Emergence of tigecycline resistance in *Escherichia coli* co-producing MCR-1 and NDM-5 during tigecycline salvage treatment

**Objective:** Here, we report a case of severe infection caused by *Escherichia coli* that harbored *mcr-1*, *bla*<sub>NDM-5</sub>, and acquired resistance to tigecycline during tigecycline salvage therapy.

**Methods:** Antimicrobial susceptibility testing, Southern blot hybridization, and complete genome sequence of the strains were carried out. The genetic characteristics of the *mcr-1* and *bla*<sub>NDM-5</sub> plasmids were analyzed. The whole genome sequencing of *mcr-1*-containing plasmid was completed. Finally, putative single nucleotide polymorphisms and deletion mutations in the tigecycline-resistant strain were predicted.

**Results:** Three *E. coli* isolates were obtained from ascites, pleural effusion, and stool of a patient; they were resistant to almost all the tested antibiotics. The first two strains separated from ascites (E-FQ) and hydrothorax (E-XS) were susceptible to amikacin and tigecycline; however, the third strain from stool (E-DB) was resistant to tigecycline after nearly 3 weeks’ treatment with tigecycline. All three isolates possessed both *mcr-1* and *bla*<sub>NDM-5</sub>. The *bla*<sub>NDM-5</sub> gene was found on the IncX3 plasmid, whereas the *mcr-1*, *fosA3* and *bla*<sub>CTX-M-14</sub> were located on the IncH12 plasmid. Mutations in *acrB* and *lon* were the reason for the resistance to tigecycline.

**Conclusion:** This is the first report of a colistin-, carbapenem-, and tigecycline-resistant *E. coli* in China. Tigecycline resistance acquired during tigecycline therapy is of great concern for us because tigecycline is a drug of last resort to treat carbapenem-resistant Gram-negative bacterial infections. Furthermore, the transmission of such extensively drug-resistant isolates may pose a great threat to public health.

**Keywords:** *mcr-1*, *bla*<sub>NDM-5</sub>, tigecycline-resistance, *acrB*, *lon*

**Introduction**

The emergence of multidrug-resistant Gram-negative bacteria, especially those producing carbapenemases, continues to threaten public health because of the limited availability of antimicrobials for treatment. Colistin and tigecycline are drugs of last resorts to treat these severe infections. The rapid transmission of the colistin-resistance gene *mcr-1* located on mobile elements has caused great concern since its initial discovery. Co-carriage of *mcr-1* and *bla*<sub>NDM-5</sub> has been reported in isolates from animals, food, and even humans since 2016.1–5 Tigecycline is effective against most carbapenemase-producing bacteria and was recently approved for clinical use in China; nevertheless, acquisition of resistance to tigecycline during therapy has been frequently reported.6–10 The mechanism of tigecycline resistance has not yet been clearly elucidated. It has been reported that resistance-nodulation-division (RND)-type transporters, primarily the AcrAB efflux pump, play an important role.11 Here, we report a case of intra-abdominal...
infection and pyothorax. Three extensively drug-resistant (XDR) *Escherichia coli* isolates with resistance to colistin and carbapenems were isolated from the patient. The first two strains, from ascites (E-FQ) and hydrothorax (E-XS), were susceptible to amikacin and tigecycline; however, the third strain from stool (E-DB) acquired resistance to tigecycline during tigecycline therapy. Fortunately, following combined antibiotic susceptibility tests in vitro, the infection was finally controlled.

To the best of our knowledge, this is the first report of a colistin-, carbapenem-, and tigecycline-resistant *E. coli* in China. The aim of our study was to clarify the resistance mechanisms of these XDR pathogens.

**Materials and methods**

**Patient and isolate data**

Three carbapenem-resistant strains of *E. coli* were isolated from a 46-year-old male patient. The patient had a history of hepatocellular carcinoma and was admitted to our hospital for intra-abdominal infection and pyothorax after liver resection in March 2017. To identify the bacteria and guide the antibiotic treatment, ascites and hydrothorax were cultured, and the patient underwent a stool screening. The first two strains cultured from ascites and hydrothorax were separated on March 8, 2017. The third, from stool, was isolated on March 28, 2017. All three isolates were identified by the VITEK 2 system (Sysmex-bioMérieux, Marcy l'Etoile, France).

**PCR and Sanger sequencing**

PCR and sequencing were used to identify the *mcr-1* and *bla*<sup>NDM-5</sup> genes in all three isolates. Mutations in *acrB* and *lon* were confirmed by PCR and Sanger sequencing (primers in Table S1).

