Genomic characteristics of clinical multidrug-resistant Proteus isolates from a tertiary care hospital in southwest China

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Multidrug-resistant (MDR) Proteus, especially those strains producing extended-spectrum β-lactamases (ESBL) and carbapenemases, represents a major public health concern. In the present work, we characterized 27 MDR Proteus clinical isolates, including 23 Proteus mirabilis, three Proteus terrae, and one Proteus faecalis, by whole-genome analysis. Among the 27 isolates analyzed, SXT/R391 ICEs were detected in 14 strains, and the complete sequences of nine ICEs were obtained. These ICEs share a common backbone structure but also have different gene contents in hotspots and variable regions. Among them, ICEPmiChn2826, ICEPmiChn2833, ICEPmiChn3105, and ICEPmiChn3725 contain abundant antibiotic resistance genes, including the ESBL gene blaCTX-M-65. The core gene phylogenetic analysis of ICEs showed their genetic diversity, and revealed the cryptic dissemination of them in Proteus strains from food animals and humans on a China-wide scale. One of the isolates, FZP3105, acquired an NDM-1-producing MDR plasmid, designated pNDM_FZP3105, which is a self-transmissible type 1/2 hybrid IncC plasmid. Analysis of the genetic organization showed that pNDM_FZP3105 has two novel antibiotic resistance islands bearing abundant antibiotic resistance genes, among which blaNDM-1 is located in a 9.0 kb ΔTnJ25 bracketed by two copies of IS26 in the same direction. In isolates FZP2936 and FZP3115, blaKPC-2 was detected on an IncN plasmid, which is identical to the previously reported pT211 in Zhejiang province of China. Besides, a MDR genomic island PmGRI1, a variant of PmGRI1-YN9 from chicken in China, was identified on their chromosome. In conclusion, this study demonstrates abundant genetic diversity of mobile genetic elements carrying antibiotic resistance genes, especially ESBL and carbapenemase genes, in clinical Proteus isolates, and highlights that the continuous monitoring on their transmission and further evolution is needed.

KEYWORDS
Proteus, SXT/R391 ICEs, blaNDM-1, blaKPC-2, plasmid
Introduction

Proteus spp., belonging to the family of Morganellaceae of the order Enterobacterales (Adeolu et al., 2016), are widespread in the environment, and also inhabit in the intestines of humans and animals (O’Hara et al., 2000). Currently, ten species were included in the genus Proteus, among which, Proteus mirabilis is the most frequently isolated from clinical samples, and represents a major cause of urinary tract infections (Schaffer and Pearson, 2015). Proteus are intrinsically resistant to polymyxins, nitrofurans, tigecycline, and tetracycline, and are naturally susceptible to aminoglycosides, fluoroquinolones, and trimethoprim-sulfamethoxazole (Korytny et al., 2016). Multidrug-resistant (MDR), to 3 or more classes of antimicrobial agents (Falagas and Karageorgopoulos, 2008) Proteus have been increasingly reported, which generally produce extended-spectrum β-lactamases (ESBLs) and AmpC-type cephalosporinase, and show co-resistance to fluoroquinolones, aminoglycosides, and sulfamethoxazole-trimethoprim (Tumbarello et al., 2012; Korytny et al., 2016). Carbanepens are one of the last-resort antibiotics to treat severe infections caused by MDR Proteus, while the identification of carbapenemases, such as KPC-2 (Hu et al., 2012), NDM-1 (He et al., 2021), OXA-48 (Fursova et al., 2015), OXA-23, and OXA-58 (Bonnin et al., 2020) in Proteus causes great clinical concern. Plasmid-mediated horizontal transfer plays an important role in the dissemination of carbapenemase genes in Proteus (Hua et al., 2020; Li et al., 2021).

Resistance genes on the chromosome are usually clustered into structures named genomic islands (GIs), which are distinct regions of a bacterial chromosome that have been acquired via horizontal transfer (Partridge et al., 2018). Integrating and conjugative elements (ICEs) are a specific GI with mobility functions that can be integrated into the host chromosome, and also excised as a circular intermediate to be self-transferred to a recipient cell via conjugation (Johnson and Grossman, 2015). The large SXT/R391 family ICEs are characterized by a conserved integrase, which catalyzes their integration into the 5′ end of the chromosomal prfC gene (encoding the peptide chain release factor 3) by site-specific recombination (Bioteau et al., 2018). SXT/R391 ICEs share 52 nearly identical core genes that mediate integration/excision, conjugative transfer, and regulation, and also contain variable DNA regions, dubbed hotspots (HS) 1–5 and variable regions (VR) 1-IV (Wozniak et al., 2009), wherein SXT/R391 acquire new variable DNA sequences that confer element-specific phenotypes, such as resistance to antibiotics (Rodriguez-Blanco et al., 2012). Several SXT/R391 elements have been identified in Proteus, and they serve as vehicles of clinical important resistance genes, such as blaCTX-M-1, blaNDM-1 (He et al., 2021), tet(X6) (Peng et al., 2020) and blcEX-M-46 (Li et al., 2018), which constitutes a severe concern. At present, SXT/R391 ICEs have been widely studied in food-producing animals in China (Li et al., 2016, 2018; Wang et al., 2021), while their prevalence and genetic characteristics in the clinical isolates are not well understood.

