 2 mg L<sup>−1</sup> CTX (for quantifying donor + transconjugants) or 50 mg L<sup>−1</sup> rifampicin and 2 mg L<sup>−1</sup> CTX (quantifying transconjugants only), or grown on LB agar plates containing (in the second conjugation round): 2 mg L<sup>−1</sup> CTX and 50 mg L<sup>−1</sup> rifampicin (quantifying donor) or 10 mg L<sup>−1</sup> nalidixic acid and 2 mg L<sup>−1</sup> CTX (quantifying transconjugants only), and incubated overnight at 37°C. The conjugation frequency was calculated as the number of transconjugants divided by the number of donors. The conjugation experiments were performed in three biological replicates with three technical replicates each.

Plasmid Profiling and PCR-Based Replicon Typing

Transconjugants from the first conjugation round showing significant antibiotic-induced increase in PT and their respective donors were selected for plasmid profiling. Plasmid DNA was purified according to the method described by Kado and Liu (1981), followed by separation on a 0.7% (W/V) agarose gel for 4 h at 150 V and stained with GelRed (Biotium, Hayward, CA, United States). The approximate molecular weight of each plasmid was determined by comparison with two reference E. coli strains, 39R861 and V517, containing multiple reference plasmids (Macrina et al., 1978; Karczmarczyk et al., 2011).
The plasmid replicon types of the ESBL plasmids were confirmed in the transconjugants from the first conjugation round using PCR-Based Replicon Typing (PBRT) 2.0 Kit (Diatheva, Fano, Italy). The total DNA was obtained by boiling lysis method exposing cells to 100°C for 10 min, and clarifying the lysate preparations by centrifugation at 15,000 × g for 10 min. One microliter of the supernatant was tested in PCR assays and the amplification products were analyzed by 2.5% agarose gel electrophoresis as described previously (Carattoli et al., 2005; Carloni et al., 2017).

Plasmid Sequencing
Plasmid DNA was isolated from the transconjugants from the first conjugation round showing significant antibiotic-induced increase in PT using PureLink HiPure Plasmid DNA Purification Kits (Invitrogen, CA, United States), and the plasmid DNA was subjected to pair-end-read format using the Nextera XT DNA Library Preparation Kit and sequenced in an Illumina MiSeq (Illumina, Inc., San Diego, CA, United States) essentially as previously described (McElhoe et al., 2014). Obtained sequences were analyzed by PlasmidFinder and BLAST in CLC Main Workbench 8.0.0 (CLC Bio, Denmark) to confirm presence of the ESBL plasmids.

RNA Extraction
Single colonies of transconjugants from the first conjugation round were grown in LB broth with shaking (125 rpm) overnight at 37°C. The cultures were diluted 1000-fold and grown with and without antibiotics to OD600 = 0.5. The antibiotic concentration was 1/2 MIC. A FastPrep cell disrupter system (Qbiogene, Illkirch, France) and RNeasy Mini Kit (Qiagen, Sollentuna, Sweden) were used to extract total RNA by mechanical disruption. Quantity of the extracted RNA was determined by A260 measurements and purity by A260/280 ratio measurements using a NanoDrop 1000 spectrophotometer (Thermo Scientific, Hvidovre, Denmark). RNA (1 μg) samples were purified by DNA digestion using TURBO™ DNase kit (2 U/μL) (Ambion, Life Technologies, Nærum, Denmark) to remove contaminating genomic DNA.

Reverse-Transcribed-Quantitative Real Time Polymerase Chain Reaction
RNA was reverse-transcribed into cDNA using the High Capacity cDNA Reverse Transcription Kit (Life Technologies, Nærum, Denmark). Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was performed using FastStart Essential DNA Green Master (Roche, Hvidovre, Denmark) and a LightCycler 96 (Roche, Hvidovre, Denmark). Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was performed using FastStart Essential DNA Green Master (Roche, Hvidovre, Denmark). Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was performed using FastStart Essential DNA Green Master (Roche, Hvidovre, Denmark). Reverse-transcribed-quantitative real time polymerase chain reaction (RT-qPCR) was performed using FastStart Essential DNA Green Master (Roche, Hvidovre, Denmark).

RESULTS
Analysis of 25 E. coli Strains and Their Plasmids
With the aim of investigating PT of different plasmid types in various E. coli strains, we obtained 25 ESBL-producing E. coli strains isolated from pigs, calves, and chicken meat. A whole genome sequence analysis was undertaken, and a total of 19 MLST types were identified among the E. coli strains (Table 1). Of the 25 strains, 17 carried blaCTX−M−1, 3 blaTEM−52B, 3 blaCTX−M−14, 1 blaCTX−M−55, and 1 blashv−12. In 20 strains, genes conferring resistance to additional non-beta-lactam antibiotics were detected. Most (n = 23) strains harbored multiple plasmid replicons (Table 1). Plasmid multilocus sequence typing (pMLST) was performed on IncI1 plasmids, leading to identification of eight pMLST types including four newly described in this study (pST293-296) that were submitted to the Plasmid MLST Databases. Three IncI1 plasmids were untypeable as one or two alleles were not detected (Table 1).