**Antimicrobial susceptibility test**

Antibiotic susceptibilities, including imipenem, meropenem, amikacin, levofloxacin, cefotaxime, aztreonam, amoxicillin–clavulanic acid, and fosfomycin, were determined by the broth microdilution method according to the guidelines provided by the Clinical and Laboratory Standards Institute. Minimum inhibitory concentrations (MICs) of tigecycline and colistin were interpreted by EUCAST guidelines (http://www.eucast.org/). MIC by Etest followed the manufacturer’s instructions for some drugs, including cefoperazone/sulbactam, ciprofloxacin, aztreonam, and trimethoprim–sulfamethoxazole. *E. coli* ATCC 25922 was used as a quality control strain for the antibiotic susceptibility test. The interactions of amoxicillin–clavulanic acid and aztreonam were assessed using the checkerboard method with slight modifications. The combinations in the 96-well plates were performed as follows: aztreonam was diluted by twofold dilutions along the x-axis of the plates (from 1/64 MIC to 2 MIC), while amoxicillin–clavulanic acid was diluted by twofold dilutions along the y-axis (from 1/32 MIC to 2 MIC). Subsequently, each well was inoculated with the tested bacterial suspension (5×10<sup>4</sup> CFU/mL). The plates were then incubated at 37°C overnight. The interaction between aztreonam and amoxicillin–clavulanic acid was determined by quantifying the fractional inhibitory concentration index (FICI) using the following formula: FIC of aztreonam = MIC of aztreonam in combination/MIC of aztreonam alone; FIC of amoxicillin–clavulanic acid = MIC of amoxicillin–clavulanic acid in combination/MIC of amoxicillin–clavulanic acid alone. FICI = FIC<sub>a</sub> + FIC<sub>b</sub>. “Synergy” was defined when FICI ≤ 0.5; 0.5 < FICI ≤ 0.75 means “partial synergy”; 0.76 < FICI ≤ 1 denotes “additive”; 1 < FICI ≤ 4 denotes “indifferent”; while “antagonistic” in cases in which the FICI > 4.

**Pulsed-field gel electrophoresis (PFGE) and Southern blotting analysis**

Genetic relationships among the isolates were evaluated by PFGE. Genomic DNA was digested with restriction enzyme XbaI (TaKaRa, Dalian, China) and was electrophoresed on a CHEF-mapper XA PFGE system (Bio-Rad, Berkeley, CA, USA) at 14°C and 6 V/cm and with alternating pulses at a 120° angle in a 5–35 seconds pulse time gradient for 22 hours. To ascertain the plasmid locations, genomic DNA was digested with S1-nuclease (TaKaRa, Kusatsu, Japan) and was electrophoresed for 18 hours at 14°C with run conditions of 6 V/cm and pulse times from 2.16 to 63.8 seconds. The DNA fragments were transferred to a positively charged nylon membrane (Millipore, Billerica, MA, USA) and then were hybridized with a digoxigenin-labeled *bla*<sup>NDM-5</sup> and *mcr-1*-specific probe. An NBT/BCIP color detection kit (Roche, Mannheim, Germany) was then used to detect the target gene. The *Salmonella enterica* serotype Braenderup H9812 was used as the size marker.

**Genomic DNA extraction and analysis**

Genomic DNA from strain E-FQ, E-XS, E-DB was sequenced with an Illumina HiSeq 2000 (Illumina Inc., San Diego, CA, USA) following a paired-end 2×100-bp protocol. The total genomic DNA of E-DB was also extracted and sequenced via the single molecule real-time technique using a PacBio RS II platform and was assembled de novo using
Canu v1.5. Gene prediction and annotation were performed with in silico online tools (http://rast.nmpdr.org/rast.cgi). Sequence comparison was performed using BLAST (http://blast.ncbi.nlm.nih.gov). Determination of plasmid replications, resistance genes, and multi-locus sequence typing (MLST) was performed using online CGE tools (http://www.genomicepidemiology.org). A comparison of genetic environment of mcr-1- and blaNDM-5-containing plasmids and their related plasmids was performed with EasyFig 2.2.2.14 The reads of E-DB were mapped against the reference genome of E-FQ and E-XS with the CLC Genomics Workbench 9 software (Qiagen, Valencia, CA, USA). The putative single nucleotide polymorphisms and deletion mutations were predicted.