The present study was conducted to characterize MDR Proteus clinical isolates from a tertiary care hospital in Sichuan province, China. We performed whole-genome sequencing and analysis to investigate their genetic features about resistance determinants, with a focus on the diversity of genetic structure of SXT/R391 ICEs and the genetic contexts of carbapenemase-encoding genes. Also, the transfer capability of ICEs and carbapenemase-encoding plasmids were determined by conjugation assays.

Materials and methods

Bacterial isolates

Proteus isolates were isolated from clinical specimens of patients at the Affiliated Traditional Chinese Medicine Hospital of Southwest Medical University, in Sichuan Province, China, from January to December of 2021. This study was approved by the Ethics Committee of the Affiliated Traditional Chinese Medicine Hospital of Southwest Medical University. Written informed consent from the patients was exempted for this study, since the present study only focused on bacteria and the strains were isolated as a part of the routine hospital laboratory procedures. Species identification was carried out by 16S rRNA gene sequencing analysis (Lane, 1991). Antibiotic susceptibility testing was performed using the Biomerieux Vitek-2 system, and the results were interpreted by the clinical breakpoints defined by the Clinical and Laboratory Standards Institute standards for Enterobacterales (CLSI, M100). Escherichia coli strain ATCC 25922 was used as the quality control. The antibiotics used in this study was provided in Supplementary Table S1. The minimum inhibitory concentrations (MICs) of meropenem against the isolates were determined using the microdilution broth methods and the disk diffusion methods of Bauer and Kirby following recommendations of the CLSI. MDR strains were defined as non-susceptibility to at least one agent in three or more of the following antibiotic groups: β-lactam-β-lactam inhibitor combinations, cephalosporins, fluoroquinolones, aminoglycosides or sulfamethoxazole and trimethoprim (Korytny et al., 2016).

Genomic DNA sequencing and data analysis

Genomic DNA was prepared using the QIamp DNA Mini Kit (Qiagen) following the manufacturer’s guidelines. Whole genome sequencing was performed on the HiSeq 2000 (Illumina, San Diego, CA, United States) platform using a paired-end library with an insert size of 150 bp by the Beijing Tsingke Bioinformatics Technology Co. Ltd. Four isolates (FZP2936, FZP3115, FZP3105 and FZP2826) were additionally sequenced on the long-read MinION sequencer (Nanopore, Oxford, United Kingdom). Both the long MinION reads and short Illumina reads were de novo-assembled using Unicycler under the conservative mode.

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(Wick et al., 2017). Pilon was employed to correct the assembled contigs with Illumina reads (Walker et al., 2014). Annotation was carried out using the RAST tools (Aziz et al., 2008) combined with BLASTp/BLASTn searches against the UniProtKB/SwissProt database (Boutet et al., 2016).

The identification of Proteus species was performed by average nucleotide identity (ANI) analysis with JSpeciesWS.1 Plasmid incompatibility types and multilocus sequence typing were identified using PlasmidFinder 2.1 (95%, minimum threshold for identity; 60%, minimum coverage) and pMLST 2.0 (Carattoli and Hasman, 2020). Antibiotic resistance genes (ARGs), insertion elements (ISs) and integrons were predicted using ResFinder (90%, minimum threshold for identity; 60%, minimum coverage; Bortolaia et al., 2020), ISfinder (Siguer et al., 2006) and INTEGRALL (Moura et al., 2009). The presence of SXT/R391 ICE was screened by targeting the conserved integrase gene (intSXT) from the whole sequenced genomes. The contigs of SXT/R391 ICE were extracted and assembled against the reference ICE in the genome of FZP3105, with gaps between contigs closed by PCR sequencing. Linear sequence alignment was performed using BLAST and visualized with Easyfig 2.2.3 (Sullivan et al., 2011).

Phylogenetic analysis

The genome sequences of other representative Proteus isolates in China from the literature were retrieved from the GenBank. Genetic relationship between the Proteus isolates in this study and these reference strains was assessed based on single nucleotide polymorphisms (SNPs) in their core genomes, as previously described with minor modification (Li et al., 2022). Briefly, Genomes were annotated using Prokka, and the generated GFF3 files were piped into Roary to create a core genome alignment. SNPs were extracted using snp-sites v2.3.2. A maximum-likelihood phylogenetic tree based on the SNPs was constructed using FastTree version 2.1.10 under the GTR model. Similarly, a likelihood phylogenetic tree based on the SNPs was constructed using the GTR model with the rifampin-resistant E. coli strain J53 as the recipients, as described previously with minor modification (Li et al., 2021). After the donor strain and recipient were grown to exponential stage (the optical density at 600 nm reaches ~0.5), mix them at a ratio of 1:1, and incubate at 37°C for 24 h. Transconjugants were selected on Luria-Bertani (LB) agar plates containing 4 μg/ml meropenem, 16 μg/ml gentamicin or 2 μg/ml tigecycline plus 400 μg/ml rifampin or 150 μg/ml sodium azide. The presence of blaNDM-1, and SXT/R391 ICE was confirmed by PCR using the primers blaNDM-F 5′-ATTACTAGGCCCTCGATTGC-3′ /blaNDM-R 5′-GCCT CTTGTCACTCGAATCG-3′, and sxtintF 5′-TCGATGATGGTGCTCTAGCTG-3′ /sxtintR 5′-TCAGTTAGCTGGCTCTGATGC-3′ (Sato et al., 2020), respectively, with the following conditions: 95°C for 5 min, and 30 cycles of amplification consisting of 30s at 95°C, 30 s at 53°C, and 1 min 30 s at 72°C, followed by a final elongation step for 5 min at 72°C.