Antibiotics Increase Conjugation Transfer Frequency
Conjugation transfer frequency was investigated using the 25 diverse strains and the previously investigated MG1655/pTF2 was included as positive control (Moller et al., 2017). PT was affected by antibiotic exposure in 8 (30.8%) out of the 26 strains. These carried IncI1/pST7/CTX−M−1, IncI1/pST49/CTX−M−1 (MG1655/pTF2), IncI1/pST293/CTX−M−1 (ESBL 10), IncI1/pST295/CTX−M−1 (ESBL 12), IncI1/pST3/CTX−M−1 (ESBL 17), IncI1/pST16/CTX−M−55 (ESBL 21), and IncFI/CTX−M−14 (n = 2, ESBL 22 and ESBL 23) (Table 2). Pre-growth with 1/2 MIC of CTX increased the PT significantly for all of these eight strains after both 30 and 60 min of cell contact compared to conjugation experiments in which the donor was grown without CTX before the conjugation. Additionally, growth with 1/2 MIC of AMP increased the PT significantly in three of the eight strains (MG1655/pTF2, MG1655/pTF2, MG1655/pTF2, and MG1655/pTF2, respectively).

The expression data were normalized to two validated reference genes, gapA and nusG (Kjeldsen et al., 2015). RT-qPCR was performed on two biological replicates with two technical replicates each, and the results were calculated by the 2−ΔΔCt method (Pfaffl, 2001).

Statistical Analysis
Statistical analysis used in this study was performed using GraphPad Prism version 7.03. Data are expressed as the means ± standard deviation (SD) from at least three independent experiments. Comparison of gene expression and conjugation frequencies with and without antibiotics were analyzed using Student's t-test with Welch’s correction. P-values ≤ 0.05 were considered statistically significant.

http://pubmlst.org/plasmid/
ESBL 4, and ESBL 22). A non-significant CIP-induced PT was observed for three other strains out of the eight (ESBL 10, 12, and 21) (Table 2). We obtained transconjugants from eight other strains (ESBL 1, 5, 6, 8, 11, 14, 16, and 18), but observed no significant antibiotic induced PT using any of the three antibiotics (Supplementary Figure 1). We did not obtain any transconjugants either without or with 1/2 MIC of AMP, CTX, or CIP exposure for the remaining 7 blaCTX-M-positive, 2 blaTEM-52R-positive, and 1 blaSHV-12-positive strains used in this study (ESBL 2, 3, 7, 9, 13, 15, 19, 20, 24, and 25). When analyzing these 10 strains from which no transconjugants were observed, the tral gene was absent in 7 strains (ESBL 2, 7, 9, 13, 19, 20, and 25), and tral, tral, and traM lacked in ESBL 19. The transconjugants were named after the ESBL number of the original donor, followed by the antibiotic used in the conjugation experiment.

Plasmid profiling and PBRT and sequencing of the transconjugants from the first conjugation round confirmed the presence of the replicons of the ESBL plasmids identified by WGS data analysis and allowed further insights into the IncI1 replicons. PBRT revealed that IncI1α and IncI1γ replicons co-existed in two transconjugants, namely 10AMP and 10CIP (transconjugant originating from ESBL10 treated with AMP or CIP, respectively). The occurrence of both replicons was verified in the donor (ESBL10) and transconjugants by analysis of the WGS data, and was also confirmed by plasmid profiling (data not shown). It is interesting to note that in transconjugant 10CTX, only the IncI1α replicon was transferred. An additional case in which the transconjugants harbored other plasmids besides the ESBL plasmid was represented by 12AMP, which harbored a non-typeable plasmid and a ColI56 plasmid besides the IncI1α ESBL encoding plasmid. In all remaining cases, the transconjugants harbored only the ESBL encoding plasmids which were either IncI1α (4CTX, 12CTX, 12AMP, 17CTX, 17AMP, 17CIP, 21CTX, 21AMP, and 21CIP) or IncFII (22CTX, 22AMP, 22CIP, 23CTX, 23AMP, and 23CIP).