### Nucleotide sequence accession numbers

The next-generation sequencings of E-FQ, E-XS, and E-DB were subjected to National Center for Biotechnology Information under accession numbers PVQI00000000, PVQH00000000, and PVQG00000000. The complete sequences of pE-DB-mcr were submitted to GenBank under accession number MH128771.

### Results

#### Patient information

A 46-year-old male patient presented to our hospital (a tertiary-care hospital in Hangzhou) with recurrent fever, pyothorax, and intra-abdominal infection in March 2017. The patient had a history of chronic hepatitis B and liver cancer. He had undergone a liver resection for the hepatocellular cancer 9 months prior. The patient had recurrent fever and abdominal pain after the surgery. A colonic fistula was found on imaging studies and during enteroscopy. The patient was admitted to our hospital (a tertiary-care hospital in Hangzhou) with recurrent fever, hydrothorax, and intra-abdominal infection in March 2017. The patient had a history of chronic hepatitis B and liver cancer. He had undergone a liver resection for the hepatocellular cancer 9 months prior. The patient had recurrent fever and abdominal pain after the surgery. A colonic fistula was found on imaging studies and during enteroscopy. The patient was treated with imipenem (0.5 g q8h for 8 days) and tigecycline (50 mg q12h for 7 days) prior to this admission.

The results of laboratory examinations were as follows: white blood cell count, 15,200/mm³; neutrophil percentage, 91%; platelet count, 79,000/mm³; alpha-fetoprotein, 1,238.3 ng/mL; C-reactive protein, 62.6 mg/L, and procalsitonin, 0.5 μg/L. A series of pathogens cultures from ascites, plural effusion, and stool grew XDR *E. coli* that were resistant to almost all tested antibiotics. The first two strains separated from ascites (E-FQ) and hydrothorax (E-XS) after admission were susceptible to amikacin and tigecycline; however, the third strain from stool (E-DB) was susceptible to amikacin but was resistant to tigecycline after nearly 3 weeks’ treatment with tigecycline (50 mg q12 h for 17 days and 100 mg q12 h for 3 days) in our hospital.

The patient underwent ileum colostomy and abdominal cavity drainage, while amoxicillin–clavulanic acid combined with aztreonam was used for 1 week according to the in vitro combined antibiotic susceptibility test (FICI 0.125). The bacterium was finally eliminated. However, the patient died in the fourth month of hospitalization in our hospital because of cancer recurrence.

### Isolate characteristics

The antimicrobial susceptibility testing results showed that three *E. coli* strains were resistant to almost all tested antibiotics (Table 1). The first two strains E-FQ and E-XS were susceptible to amikacin (MIC = 8 mg/L) and tigecycline (MIC = 1 mg/L); however, the third strain E-DB was resistant to tigecycline (MIC = 8 mg/L). PCR and sequencing identified that all three isolates contained genes mcr-1 and blaNDM-5. These isolates expressed indistinguishable band patterns by XbaI PFGE analysis (Figure S1). In addition to mcr-1 and blaNDM-5, the three strains were found to harbor resistance genes for aminoglycosides (*aph*4-*Ia, aadA1, streB, strA, aacA4, aac(3)-IId and aac(3)-IV a), beta-lactam (*bla*), fluoroquinolone (*aac(6’)*Ib-cr), fosfomycin (*fosA*), phenicol (*floR*), sulfonamide (*sul1 and sul2*), trimethoprim (*dfrA1*), and tetracycline (*tetA*), that were consistent with an extremely drug-resistant profile. Plasmid incompatibility