Results and discussion

Sources, resistance phenotype, and genotypic diversity of MDR Proteus isolates

54 Proteus strains were isolated from clinical specimens, and 27 of them were identified as MDR strains (Table 1). These MDR isolates were obtained from urine (n = 13, 48.1%), wound secretion (n = 8, 29.6%), sputum (n = 2, 7.4%), blood (n = 2, 7.4%), sanies (n = 1, 3.7%), and drainage (n = 1, 3.7%). ANI analysis indicated that 27 MDR Proteus strains belong to P. mirabilis (n = 23), Proteus terrae (n = 3), and Proteus faecis (n = 1). In addition to their intrinsic resistance profiles, these MDR Proteus isolates also showed high levels of resistance to ampicillin (n = 27, 100%), cefaloti (n = 19, 70.4%), ceftriaxone (n = 19, 70.4%), ampicillin/sulbactam (n = 18, 66.6%), sulfamethoxazole-trimethoprim (n = 16, 60.0%), meropenem (n = 3, 11.1%), and tobramycin (n = 11, 40.7%). Some MDR isolates were also resistant to aztreonam (n = 6, 22.2%), norfloxacin (n = 9, 33.3%), impenem (n = 5, 18.5%), meropenem (n = 3, 11.1%), cefazidime (n = 2, 7.4%), and pipercillin/tazobactam (n = 2, 7.4%; Table 1). Notably, all strains remained susceptible to amikacin.

We sequenced the genomes of all 27 MDR isolates on the Illumina platform (Supplementary Table S2). Genomic analysis revealed that 57 different ARGs were detected in the 27 MDR Proteus strains, and 23 (85.2%) of them were carried at least 13 ARGs (Figure 1). Of the detected β-lactamase genes, blaCTX-M-65, the only ESBL-producing gene, was the most prevalent (n = 13, 48.1%), followed by the non-ESBL blaOXA-1 (n = 12). Carbapenemase gene blaKPC-2 was found in two isolates, FZP2936 and FZP3115, in combination with blaTEM-1β-blaNDM-1. Was only detected in one isolate, FZP3105, which also harbors blaCTX-M-65, blaOXA-1, and blaOXA-19. Accordingly, FZP2936, FZP3115, FZP3105, and one blaCTX-M-65-positive isolate FZP2826 were selected for further...
| Strain  | Species               | Specimen       | Gender/age (year) | ICE/variants of PmGRI1 | Resistance phenotype |
|---------|-----------------------|----------------|-------------------|------------------------|---------------------|
| FZP1665 | Proteus mirabilis     | Urine          | F/49              | PmGRI1                 | AMP, CIP, GEN, NIT, SXT, TCY, TGC |
| FZP2056 | Proteus mirabilis     | Urine          | M/75              | ICE                    | AMP, CIP, CZO, LVX, MEF, NIT, NOR, SAM, SXT, TCY, TGC |
| FZP2095 | Proteus mirabilis     | Wound secretion| M/56              |                        | AMP, ATM, CAZ, CIP, CRO, CTX, CXM, CZO, FEP, FOX, SAM, SXT, TGC |
| FZP2128 | Proteus mirabilis     | Wound secretion| M/41              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NIT, SAM, SXT, TCY, TGC, TOB |
| FZP2958 | Proteus mirabilis     | Urine          | M/70              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NIT, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP3803 | Proteus mirabilis     | Wound secretion| M/35              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NIT, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP4264 | Proteus mirabilis     | Urine          | F/49              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, NIT, PIP, SXT, TCY, TGC, TOB |
| FZP4349 | Proteus mirabilis     | Wound secretion| F/58              | ICE                    | AMP, CIP, CXA, CXM, CZO, NIT, SXT, TCY, TGC |
| FZP4423 | Proteus mirabilis     | Blood          | M/82              | ICE                    | AMC, AMP, CIP, CPD, CIP, CTX, CXM, CZO, GEN, IPM, LVX, MEF, NOR, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP4515 | Proteus mirabilis     | Wound secretion| M/49              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NOR, PIP, SXT, TCY, TGC |
| FZP4515 | Proteus mirabilis     | Wound secretion| M/49              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NOR, PIP, SXT, TCY, TGC |
| FZP4515 | Proteus mirabilis     | Wound secretion| M/49              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NOR, PIP, SXT, TCY, TGC |
| FZP1665 | Proteus mirabilis     | Urine          | M/66              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, LVX, MEF, NOR, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP2024 | Proteus mirabilis     | Urine          | M/77              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, NIT, SXT, TCY, TGC |
| FZP2561 | Proteus mirabilis     | Urine          | F/49              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, NIT, PIP, SXT, TCY |
| FZP2833 | Proteus mirabilis     | Drainage       | F/49              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, PIP, SAM, SXT, TCY, TGC |
| FZP2937 | Proteus mirabilis     | Wound secretion| M/32              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, NOR, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP3043 | Proteus mirabilis     | Urine          | F/37              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, NIT, SAM, SXT, TCY, TGC, TOB |
| FZP3105 | Proteus mirabilis     | Sanies         | M/32              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, IPM, LVX, MEF, MEM, NOR, PIP, SAM, SXT, TCY, TGC |
| FZP3320 | Proteus mirabilis     | Urine          | F/56              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, MEM, NOR, PIP, SAM, SXT, TCY, TGC |
| FZP3364 | Proteus mirabilis     | Wound secretion| F/82              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, MEM, NOR, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP3406 | Proteus mirabilis     | Urine          | F/52              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, GEN, NIT, SAM, SXT, TCY, TOB |
| FZP3725 | Proteus mirabilis     | Blood          | M/81              | ICE, PmGRI1            | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, LVX, MEF, PIP, SAM, SXT, TCY, TGC |
| FZP2826 | Proteus mirabilis     | Urine          | M/58              | ICE                    | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, FEP, IPM, LVX, MEF, NOR, PIP, SAM, SXT, TCY, TGC, TOB |
| FZP2936 | Proteus mirabilis     | Urine          | M/58              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, ETP, GEN, IPM, MEM, MFX, NIT, NOR, PIP, SAM, SXT, TCY, TGC, TOB, TZP |
| FZP3115 | Proteus mirabilis     | Wound secretion| M/58              | PmGRI1                 | AMP, CIP, CIP, CPD, CRO, CTX, CXM, CZO, ETP, GEN, IPM, MEM, MFX, NIT, NOR, PIP, SAM, SXT, TCY, TGC, TOB, TZP |