### Table 2: Fold changes of antibiotic (CTX, AMP, or CIP)-induced increased conjugation transfer frequency using ESBL E. coli as donors and JS3-2 as recipient.

| Donor | Conjugation time (min) | Conjugation transfer frequencya (Control) | Conjugation transfer frequencya (CTX) | Fold changeb | Conjugation transfer frequencya (AMP) | Fold changeb | Conjugation transfer frequencya (CIP) | Fold changeb |
|-------|------------------------|------------------------------------------|--------------------------------------|-------------|--------------------------------------|-------------|--------------------------------------|-------------|
| MG1655/pTF2 (IncI1/CTX-M-1) | 30 | 4.63 ± 10-3 | 2.93 ± 10-2 | 6.3** | 1.30 ± 10-2 | 2.8* | 5.73 ± 10-3 | 1.2 |
| | 60 | 1.11 ± 10-2 | 0.23 ± 10-2 | 3.7** | 6.11 ± 10-2 | 5.5* | 6.97 ± 10-2 | 6.3* |
| ESBL 4 (IncI1/CTX-M-1) | 30 | 4.11 ± 10-4 | 1.04 ± 10-3 | 2.5 | 3.63 ± 10-3 | 8.8 | 1.39 ± 10-3 | 4.7 |
| | 60 | 5.71 ± 10-4 | 0.38 ± 10-3 | 3.3* | 6.53 ± 10-3 | 11.3* | 4.67 ± 10-3 | 8.1* |
| ESBL 10 (IncI1/CTX-M-1) | 30 | 1.04 ± 10-2 | 3.62 ± 10-2 | 3.5 | 2.93 ± 10-2 | 2.8* | 5.73 ± 10-2 | 2.3 |
| | 60 | 4.11 ± 10-2 | 1.22 ± 10-2 | 8.7* | 2.97 ± 10-2 | 2.5* | 5.26 ± 10-2 | 4.5 |
| ESBL 12 (IncI1/CTX-M-1) | 30 | 4.17 ± 10-4 | 1.77 ± 10-3 | 12.0* | 6.63 ± 10-4 | 3.3 | 1.47 ± 10-4 | 1.7 |
| | 60 | 1.09 ± 10-4 | 0.17 ± 10-4 | 1.2 | 3.35 ± 10-5 | 3.8 | 0.19 ± 10-4 | 3.8 |
| ESBL 17 (IncI1/CTX-M-1) | 30 | 2.61 ± 10-5 | 5.61 ± 10-5 | 2.1* | 3.55 ± 10-5 | 1.2 | 3.20 ± 10-5 | 1.2 |
| | 60 | 1.07 ± 10-5 | 0.20 ± 10-4 | 1.0 | 4.20 ± 10-5 | 0.8 | 3.29 ± 10-5 | 0.8 |
| ESBL 21 (IncI1/CTX-M-55) | 30 | 2.73 ± 10-3 | 8.96 ± 10-3 | 3.3 | 8.03 ± 10-3 | 3.4 | 9.34 ± 10-3 | 3.4 |
| | 60 | 5.41 ± 10-3 | 1.62 ± 10-2 | 3.3* | 1.45 ± 10-2 | 2.4 | 1.30 ± 10-2 | 2.4 |
| ESBL 22 (IncF/CTX-M-14) | 30 | 1.61 ± 10-4 | 5.91 ± 10-4 | 3.7* | 9.3 ± 10-4 | 6.0* | 9.67 ± 10-4 | 6.0* |
| | 60 | 0.94 ± 10-4 | 1.81 ± 10-4 | 1.1 | 2.7 ± 10-3 | 4.5* | 0.94 ± 10-4 | 4.5* |
| ESBL 23 (IncF/CTX-M-14) | 30 | 0.27 ± 10-4 | 7.35 ± 10-4 | 2.7* | 3.11 ± 10-4 | 1.1 | 2.87 ± 10-4 | 1.1 |
| | 60 | 0.17 ± 10-4 | 0.55 ± 10-4 | 0.9 | 0.28 ± 10-4 | 0.22 | 0.24 ± 10-4 | 0.22 |

*The results shown are from two biological replicates with three technical replicates each, the data shown represent the mean ± standard deviation. bThe asterisks indicate statistical significance between antibiotic treated and control at level: *p ≤ 0.05, **p ≤ 0.01.
Expression of tra Genes Is Induced by Antibiotics in the Transconjugants

Previous data had shown that increased PT was linked to up-regulation of the transfer (tra) genes (Møller et al., 2017). In order to investigate whether the tra genes on the conjugative ESBL encoding plasmids able to transfer were up-regulated in response to exposure to antibiotics, we exposed the transconjugants from the first round of conjugation to the three antibiotics at a concentration of 1/2 MIC. We observed that several tra genes were significantly up-regulated in all of the eight transconjugants when exposed to CTX, AMP, and CIP (Figure 1). To rule out that the observed increase in expression levels was caused by antibiotic induced increases in plasmid copy number, we performed RT-qPCR. We found that the ratio between a plasmid encoded gene (traF) and a chromosomally encoded gene (dhs) did not change significantly due to antibiotic exposure, supporting that the antibiotics do not affect copy-number (Supplementary Figure 2).