### Table 1 Sources and MICs for the three Escherichia coli isolates and E. coli ATCC 25922

<table>
<thead>
<tr>
<th>Isolates</th>
<th>Date of isolation</th>
<th>Source</th>
<th>MLST</th>
<th>MIC (mg/L)</th>
<th>IMP</th>
<th>MEM</th>
<th>CST</th>
<th>TGC</th>
<th>AK</th>
<th>LVX</th>
<th>CTX</th>
<th>FEP</th>
<th>FOS</th>
<th>AT</th>
<th>CI*</th>
<th>CPS</th>
<th>TS*</th>
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<tbody>
<tr>
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<td>March 8, 2017</td>
<td>Human ascites</td>
<td>2736</td>
<td>16</td>
<td>32</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>64</td>
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<td>1.024</td>
<td>12</td>
<td>&gt;32</td>
<td>&gt;256</td>
<td>&gt;32</td>
<td></td>
</tr>
<tr>
<td>E-XS</td>
<td>March 8, 2017</td>
<td>Human hydrothorax</td>
<td>2736</td>
<td>16</td>
<td>32</td>
<td>8</td>
<td>1</td>
<td>8</td>
<td>64</td>
<td>&gt;256</td>
<td>256</td>
<td>1.024</td>
<td>12</td>
<td>&gt;32</td>
<td>&gt;256</td>
<td>&gt;32</td>
<td></td>
</tr>
<tr>
<td>E-DB</td>
<td>March 28, 2017</td>
<td>Human feces Laboratory</td>
<td>2736</td>
<td>32</td>
<td>32</td>
<td>4</td>
<td>8</td>
<td>8</td>
<td>128</td>
<td>&gt;256</td>
<td>&gt;256</td>
<td>1.024</td>
<td>16</td>
<td>&gt;32</td>
<td>&gt;256</td>
<td>&gt;32</td>
<td></td>
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<tr>
<td>ATCC 25922</td>
<td></td>
<td></td>
<td>0.125</td>
<td>&lt;0.125</td>
<td>0.25</td>
<td>0.125</td>
<td>2</td>
<td>&lt;0.125</td>
<td>0.125</td>
<td>&lt;0.125</td>
<td>1</td>
<td>1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Notes:** MICs were interpreted by Etest by following the manufacturer’s instructions for CPS, CI, TS, and AT.

**Abbreviations:** MLST, multi-locus sequence typing; MICs, minimum inhibitory concentration(s); IMP, imipenem; MEM, meropenem; CST, colistin; TGC, tigecycline, AK, amikacin, LVX, levofloxacin; CTX, cefotaxime, FEP, cefepime, FOS, fosfomycin; AT, aztreonam; CI, ciprofloxacin, CPS, cefoperazone/sulbactam; TS, trimethoprim–sulfamethoxazole.
typing detected the presence of the plasmid replicons IncA/C2, IncFIA, IncHI2A, IncHI2, IncX3, and IncFIB. MLST demonstrated that the isolates belonged to sequence type 2736. The reads of strain E-DB were mapped against the reference genome of E-FQ and E-XS. The difference between E-DB and E-FQ was less than five single nucleotide variants (Table 2), meanwhile, the difference between E-DB and S-XS was also less than five.

Genetic context of plasmids
S1 nuclease PFGE and Southern blot showed the presence of mcr-1 in an ~244–330 kb plasmid and bla\textsubscript{NDM-5} in an ~33–55 kb plasmid (Figure S1), belonging to IncHI2 and IncX3 by analysis of whole genome sequencing. mcr-1-containing plasmid pE-DB-mcr was 230,365 bp in size with an average G+C content of 51%. In the multidrug resistance (MDR) region, it harbored resistance genes for aminoglycosides (aph(4)-Ia, aac(6)-Ia, and aac(3)-IvA), fosfomycin (fosA), phenicol, sulfonamide (sul2) and beta-lactam (bla\textsubscript{CTX-M-14}). The horizontal diffusion of these resistance genes is particularly worrisome because the regimen of colistin plus fosfomycin remains a therapeutic alternative in cases of multidrug-resistant bacteria. There is increasing evidence that IS\textsubscript{Apl1}-bearing plasmids deposited in GenBank suggest that IncHI2 is one of the most dominant replicon types of plasmid carrying mcr-1 plasmids (45/219, 20.5%); IncHI2 plasmids had the largest size, with sequence lengths up to 267,486 bp that were always integrated alongside a large MDR gene cassette. In our study, we found that the mcr-1-containing plasmid also harbored the fosfomycin resistance gene fosA3 and beta-lactam bla\textsubscript{CTX-M-14}. The horizontal diffusion of these resistance genes is particularly worrisome because the regimen of colistin plus fosfomycin remains a therapeutic alternative in cases of multidrug-resistant bacteria. There is increasing evidence that IS\textsubscript{Apl1} plays a pivotal role in the mobilization of mcr-1. Tn6330, with the structure IS\textsubscript{Apl1}-mcr-1-orf-IS\textsubscript{Apl1}, was proposed as a key element mediating translocation of mcr-1.

Tigecycline resistance
Antimicrobial susceptibility showed that E-FQ and E-XS were sensitive to tigecycline (MIC 1 mg/L), while strain E-DB was resistant to tigecycline (MIC 8 mg/L). The putative single nucleotide and deletion mutations in E-DB were predicted. acrB and lon were the only two genes found in both mappings (Table 2). Mutations in acrB and lon were confirmed by PCR and Sanger sequencing. As to acrB, it is a single nucleotide mutation that causes a change of arg to cys (R620C). As to lon, it is a single nucleotide polymorphism mutation that resulted in the change of pro to leu (P403L) (Table 2).