AMC, amoxicillin/clavulanic acid; AMP, ampicillin; ATM, aztreonam; CIP, ciprofloxacin; CAZ, ceftazidime; CEP, cefalotin; CPD, cefpodoxime; CRO, ceftriaxone; CSL, cefoperazone/sulbactam; CTT, cefotetan; CTX, cefotaxime; CXA, cefuroxime axetil; CXM, cefuroxime; CZO, cefazolin; CZX, ceftizoxime; ETP, ertapenem; FEP, cefepime; FOX, cefotaxim; GEN, gentamicin; IPM, imipenem; LVX, levofloxacin; MEM, meropenem; MEF, moxifloxacin; NIT, nitrofurantoin; NOR, norfloxacin; PIP, piperacillin; SAM, ampicillin/sulbactam; SXT, trimethoprim/sulfamethoxazole; TCY, tetracycline; TGC, tigecycline; TOB, tobramycin; TZP, piperacillin/tazobactam.
analyses by whole-genome sequencing using the Nanopore technology. Three different quinolone resistance genes were present: \(\text{aac}(6')\text{-Ib-cr}\) (14/27, 51.9%), \(\text{qnrA}1\) (2/27, 7.4%), and \(\text{qnrD}1\) (8/27, 29.6%). Aminoglycoside resistance genes were commonly detected in these MDR \textit{Proteus} strains, with \(\text{aadA}1\) (21/27, 77.7%), \(\text{aph}(3'\text{-})\text{-Ib}\) (17/27, 63.0%), and \(\text{aph}(6)-\text{Id}\) (17/27, 63.0%) being the most prevalent. All the isolates, except for FZP1097 and FZP3364, contained both \(\text{sul}\) and \(\text{dfrA}\) genes, conferring co-resistance to sulfamethoxazole-trimethoprim. Albeit only one known resistance gene \(\text{hugA}\) (encoding a class A \(\beta\)-lactamase conferring third-generation cephalosporin-resistance) was detected, FZP1097 was resistant to several antimicrobial agents, such as ciprofloxacin and sulfamethoxazole-trimethoprim in addition to cefotaxime, indicating that some unknown resistance mechanisms may be involved in its MDR phenotype.

Population structure analysis of MDR \textit{Proteus} isolates

To understand the genetic relationship, the 27 MDR \textit{Proteus} strains were compared to other \textit{Proteus} isolates from different geographic locations in China, and a core genome-based phylogenetic tree was constructed, which revealed two distinct groups (Figure 1). The large group includes all the \textit{P. mirabilis} strains with a high degree of whole-genome homogeneity (0–5,555 SNPs, Supplementary Table S3), and the other one consists of different non-\textit{P. mirabilis} strains of the \textit{Proteus} genus. From the phylogenetic tree, some isolates from different patients are tightly clustered, for example FZP2833/FZP3803 and FZP4349/FZP3364, hinting a common origin and cryptic transmission in this hospital. In addition, clonal relationship was also noticed between some isolates in this study and those from food animals or humans in other locations of China (Supplementary Table S4), for example FZP2958/XH1653 (\textit{P. mirabilis}, humans, Zhejiang of China, 2015) and FZP2095/ZF2 (\textit{P. terrae}, animals, Jiangsu of China, 2018), suggesting the possible circulation and clonal transmission of these MDR \textit{Proteus} strains amongst animals and humans across China. Active monitoring the spread of MDR \textit{Proteus} strains in the context of ‘One Health’ (environmental, animal, and human sectors) is an essential part in combating antimicrobial resistance.