To investigate the involvement of the SOS response in antibiotic induced PT, we analyzed the expression of recA and sfiA. RT-qPCR analysis showed that expression of these two genes was not affected significantly by growth in the presence of CTX and AMP (Figure 2). In contrast, CIP treatment increased the expression of recA significantly in all of these eight strains, and increased the level of sfiA significantly in seven of these strains (Figure 2).

Antibiotics Induce Conjugation From Transconjugants

In order to investigate whether the up-regulated tra genes in transconjugants from the first round of conjugation would also lead to increased PT frequency, a second conjugation experiment was performed using the transconjugants as donors and E. coli J53-1 as recipient strain. The experiment showed that the CTX treatment increased significantly PT of IncI1/pST7/CTX-M-1 plasmid (4CTX), IncI1/pST293/CTX-M-1(10CTX), IncI1/pST295/CTX-M-1 (12CTX), IncI1/pST3/CTX-M-1 (17CTX), IncI1/pST16/CTX-M-55 (21CTX), and IncF/CTX-M-14 plasmids (22CTX), compared to conjugation without pre-growth in the presence of CTX (Table 3). The AMP treatment also showed significantly increased PT in IncI1/pST7/CTX-M-1 plasmid (4AMP) and IncI1/pST3/CTX-M-1 (17AMP). Additionally, CIP treatment increased significantly PT of IncI1/pST7/CTX-M-1 plasmid (4CIP), IncI1/pST3/CTX-M-1 (17CIP), IncI1/pST16/CTX-M-55 (21CIP), and IncF/CTX-M-14 plasmids (22CIP). For the two remaining plasmids we observed a, non-significant, increased antibiotic induced PT (Table 3).

DISCUSSION

The increasing occurrence of antibiotic resistance (AMR) among pathogenic bacteria is considered a major problem for public health in recent decades; and conjugation plays a major role in the spread of AMR (Zatyka and Thomas, 1998; Pfaffl, 2001; Sørensen et al., 2005; Bennett, 2008; Norman et al., 2009; Thomsen et al., 2016; von Wintersdorff et al., 2016). Even though many studies have focused on characterizing the impact of antibiotics on PT, the conclusions are not comparable (Barr et al., 1986; Bahl et al., 2004; Feld et al., 2008; Lopatkin et al., 2016). The possible reasons for these divergent results may be caused by different types and concentrations of antibiotics, different plasmids, and strains tested. A recent publication supports this hypothesis, as they also pointed out that antibiotics contribute to the increased HGT, but the level of HGT also relied on which donor and recipient cells were involved (Lu et al., 2017).