Discussion
To our knowledge, this is the first report of an isolate harboring mcr-1 and bla\textsubscript{NDM-5} acquiring resistance to tigecycline during therapy. Since its discovery in Southern China, mcr-1 has already been discovered in >40 countries/regions covering five of seven continents and where it plays an important role in polymyxin resistance. Searches for mcr-1-bearing plasmids deposited in GenBank suggest that IncHI2 is one of the most abundant replicon types of plasmids carrying mcr-1 plasmids (45/219, 20.5%); IncHI2 plasmids had the largest size, with sequence lengths up to 267,486 bp that were always integrated alongside a large MDR gene cassette.

Table 2 The putative single nucleotide variants and deletion mutations in Escherichia coli E-DB compared with E-FQ and E-XS

<table>
<thead>
<tr>
<th>Reference position</th>
<th>Gene and product</th>
<th>Nucleotide sequence</th>
<th>Amino acid sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>E. coli E-FQ</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>contig_44 3,490</td>
<td>acrB, multidrug efflux RND transporter permease subunit</td>
<td>C1858T</td>
<td>R620C</td>
</tr>
<tr>
<td>contig_44 25,952</td>
<td>lon, ATP-dependent protease Lα</td>
<td>G1148A</td>
<td>P403L</td>
</tr>
<tr>
<td>contig_128 9,752</td>
<td>ISL3, transposase</td>
<td>G191T</td>
<td>A64S</td>
</tr>
<tr>
<td>contig_12 1,269</td>
<td>radC, DNA repair protein</td>
<td>G174A</td>
<td>V247I</td>
</tr>
<tr>
<td>contig_147 2,477</td>
<td>Not an ORF</td>
<td>T2477C</td>
<td>None</td>
</tr>
<tr>
<td>E. coli E-XS</td>
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<td></td>
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</tr>
<tr>
<td>contig_77 52,399</td>
<td>acrB, multidrug efflux RND transporter permease subunit</td>
<td>C1858T</td>
<td>R620C</td>
</tr>
<tr>
<td>contig_77 29,937</td>
<td>lon, ATP-dependent protease Lα</td>
<td>G1148A</td>
<td>P403L</td>
</tr>
<tr>
<td>contig_43 206</td>
<td>orfA, IS1 transposase</td>
<td>T204C</td>
<td>R688</td>
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<td>contig_43 209</td>
<td>orfA, IS1 transposase</td>
<td>C207T</td>
<td>H69H</td>
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<tr>
<td>contig_53 80</td>
<td>Not an ORF</td>
<td>A80G</td>
<td>None</td>
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</tbody>
</table>

Abbreviation: ORF, open reading frame.
genes, respectively. Brown arrows indicate genes for hypothetical proteins as well as IS\_apl\_Kpn26. Light gray shades indicate shared regions with a high degree of homology.

The genetic environment of mcr-1 and blanDM-5. The genetic environment of mcr-1 bearing IncHII plasmid and the genetic information (green and purple circles). (C) The genetic environment of IncHI2. The plasmid backbone was divided by an IS\_apl\_Kpn26 into two parts, with a copy of IS\_apl\_Kpn26 and one part of IS\_apl\_ that was highly similar to plasmid pls1. (C) The genetic environment of IncHI2. The IS\_apl\_Kpn26 was inserted by an IS\_apl\_Kpn26 in two parts, and we infer that this insertion may increase the stability of Tn6330 in the backbone of IncHI2.