Genetic features of ICE in MDR \textit{Proteus} isolates

Fourteen out of the 27 \textit{Proteus} strains were positive for the \(\text{int}_{\text{SXT}}\) gene (>97% identity to that of SXT), and all of these SXT/
R391 ICE-harboring isolates were P. mirabilis, except one P. terrae (Table 1). Based on the genome data, the complete sequences of nine SXT/R391 ICEs were successfully assembled, ranging in size from 83,975 bp to 130,923 bp, and five were fragmented in two or more contigs (Figure 2). Among them, ICEPmiChn2826 and ICEPmiChn2833 are almost identical, with only several different bases in variable regions. ICEPmiChn2024 and ICEPmiChn2561 are only differed by an insertion of a 1,063 bp truncated IS26 between traI and traD in the former.

Analysis of genetic organization showed that the nine ICEs shared a common backbone structure with most SXT/R391 ICEs, but also contained DNA sequences that are relatively specific for individual elements, comprising five hotspots (HS1-5) and three variable regions (VRII, VRIII and VRII; Figure 2), as described previously (Wozniak et al., 2009). Four ICEs (ICEPmiChn2024, ICEPmiChn2561, ICEPmiChn4280, and ICEPteChn4349) harbor a mutL gene (encoding a putative DNA mismatch repair protein) in the VRII (xis-int), and ICEPmiChn4280 solely has a mer operon in the VRIV (traG-ex). All nine ICEs harbored VRIII (disrupting the rumB gene), and three ICEs (ICEPmiChn2826, ICEPmiChn2833, and ICEPmiChn3803) have a MDR region bearing ARGs floR, strB/A, and sul2, conferring resistance to chloramphenicol, streptomycin, and sulfamethoxazole, respectively. Specially, ICEPmiChn4280 has a multi-drug resistance RND efflux pump gene cluster, tmexCD3-toprJ3, in VRIII (Wang et al., 2021). HS1 (s043-traL), HS2 (traA-s054), HS4 (traN-s063), and HS5 (s026-traI) were detected in all nine ICEs, while HS3 (s073-traE) was only found in ICEPmiChn2024 and ICEPmiChn2561, with inserted genes encoding an integrase, a dihydrofolate reductase-like protein and a hypothetical protein. Gene clusters encoding diverse restriction-modification (RM) systems conferring resistance to bacteriophages (Balado et al., 2013) are commonly found in HS5 in these ICEs, including the BREX phage resistance system (brxA-brxB-brxC-pglX-pglZ-brxL; Slattery et al., 2020). Additionally, genes encoding endonuclease, ATPase, methyltransferase, helicase, and the mrr restriction system are also detected in this region (Figure 2).

Abundant ARGs are present in HS4 in ICEPmiChn2826, ICEPmiChn2833, ICEPmiChn3105, and ICEPmiChn3725. Structural comparison showed that the four ICEs share similar gene contents in the HS4 region, which is also similar to that in ICEPmiChnXH1653 detected in a P. mirabilis strain from a urine sample of a patient in Zhejiang, China, in 2015 (He et al., 2021; Figure 3), indicating a common origin. Like the scenario in ICEPmiChnXH1653, the HS4 in ICEPmiChn3105 and ICEPmiChn3725 are also present as an ISppu12-mediated region, but the two tandem copies of a blaNDM-1-bearing ISCR1 element downstream of sul1, and two copies of IS26 interrupting tnpR are not detected. It has been known that a second IS26 preferentially inserted into adjacent positions of the existing IS26, and the recombination between the two copies of IS26 would cause deletion, insertion or translocation (Partridge et al., 2018). The

![Genetic organization of the ICEs in this study. The upper line shows the backbone of SXT/R391 ICEs with conserved core genes. Arrows in orange, purple, green and amaranth indicate genes involved in site-specific excision and integration (xis and int), DNA repair (rumAB), conjugative transfer (tra) and regulation (setCDR). Other core genes are indicated by light gray arrows. Under the backbone, hotspots (HS1–HS5) and variable regions (VRII–IV) are shown, with thin arrows indicating the sites of their insertion. ARGs, insertion sequences, BREX phage resistance system, and restrictive modification systems are also detected in this region.](10.3389/fmicb.2022.977356)
HS4 in ICEPmiChn2826 (ICEPmiChn2833) seems to be a variant of ICEPmiChn3105 with the deletion of 9,781 bp HS4 region and 15,456 bp of adjacent backbone, very likely resulting from an insertion of the second IS26 inside the traG and the subsequent recombination action of IS26 (Figure 3). The HS4 in ICEPmiChn3105 and ICEPmiChn3725 are highly similar (99.97% nucleotide identity at 92% coverage), and an IS26-mediated excision of a 4,311 bp region bearing aph(3′)-Ia and fosA3 downstream of tetR, and an 813 bp deletion upstream of tnpR represent two major modular differences of them (Figure 3). This finding suggests that these MDR ICEs are undergoing rapid evolution within healthcare environments.