Extended spectrum beta-lactamases genes, and in particular CTX-M beta-lactamases, have become the most prevalent cephalosporin-resistance determinants in Enterobacteriaceae (Hawkey and Jones, 2009), and it was considered important to understand whether antimicrobial treatments may contribute to the spread of these genes. Therefore, 25 E. coli strains harboring different naturally occurring ESBL plasmids were selected to perform conjugation experiments. MG1655/pTF2 was included as we have previously shown that CTX could induce PT in this strain (Møller et al., 2017). Our main aim was therefore to investigate the effect of β-lactams on PT of the different ESBL plasmids. We chose to include CIP, as another antibiotic class to see whether we would get similar results, as CIP has previously been shown to increase PT (Shun-Mei et al., 2018). The strains encompassed all diverse ESBL genes and plasmids observed in ESBL positive E. coli from AMR surveillance in livestock in Denmark in 2015, and consequently showed uneven distribution of resistance genes, reflecting that blaCTX-M−1 was the most common ESBL gene in E. coli from livestock (Meunier et al., 2006; Rodríguez et al., 2009). We found that antibiotic treatment significantly increased PT in eight ESBL-producing E. coli. In addition, at least four other blacT×M−1 positive strains with IncI1 plasmid revealed an increased PT upon antibiotic exposure, however, not statistically significantly. We observed a high prevalence of antibiotic induced PT from strains with IncI1 plasmids, however, this was also the most common ESBL plasmid in our collection, present in 18 out of our 26 strains investigated. We observed no PT from 10 strains. Previous studies have shown that traF is essential in the formation of mating apparatus and conjugative transfer; and modification of any region of traF resulted in a loss of conjugal function (Waters et al., 1992; Lento et al., 2016; Huang et al., 2019). Analysis of the 10 strains from which no transconjugants was observed, showed that in 7 strains (ESBL 2, 7, 9, 13, 19, 20, and 25) the traF gene was absent. Additionally, ESBL19 lacked tral, traL, and traM. These deletions may be the cause of the lack of conjugation of these plasmids. Three of the plasmids (from ESBL 3, 15, and 24) encoded the tra genes and could potentially conjugate, but below detection limit. When examining the transconjugants, we investigated expression levels of five transfer genes involved in the conjugation process. traF and traL are involved in the pilus synthesis and assembly system, tral and traM participate in the DNA transfer.
FIGURE 1 | Expression of genes involved in conjugation in the transconjugants from the first round of conjugation. Cell samples grown with no or 1/2 MIC of antibiotics (+CTX, AMP, or CIP) were used for RT-qPCR. Data are presented as fold change relative to control without antibiotics. The combination of strain number with antibiotic name represents the transconjugants from the first round of conjugation obtained after exposure to the specified antibiotic. The expression data were normalized to the reference genes, gapA and nusG. The results shown are means of three biological replicates with two technical replicates each and the error bars represent standard errors of the means. The stars indicate statistical significance at different levels: ∗P ≤ 0.05, ∗∗P ≤ 0.01.
FIGURE 2 | Expression of SOS-responsive genes in the transconjugants from the first round of conjugation. Cell samples grown with no or 1/2 MIC of antibiotics (+CTX, AMP, or CIP) were used for RT-qPCR. Data are presented as fold change relative to control without antibiotics. Note that the different figures are not drawn to the same scale. The combination of strain number with antibiotic name represents the transconjugants from the first round obtained after exposure to the specified antibiotic. The expression data were normalized to two validated reference genes, \textit{gapA} and \textit{nusG}. The results shown are means of three biological replicates with two technical replicates each and the error bars represent standard errors of the means. The stars indicate statistical significance at different levels: $^* P \leq 0.05$, $^** P \leq 0.01$. 
system, and pilS is one of the pilus structural components. These genes were selected as they all play an important role in conjugation and have previously been shown to be up-regulated in bacteria exposed to CTX (Moller et al., 2017). We found these genes to be up-regulated in the transconjugants when exposed to antibiotics. It was furthermore demonstrated that this up-regulation of transfer associated genes correlated with an increase in PT frequency. Although the increased PT for some strains exposed to specific antibiotics did not increase beyond a statistically significant threshold, this non-significant increased PT still correlated to increased expression of the tra genes, supporting that antibiotics also induced PT in these strains (Table 3 and Figure 1). That we could show antibiotic increased PT when using the transconjugants as donors, indicated that antibiotic-increased PT was independent of the strains harboring the ESBL plasmids. When comparing the PT of the two rounds of conjugation, we observed that the frequency of PT of the second round of conjugation was higher than that of the first round of conjugation. This might be because the strains used as donors (E. coli J53-2 harboring plasmids) and recipients (E. coli J53-1) in the second round have high homology being resistant derivatives of the same K12 strain.

Previously, it had been shown that induction of the SOS-response by antibiotics enhanced transfer of plasmids and ICEs in Vibrio cholerae, Aeromonas hydrophila, and S. aureus (Barr et al., 1986; Beaber et al., 2004; Cantas et al., 2012). However, we have shown that CTX exposure increases transfer of a blaCTX−M−1 encoding IncI1 plasmid in E. coli in a SOS-independent manner (Moller et al., 2017). We therefore also investigated the involvement of the SOS response, by comparing the expression of two genes associated with the SOS response, recA and sfaA, in the transconjugants with and without antibiotic exposure. We found that none of the typical SOS-response genes were induced by growth in the presence of CTX and AMP, but they were induced when the strains were exposed to CIP. This result supports previous findings showing that quinolones/CIP induce the SOS response (Piddock and Wise, 1987; Beaber et al., 2004). However, a recent study showed that even though CIP increased the expression of genes associated with the SOS response, there was no correlation with conjugation efficiency, and they concluded that sub-MIC CIP may promote conjugational PT by up-regulating the expression of conjugation associated genes via an SOS-independent mechanism (Shun-Mei et al., 2018). Since the strains are CTX and AMP resistant, but CIP sensitive, it could be suggested that only when exposed to CIP, the strains induce the SOS response to modulate their susceptibility. It is therefore currently not possible to conclude whether the CIP-induced PT and SOS-response occurs through dependent or independent regulatory pathways.

Previously published results (Moller et al., 2017) revealed that the resistance gene (blaCTX−M−1) was required for CTX-induced tra-expression and PT. The mechanism behind the up-regulation remains elusive. A proteomic study of MG1655/pTF2 exposed to CTX only showed up-regulation of plasmid-encoded transfer-related proteins, the CTX-M-1 β-lactamase, and a hypothetical protein in response to the antibiotic. However, a proteomic approach may be to insensitive to determine the pathways involved, since various regulatory genes (e.g., two-component systems, regulatory RNAs, and regulatory proteins) may not show significant up-regulation, but yet be changed in expression. Further investigations on the regulatory mechanism(s) are therefore relevant.