Tigecycline belongs to the group called glycylcyclines that binds to 16S rRNA and prevents mRNA decoding. Compared with earlier class tetracyclines, the bulky side chain attached to the C-9 position of ring D is the key to escape common tetracycline resistance mechanisms, for example, tetracycline efflux pumps and ribosomal protection. Tigecycline is also not affected by most of the common mechanisms that affect other classes of antimicrobial agents, including target site modifications, enzymatic degradation of the drug molecule, and DNA gyrase mutations. However, the reports of clinical tigecycline resistance were published soon after its use in medical practice. Tigecycline resistance occurring in carbapenem-resistant Enterobacteriaceae (CRE) during therapy has been reported in many cases. It had been reported that Tet proteins can mutate to acquire high-level tigecycline resistance, but in most cases, tigecycline resistance is due to the constitutive expression of RND efflux pumps; for example, AcrAB in E. coli. In E. coli, the tripartite efflux system AcrAB-ToIC is the pump in charge of the efflux of dyes, bile salts, detergents, and a large group of antibiotics, including the tetracyclines. AcrB is an RND pump situated in the inner membrane that functions in a complex with two other proteins, an outer membrane channel TolC and a periplasmic adaptor protein AcrA. AcrB is the major site for substrate recognition and energy transduction of the entire tripartite system. In one study, the author found a trend toward higher acrB expression as tigecycline MIC increases. Several global transcriptional regulators of the araC family, ramA, marA, soxS, and rra, may participate in tigecycline resistance via AcrAB efflux pump activation, of which ramA, marA, and soxS are regulated by the lon. Nicoloff et al found that about half of the selected spontaneous tetracycline- or chloramphenicol-resistant E. coli mutants carried additional lon mutations. lon mutation causes low MDR by stabilizing MarA and SoxS, two transcriptional activators, whose induction increases resistance to many antimicrobial drugs. The first-step lon mutation greatly increases the rate of development of higher resistance via acquisition of new sets of mutations. In this case, with tigecycline treatment, mutations in acrB (R620C) and lon (P403L) in strain E-DB caused activation of the AcrAB-ToIC efflux pump, leading to the high MIC of 8 mg/L for tigecycline.
CRE are often associated with severe health care-associated infections with high mortality ranging from 32.1% to 71%. NDM-type beta-lactamase are the second most common carbapenemases found among CRE in China. These carbapenemases cannot be inhibited by beta-lactamase inhibitors but remain susceptible to aztreonam, although most metallo-beta-lactamase-producing CRE isolates also harbor extended-spectrum beta-lactamases that confer resistance to aztreonam. Therefore, many NDM producers remain susceptible only to tigecycline, colistin, and to a lesser extent to fosfomycin. In our disseminated infection case, the colistin- and tigecycline-resistant CRE made the treatment more complex and challenging, and even more worrisome, because the *E. coli* harbored *fosA*3 and *bla*<sub>CTX-M-14</sub>. In addition to the surgical intervention, amoxicillin–clavulanic acid combined with aztreonam was used on the basis of the combined antibiotic susceptibility test, and the bacterium was finally eliminated. This regimen may promisingly combat *E. coli* harboring *mcr*<sub>1</sub> and *bla*<sub>NDM</sub>–5. Combined antibiotic susceptibility test in vitro is vital in the so-called “post-antibiotic” era.

**Conclusion**

This is the first report of a colistin-, carbapenem- and tigecycline-resistant *E. coli* in China. Tigecycline resistance acquired during tigecycline therapy is a great concern because tigecycline is a drug of the last resort for carbapenem-resistant Gram-negative bacteria infections. Furthermore, the transmission of such extensively drug-resistant isolates may pose a great threat to public health. Appropriate treatment schemes that effectively kill the XDR pathogen are imperative.

**Ethics statement**

The clinical isolates of *E. coli* E-FQ, E-XS, E-DB were part of routine hospital laboratory procedure. The patient’s next of kin had provided written informed consent for details to be used in the manuscript.

**Acknowledgments**

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**Author contributions**

Conceived and designed the experiments: YY and PZ; performed the experiments: QW, PZ, YW, and DZ; analyzed the data: DZ, FZ, and YJ; wrote the manuscript: QW, PZ, XL, and XD. All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


Supplementary materials

Figure S1 Escherichia coli isolates E-FQ, E-XS, and E-DB expressed indistinguishable band patterns by XbaI PFge analysis. Analysis by S1 nuclease PFGE and Southern blot showed the presence of the mcr-1 plasmid in a ~244–330 kb plasmid and blcaDNm-5 in a ~33–55 kb plasmid.
Abbreviation: PFGE, pulsed-field gel electrophoresis.

Table S1 Primers used in mapping the mcr-1 genetic environment and confirming mutations in acrB and lon

<table>
<thead>
<tr>
<th>Name of primers</th>
<th>Sequence</th>
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<tr>
<td>140-27-F</td>
<td>AAGAAAAACGGCAACACTCGC</td>
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<tr>
<td>140-27-R</td>
<td>TCGGGCATCTGGCAAAAGAAT</td>
</tr>
<tr>
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<td>TGGAAATGGGAACCTGTCG</td>
</tr>
<tr>
<td>S-140-R</td>
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</tr>
<tr>
<td>AcrB-RND-F</td>
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<tr>
<td>La-F</td>
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