By BLASTn analysis, the nine SXT/R391 ICEs exhibited similarity to many different ICEs recovered from food animals and humans in different locations of China (Supplementary Table S5), indicating wide spread of them. The core gene phylogenetic analysis showed that ICEPmiChn2826, ICEPmiChn2833, and ICEPmiChn3803 are closely related to the reference MDR ICEs, ICEPmiChn1 (P. mirabilis, chicken, Hubei of China), ICEPmiChnBCP11 (P. mirabilis, swine, Sichuan of China), and ICEPmiChnChSC1905 (P. mirabilis, swine, Sichuan of China; Figure 4; Supplementary Table S6). ICEPmiChn3105 is clustered with ICEPmiChnCC15031 that was detected in a P. mirabilis strain recovered from the intestinal contents of humans in Zhejiang, China. The phylogenetic analysis indicated that ICEPteChn4349 serves as a potentially novel SXT/R391 element as it forms a distinct clade separated from other representative ICEs (Figure 4; Supplementary Table S6). ICEPmiChn2024, ICEPmiChn2561, and ICEPmiChn4280 are more distantly related to the ICEs in this study. ICEPmiChn4280 has the most closely genetic relationship with ICEPmiChnRGF134-1 (P. mirabilis, swine, Jiangsu of China), and ICEPmiChn2024 (ICEPmiChn2561) is closest to ICEPteChnS24-2-1 (P. terrae, Cacatua, Guangzhou of China), with 0 and 2 SNPs, respectively (Figure 4; Supplementary Table S6). These findings suggest the cryptic dissemination of these ICEs in Proteus strains on a China-wide scale.

According to the species tree (Figure 1), we found that strains FZP2024 and FZP2561 are little related, but share almost identical ICEs (ICEPmiChn2024 and ICEPmiChn2561). The closely related strains FZP2826 and FZP3803 bear ICEs (ICEPmiChn2826 and ICEPmiChn3803) with markedly different HS4 and HSS contents. Besides, strains FZP4349 and FZP3364 are tightly clustered, and the latter is devoid of an SXT/R391 element on the chromosome while the former possesses the ICEPteChn4349. These findings suggest the independent acquisition and horizontal transmission of SXT/R391 ICEs across the Proteus population (Sato et al., 2020).
We also detected the circular intermediate of ICEs by using the primers LE4 and RE4 as previously described (Lei et al., 2016). Results showed that the circular form of ICEs could be detected in all 14 ICE-harboring strains, revealing a potential transmission pattern of them. To determine the transfer ability of these ICEs, we selected three ICE-carrying isolates (FZP2826, FZP3725 and FZP4280) for the conjugation experiments. Results showed that ICE \textit{PmiChn3725} could be transferred to \textit{E. coli} J53 at a frequency of \(9 \times 10^{-5}\) transconjugants per recipient cell. Antimicrobial susceptibility analysis showed that the acquisition of ICE\textit{PmiChn3725} enables \textit{E. coli} J53 to become resistant to gentamicin, cefotaxime, aztreonam, ampicillin/sulbactam, and trimethoprim/sulfamethoxazole (Supplementary Table S7), despite that resistance to aztreonam conferred by CTX-M-65 in FZP3725 remains within the sensitivity range. We hypothesize that the hydrolysis of aztreonam by CTX-M-65 is possibly inhibited in FZP3725 by an unknown regulatory mechanism, but it works normally in \textit{E. coli} J53. The conjugation of ICE\textit{PmiChn2826} and ICE\textit{PmiChn4280} were failed despite repeated attempts, suggesting that both ICEs are not self-transmissible. The non-transferability of ICE\textit{PmiChn2826} may be caused by the truncated \textit{traG}, and the deleted \textit{traF} and \textit{traH} (Figure 3).