### CONCLUSION

In conclusion, our findings reveal that exposure to different antibiotics can increase conjugative transfer of different naturally occurring β-lactam resistance plasmids in a strain-independent manner. Thus, antibiotic-induced conjugation transfer of ESBL plasmids appears to be a more common phenomenon in E. coli, at least for plasmids carrying the CTX-M bla-types. Since increased conjugation frequency will be an adverse effect of drug treatment of humans and animals, there is an urgent need to investigate the mechanisms underlying antibiotic-mediated increased conjugative PT, to be able to counteract the spread of resistance. Our findings have implications for evaluating
the risks of antibiotic use, and for developing new antibiotic treatment protocols.

**DATA AVAILABILITY**

The raw data supporting the conclusions of this manuscript will be made available by the authors, without undue reservation, to any qualified researcher.

**AUTHOR CONTRIBUTIONS**

GL, LT, VB, and JO designed the study. GL and KB carried out the experiments. GL, VB, and LT analyzed the results. GL and LT wrote the manuscript. All authors commented on and approved the final version of the manuscript.

**REFERENCES**

Al-Bayssari, C., Dabbousi, F., Hamze, M., and Rolain, J. M. (2015). Detection of expanded-spectrum beta-lactamases in Gram-negative bacteria in the 21st century. *Expert Rev Anti Infect Ther* 13, 1139–1158. doi: 10.1586/14787210.2015.1066247

Aminov, R. I. (2011). Horizontal gene exchange in environmental microbiota. *Front Microbiol* 2:158. doi: 10.3389/fmicb.2011.00158

Appelbaum, P. C., Trichardt, P. F., Krizsanovich, K., Coetzee, J. N., and Hugon. Barlow, M. (2009). What antimicrobial resistance has taught us about horizontal gene transfer. *Methods Mol Biol* 532, 397–411. doi: 10.1007/978-1-60327-853-9_23

Barr, V., Barr, K., Millar, M. R., and Lacey, R. W. (1986). Beta-lactam antibiotics increase the frequency of plasmid transfer in Staphylococcus aureus. *J Antimicrob Chemother* 17, 409–413. doi: 10.1093/jac/17.4.409

Beaber, J. W., Hochhut, B., and Waldor, M. K. (2004). SOS response promotes conjugative transposon Tn916 in the guts of gnotobiotic rats. *Appl Environ Microbiol* 70, 758–764. doi: 10.1128/ae.70.2.758-764.2004

Bankievel, A., Nuri, S., Antipov, D., Gurevich, A. A., Dvorkin, M., Kulikov, A. S., et al. (2012). SPADES: a new genome assembly algorithm and its applications to single-cell sequencing. *J Comput Biol* 19, 455–477. doi: 10.1089/cmb.2012.0021

Barlow, M. (2009). What antimicrobial resistance has taught us about horizontal gene transfer. *Methods Mol Biol* 532, 397–411. doi: 10.1007/978-1-60327-853-9_23

Bolger, A. M., Lohse, M., and Usadel, B. (2014). Trimmomatic: a flexible trimmer for Illumina sequence data. *Bioinformatics* 30, 2114–2120. doi: 10.1093/bioinformatics/btu170

Cantas, L., Midtflyng, P. J., and Sorum, H. (2012). Impact of antibiotic treatments on the expression of the R plasmid tra genes and on the host innate immune activity during pRAS1 bearing Aeromonas hydrophila infection in eel (Danio rerio). *BMC Microbiol* 12:37. doi: 10.1186/1471-2180-12-37

Carattoli, A., Bertini, A., Villa, L., Falbo, V., Hopkins, K. L., and Threlfall, E. J. (2005). Identification of plasmids by PCR-based replicon typing. *J Microbiol Methods* 63, 219–228. doi: 10.1016/j.mimet.2005.03.018

Carloni, E., Andreoni, F., Omiccioli, E., Villa, L., Magnani, M., and Carattoli, A. (2017). Comparative analysis of the standard PCR-Based Replicon Typing (PBRT) with the commercial PBRT-KIT. *Plasmid* 90, 10–14. doi: 10.1016/j.plasmid.2017.01.005

de la Cruz, F., Frost, L. S., Meyer, R. J., and Zechner, E. L. (2010). Conjugative DNA metabolism in Gram-negative bacteria. *FEMS Microbiol Rev* 34, 18–40. doi: 10.1111/j.1574-6976.2009.00195.x

Feld, L., Schjoining, S., Hammer, K., Licht, T. R., Danielsen, M., Krogfelt, K., et al. (2008). Selective pressure affects transfer and establishment of a Lactobacillus plantarum resistance plasmid in the gastrointestinal environment. *J Antimicrob Chemother* 61, 485–492. doi: 10.1093/jac/dkn033

Halary, S., Leigh, J. W., Cheaib, B., Lopez, P., and Baptiste, E. (2010). Network analyses structure genetic diversity in independent genetic worlds. *Proc Natl Acad Sci U S A* 107, 127–132. doi: 10.1073/pnas.0908978107