**Genetic features of the \textit{bla}_{NDM-1}-harboring plasmid pNDM_FZP3105**

\textit{P. mirabilis} FZP2937 and FZP3105 were recovered from the same patient, and were sampled 18 days apart, from wound secretion and sanies, respectively. FZP2937 and FZP3105 are clonally related as they share identical core genomes (0 SNP). Despite that, the two strains exhibited different phenotypic resistance (Table 1), and the most noteworthy feature is that FZP3105 shows resistant to meropenem and ceftazidime, to which FZP2937 is susceptible. Genome data revealed that the later strain FZP3105 additionally acquired an NDM-1-producing MDR plasmid pNDM_FZP3105. This plasmid is 205,118 bp in size, and belongs to IncC ST3 (A053-parA-parB-repA allele number 1–2–2–2) incompatibility group. BLASTn analysis showed that it has >99.8% nucleotide identity (≥77% coverage) to pCMC307\_P2 (CP079626, \textit{Klebsiella pneumoniae}, India), p13ARS\_GMH0099.
pNDM_FZP3105 is a type 1/2 hybrid IncC plasmid, as it contains orfI832 (characteristic of type of 1), and $\Delta$rs2 (characteristic of type 2), lacking of two additional sequences i1 and i2 (Harmer and Hall, 2014). The backbone of pNDM_FZP3105 is 112,618bp in size with an average 51.05% G + C content. Nucleotide (nt) 56,193–2014 lacking of two additional sequences i1 and i2 (a type 1/2 hybrid IncC plasmid, as it contains $\Delta$rs2, Aeromonas hydrophila, South Korea), and nt 1–2,522, 11,597–28,308, 163,334–165,885, and 185,018–205,098 showed 99.98% identity to that of type 1 IncC reference plasmid pR148 (X1414747, Aeromonas hydrophila). Compared to the prototype ARI-A of a complex transposon immediately upstream of the 452 bp antibiotic resistance island (ARI)-A, ARI-B, and the IS$_{\text{Cmr}}$ (increased meropenem resistance in EC600 by at least 8-fold E. coli 163,334–165,885, and 185,018–205,098 showed 99.98% identity with an average 51.05% G + C content. Nucleotide (nt) 56,193–2014 lacking of two additional sequences i1 and i2 (a type 1/2 hybrid IncC plasmid, as it contains $\Delta$rs2, Aeromonas hydrophila, South Korea), and nt 1–2,522, 11,597–28,308, 163,334–165,885, and 185,018–205,098 showed 99.98% identity to that of type 2 IncC reference plasmid pR55 (JQ010984, K. pneumoniae, France; Supplementary Figure S1). Conjugation assays showed that pNDM_FZP3105 was able to conjugate into E. coli EC600, and the acquisition of pNDM_FZP3105 greatly increased meropenem resistance in EC600 by at least 8-fold (Supplementary Table S8).

pNDM_FZP3105 carries three accessory modules, namely the antibiotic resistance island (ARI)-A, ARI-B, and the IS$_{\text{Cmr}}$-aac(3)-IId-tmrB module. ARI-A (nt 127,160 to 163,333) is located immediately upstream of the 452 bp rhs2 remnant (Figure 5B). Compared to the prototype ARI-A of a complex transposon structure bounded by 38-bp inverted repeats (IRs) of $\Delta$n1696 top and pDU mer modules interrupted by either IS4321 or IS0705 (Harmer and Hall, 2015), like the case in pRMH760 (KF976746, K. pneumoniae, Australia), ARI-A in pNDM_FZP3105 has only retained the IS4321-top AR structure, though it is interrupted by the Tn6260 that is truncated by an ISPn3 inside of topA. Instead of In34 in ARI-A in pRMH760, an $\Delta$n633-like segment intII-dfrA1-arr2-bla$_{\text{kar1}}$-aadA1-quad1-sul1-ISCR1 was present in pNDM_FZP3105, followed by a 6.5-kb region carrying an ISV9, a DNA repair ATPase and three hypothetical proteins, and a 10.7-kb module ISCR1-ISEc28-rrmB-ISEc29-msr(E)-mph(E)-IS15. BLASTn analysis showed that ARI-A in pNDM_FZP3105 has 99.9% nucleotide identity at 59% coverage to pT211-top, E. coli (United States), but with genetic element insertions, deletions, and replacements (Figure 5B). These findings suggest that ARI-A in pNDM_FZP3105 is a novel MDR mosaic region. lnu(G), which confers resistance to lincomycin by nucleotidylation, is commonly found on the chromosomes of Enterococcus, and also on the plasmids surrounding $\text{luxS}$ gene on the chromosome, as described in P. mirabilis strain T21 (Hua et al., 2020). The finding suggests the likelihood of a wide dissemination of the pT211-like blac$_{\text{KPC-2}}$-bearing plasmids in proteus in China, which warrant more surveillance. Given that FZP2936 and FZP3115 have close genetical relationship to strain T21 (7 SNPs; 1,457 bp downstream of repA, breaking and truncating the orf190 gene (encoding a hypothetical protein) downstream of cysH (Supplementary Figure S1). An insertion in this location is barely seen in IncC plasmids, and similar (>99% coverage, >99% identity) aac(3)-IId-tmrB module is only identified in pVF3N-blaoXA-193K (CP089604, Vibrio furnissii, hospital sewage, China) by BLASTn analysis. The mobilization mechanism of the IS$_{\text{Cmr}}$-aac(3)-IId-tmrB module remains unknown.

#### Characteristics of blac$_{\text{KPC-2}}$-carrying Proteus mirabilis FZP2936 and FZP3115

FZP2936 and FZP3115 are two subsequent isolates from a single patient, from urine and wound secretion, respectively, 19 days apart. The two strains share identical core genomes as well as resistance profiles, showing the clonal nature of them. FZP2936 and FZP3115 both have a 4,245,458-bp circular chromosome and two closed plasmids. 15 ARGs were identified in both strains, and all of them, except blac$_{\text{KPC-2}}$, are located on the chromosome. blac$_{\text{KPC-2}}$ is present on 24,225-bp IncN plasmids pKPC_FZP2936 and pKPC_FZP3115, which are identical to the previously reported pT211 (CP017083) from a clinical P. mirabilis strain T21 in 2013, in Zhejiang province (approximately 1,830 km apart in distance from Sichuan), China (Hua et al., 2020). The finding suggests the likelihood of a wide dissemination of the pT211-like blac$_{\text{KPC-2}}$-bearing plasmids in proteus in China, which warrant more surveillance. Given that FZP2936 and FZP3115 have close genetical relationship to strain T21 (7 SNPs; Figure 1), and that pT211 is not self-transferable as previously revealed (Hua et al., 2020), the diffusion of pT211-like plasmids is likely due to expansion of bacterial clones. The dfrA1-sat2-ada1 gene cassettes in both strains are included in the Tn7-like transposon 20bp downstream of the glmS gene on the chromosome, as described in P. mirabilis strain T21 (Hua et al., 2020). The remainder of the chromosomal ARGs were mainly clustered in a 64,829 bp MDR genomic island PmGR11 (Lei et al., 2020), which was later identified as a variant of PmGR11-YN9 (Ma et al., 2021; MW699445, P. mirabilis, animal, China) by BLASTn analysis. Compared to the configuration of PmGR11-YN9, the PmGR11 in FZP2936 (FZP3115) have two IS26-mediated excision situating 1,339 bp upstream and 533 bp downstream of blac$_{\text{KPC-2}}$, with 14,053 bp and 20,022 bp in size, respectively, resulting in the loss of several ARGs (Figure 6). Besides, the
FIGURE 5
Genetic features of pNDM_FZP3105. (A) Circular comparison of pNDM_FZP3105 with pCMC307P_P2, p13ARS_GMH0099 and pNDM-1_Dok01. Arrows on the outer ring indicate deduced ORFs and their orientations. Genes for replication (repA), conjugal transfer (tra), and plasmid maintenance (stbA, parAB) are highlighted in green. ARGs are highlighted in red. Two antibiotic resistance islands, ARI-A and ARI-B, are indicated.

(B) Organization of the ARI-A of pNDM_FZP3105, and comparisons to related regions. (C) Organization of the ARI-B of pNDM_FZP3105, and comparisons to related regions. Genes are denoted by arrows, and ARGs, integrase genes, IS26, and other transposase genes are highlighted in red, blue, yellow, and olive, respectively. The Tn6260 tnp and pDU mer modules are indicated in purple, and Tn6260 is in green. Regions of >90% identity are indicated by grey shading. Δ represents truncated genes.

PmGRI1 in this study was almost identical to that in P. mirabilis strain T21, except an IS26-mediated deletion of aph(3')-Ia 1,491 bp downstream of blaTEM-1B, and a tnsB-mediated reversal of a 7,275 bp region bearing a class I integron with the cassette array dfrA17-addA5-qacEΔ1-sul1 (Figure 6). These results highlight the high plasticity of PmGRI1, and also reinforce the great ability of IS26 to
accumulate ARGs, and to promote the diversity of the multidrug resistance regions in mobile genetic elements (He et al., 2021).

PmGRI1 is a new GI carrying various resistance genes initially identified in a P. mirabilis strain from chicken in 2018 (Lei et al., 2020), and was later detected in several P. mirabilis isolates of swine and chicken origins in different locations of China (Ma et al., 2021). To determine the prevalence of PmGRI1 in Proteus isolates in the same hospital, we detected the PmGRI1 by searching the integrase gene int in the remaining 25 strains, and found that 10 of them were positive for the PmGRI1 integrase gene (Table 1). This finding indicates the possible prevalence of PmGRI1 variants in Proteus strains in both the animals and humans. More data is needed to better understand the epidemiology and dynamic evolution of PmGRI1 in China.

**Conclusion**

In this study, we demonstrated the genetic diversity of MDR Proteus strains and spotlighted the important roles of GIs, especially SXT/R391 ICEs, and plasmids in capturing and spreading ARGs in Proteus. Our study also raises concern that these MDR Proteus strains might be circulating and undergoing rapid evolution amongst animals and humans across China. From ‘One Health’ perspective, active surveillance for MDR Proteus in the environment is urgently needed.

**Data availability statement**

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found in the article/Supplementary material.

**Author contributions**

YL: conceptualization, formal analysis, and writing-original draft. QL, YQ, and HC: methodology, resources, and formal analysis. XD, CF, and LZ: methodology and software. YZ and JS: writing-review and editing. LZ: conceptualization, writing-review and editing, and supervision. All authors contributed to the article and approved the submitted version.

**Funding**

This work was supported by National Natural Science Foundation of China (31900125), Scientific and technological project in Sichuan Province (2022JDRC0144), the Joint Funds of the Luzhou and Southwest Medical University Natural University Natural Science Foundation (2019LZXNYJD47 and 2020LZXNYJD34), and Central Government Funds for Guiding Local Scientific and Technological Development of Sichuan Province (2021ZYD0084). The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

**Conflict of interest**

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.

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**Supplementary material**

The Supplementary material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2022.977356/full#supplementary-material
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