Hawkey, P. M., and Jones, A. M. (2009). The changing epidemiology of resistance. *J Antimicrob Chemother* 64(Suppl. 1), i3–i10. doi: 10.1093/jac/dkp256

Huang, H., Liao, J., Zheng, X., Chen, Y., and Ren, H. (2019). Low-level free nitrous acid efficiently inhibits the conjugative transfer of antibiotic resistance by altering intracellular ions and disabling transfer apparatus. *Water Res* 158, 383–391. doi: 10.1016/j.watres.2019.04.046

Johnson, C. M., and Grossman, A. D. (2015). Integrative and Conjugative Elements (ICEs): What They Do and How They Work. *Annu Rev Genet* 49, 577–601. doi: 10.1146/annurev-genet-112414-055018

Kado, C. I., and Liu, S. T. (1981). Rapid procedure for detection and isolation of large and small plasmids. *J Bacteriol* 145, 1365–1373.

Karzmarczyk, M., Abbott, Y., Walsh, C., Leonard, N., and Fanning, S. (2011). Characterization of multidrug-resistant *Escherichia coli* isolates from animals presenting at a university veterinary hospital. *Appl Environ Microbiol* 77, 7104–7112. doi: 10.1128/AEM.00599-11

Kjeldsen, T. S., Overgaard, M., Nielsen, S. S., Bortolaia, V., Jelsbak, L., Sommer, M., et al. (2015). CTX-M-1 beta-lactamase expression in *Escherichia coli* is dependent on cefotaxime concentration, growth phase and gene location. *J Antimicrob Chemother* 70, 62–70. doi: 10.1093/jac/dku332

Lento, C., Ferraro, M., Wilson, D., and Audette, G. F. (2016). HDX-MS and deletion analysis of the type 4 secretion system protein TraF from the *Escherichia coli* F plasmid. *FEBS Lett* 590, 376–386. doi: 10.1002/1873-3468.12066

Lopatkin, A. J., Huang, S., Smith, R. P., Srimani, J. K., Sysoeva, T. A., Bewick, S., et al. (2016). Antibiotics as a selective driver for conjugation dynamics. *Nat Microbiol* 1, 16044. doi: 10.1038/nmicrobiol.2016.44

Lu, Y., Zeng, J., Wang, L., Lan, K. E. S., Wang, L., Xiao, Q., et al. (2017). Antibiotics Promote *Escherichia coli*-Pseudomonas aeruginosa Conjugation through Inhibiting Quorum Sensing. *Antimicrob Agents Chemother* 61, e1284–e1217. doi: 10.1128/AAC.01284-17

Macrina, F. L., Kopecko, D. J., Jones, K. R., Ayers, D. J., and Mccowen, S. M. (1978). A multiple plasmid-containing *Escherichia coli* strain: convenient source of size reference plasmid molecules. *Plasmid* 1, 417–420. doi: 10.1016/0147-619x(78)90056-2

**FUNDING**

GL was supported by a Ph.D. fellowship from the China Scholarship Council.

**ACKNOWLEDGMENTS**

Tony P. Bonnelycke and Dan F. Ryttov are thanked for valuable technical assistance.

**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.02119/full#supplementary-material
Malloy, A. M., and Campos, J. M. (2011). Extended-spectrum beta-lactamases: a brief clinical update. *Pediatr Infect Dis J* 30, 1092–1093. doi: 10.1097/inf.0b013e31823ce9fd

Mazel, D., and Davies, J. (1999). Antibiotic resistance in microbes. *Cell Mol Life Sci* 56, 742–754.

McElho, J. A., Holland, M. M., Makova, K. D., Su, M. S., Paul, I. M., Baker, C. H., et al. (2014). Development and assessment of an optimized next-generation DNA sequencing approach for the mtgenome using the Illumina MiSeq. *Forensic Sci Int Genet* 13, 20–29. doi: 10.1016/j.fsigen.2014.05.007

Meunier, D., Jouy, E., Lazizzera, C., Kobisch, M., and Madec, J. Y. (2006). CTX-M-1- and CTX-M-15-type beta-lactamases in clinical *Escherichia coli* isolates recovered from food-producing animals in France. *Int J Antimicrob Agents* 28, 402–407. doi: 10.1016/j.ijantimicag.2006.08.016

Moller, T. S. B., Liu, G., Boysen, A., Thomsen, L. E., Luthje, F. L., Mortensen, S., et al. (2017). Treatment with Cefotaxime Affects Expression of Conjugation Associated Proteins and Conjugation Transfer Frequency of an IncI1 Plasmid in *Escherichia coli*. *Front Microbiol* 8:2365. doi: 10.3389/fmictb.2017.02365

Norman, A., Hansen, L. H., and Sorensen, S. J. (2009). Conjugative plasmids: vessels of the communal gene pool. *Philos Trans R Soc Lond B Biol Sci* 364, 2275–2289. doi: 10.1098/rstb.2009.0037

Orlek, A., Phan, H., Sheppard, A. E., Doumith, M., Ellington, M., Petò, T., et al. (2017). A curated dataset of complete Enterobacteriaceae plasmids compiled from the NCB! nucleotide database. *Data Brief* 12, 423–426. doi: 10.1016/j.db.2017.04.024

Overdevest, I., Willemsen, I., Rijnsburger, M., Eustace, A., Xu, L., Hawkey, P., et al. (2011). Extended-spectrum beta-lactamase genes of *Escherichia coli* in chicken meat and humans. *The Netherlands. Emerg Infect Dis* 17, 1216–1222. doi: 10.3201/eid1707.110209

Pfaff, M. W. (2001). A new mathematical model for relative quantification in real-time RT-PCR. *Nucleic Acids Res* 29, e45.

Piddock, L. J., and Wise, R. J. F. M. (1987). Induction of the SOS response in *Escherichia coli* by 4-quinolone antimicrobial agents. *FEMS Microbiology Letters* 41, 289–294. doi: 10.1016/0168-6445(87)90255-2

Rodriguez, L., Barownick, W., Helmut, R., Mendoza, M. C., Rodicio, M. R., Schroeter, A., et al. (2009). Extended-spectrum [beta]-lactamases and AmpC [beta]-lactamases in cefotiofur-resistant *Salmonella enterica* isolates from food and livestock obtained in Germany during 2003-07. *J Antimicrob Chemother* 64, 301–309. doi: 10.1093/jac/dkp195

Shun-Mei, E., Zeng, J. M., Yuan, H., Lu, Y., Cai, R. X., and Chen, C. (2018). Sub-inhibitory concentrations of fluoroquinolones increase conjugation frequency. *Microb Pathog* 114, 57–62. doi: 10.1016/j.micpath.2017.11.036

Sorensen, S. J., Bailey, M., Hansen, L. H., Kroer, N., and Wuertz, S. (2005). Studying plasmid horizontal transfer in situ: a critical review. *Nat Rev Microbiol* 3, 700–710. doi: 10.1038/nrmicro1232

 Stevens, A. M., Shoemaker, N. B., Li, L. Y., and Salyers, A. A. (1993). Tetracycline regulation of genes on *Bacteroides* conjugative transposons. *J Bacteriol* 175, 6134–6141. doi: 10.1128/JB.175.19.6134-6141.1993

Taylor, D. E., and Levine, J. G. (1980). Studies of temperature-sensitive transfer and maintenance of H incompatibility group plasmids. *J Gen Microbiol* 116, 475–484. doi: 10.1009/00221287-116-2-475

Thomsen, M. C., Ahrenfeldt, J., Cisneros, J. L., Jurtz, V., Larsen, M. V., Hasman, H., et al. (2016). A Bacterial Analysis Platform: An Integrated System for Analysing Bacterial Whole Genome Sequencing Data for Clinical Diagnostics and Surveillance. *PLoS One* 11:e0157718. doi: 10.1371/journal.pone.0157718

von Wintersdorff, C. J., Penders, J., Van Nierkerk, J. M., Mills, N. D., Majumder, S., Van Alphen, L. B., et al. (2016). Dissemination of Antimicrobial Resistance in Microbial Ecosystems through Horizontal Gene Transfer. *Front Microbiol* 7:173. doi: 10.3389/fmicb.2016.00173

Waters, V. L., Strack, B., Pansegrau, W., Lanka, E., and Guiney, D. G. (1992). Mutational analysis of essential IncP alpha plasmid transfer genes traF and traG and involvement of traF in phage sensitivity. *J Bacteriol* 174, 6666–6673. doi: 10.1128/jb.174.20.6666-6673.1992

Wayne, P. A., Clinical, and Laboratory Standards Institute. (2015). *CLSI Performance Standards for Antimicrobial Susceptibility Testing; Twenty-Fifth Informational Supplement*. CLSI Document M100-S25. Wayne, PA: Clinical and Laboratory Standards Institute.

Whittle, G., Shoemaker, N. B., and Salyers, A. A. (2002). Characterization of genes involved in modulation of conjugal transfer of the *Bacteroides* conjugative transposon CTnDOT. *J Bacteriol* 184, 3839–3847. doi: 10.1128/jb.184.14.3839-3847.2002

Zatyka, M., and Thomas, C. M. (1998). Control of genes for conjugative transfer of plasmids and other mobile elements. *FEMS Microbiol Rev* 21, 291–319. doi: 10.1016/0168-6445(97)00061-2

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Liu, Bogaj, Bortolaia, Olsen and Thomsen. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.