First description of antimicrobial resistance in carbapenem-susceptible *Klebsiella pneumoniae* after imipenem treatment, driven by outer membrane remodeling

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**Abstract**

**Background:** The emergence of carbapenem-resistant *Klebsiella pneumoniae* (CRKP) poses a looming threat to human health. Although there are numerous studies regarding porin alteration in association with the production of ESBLs and/or AmpC β-lactamase, a systematic study on the treatment-emergence of porins alteration in antibiotic resistance does not yet exist. The aim of this study was to investigate the underlying mechanism of resistance of *K. pneumoniae* during carbapenem treatment.

**Results:** Here, we report three strains (FK-2624, FK-2723 and FK-2820) isolated from one patient before and after imipenem treatment during hospitalization. Antibiotic susceptibility testing indicated that the first isolate, FK-2624, was susceptible to almost all tested antimicrobials, being resistant only to fosfomycin. The subsequent isolates FK-2723 and FK-2820 were multidrug resistant (MDR). After imipenem therapy, FK-2820 was found to be carbapenem-resistant. PCR and Genome Sequencing analysis indicated that *oqxA*, and *fosA5* were identified in all three strains. In addition, FK-2624 also harbored *bla*<sub>SHV-187</sub> and *bla*<sub>TEM-116</sub>. The *bla*<sub>SHV-187</sub> and *bla*<sub>TEM-116</sub> genes were not detected in FK-2723 and FK-2820. *bla*<sub>DTAH-1</sub>, *qnrB4*, *aac (6′)-Ic*, and *bla*<sub>DTAH-12</sub>, *EreA2*, *CatA2*, *Sul1*, and *tetD*, were identified in both FK-2723 and FK-2820. Moreover, the genes *bla*<sub>DTAH-1</sub>, *qnrB4*, *aac (6′)-Ic* were co-harbored on a plasmid. Of the virulence factors found in this study, *ybtA*, *ICEKp6*, *mrkD*, *entB*, *iroN*, *mpa2-6*, *wzi16*, and capsular serotype K57 were found in the three isolates. The results of pairwise comparisons, multi-locus sequencing typing (MLST) and pulsed-field gel electrophoresis (PFGE) revealed high homology among the isolates. Sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) results showed that isolate FK-2820 lacked OmpK36, with genome sequence data validating that there was a premature stop codon in the *ompK36* gene and real-time RT-PCR suggesting high turnover of the *ompK36* non-sense transcript in FK-2820, with the steady-state mRNA level 0.007 relative to the initial isolate.

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Background

*Klebsiella pneumoniae* is a serious hospital-acquired pathogen causing infections such as urinary tract infections, pneumonia and bloodstream infections [1, 2]. The abuse of expanded-spectrum cephalosporins for the remedy of these organisms has contributed to the appearance of extended-spectrum β-lactamases (ESBLs) and AmpC-producing *K. pneumoniae*. Carbenepenems had been considered as the last resort for the treatment of infections caused by multidrug-resistant (MDR) isolates, due to their broadest antibacterial spectrum compared to other β-lactams. However, carbapenem-resistant *K. pneumoniae* (CRKP) have increasingly emerged under antibacterial drug selection pressure. The acquisition of plasmid-borne genes encoding carbapenemases is the mainly mechanism of CRKP since the first isolation of CRKP in America in 1996 [3]. Recently, it has been shown that carbapenem-resistance can also occur in the absence of carbapenemases, resulting instead from porin alteration so long as it occurs in association with the production of ESBLs and/or AmpC β-lactamases [4]. The porin OmpK36 is a nonspecific, passive diffusion pore in the outer membrane of *K. pneumoniae*, belonging to the OmpC porin group [5]. Several reports now support the contention that loss of OmpK36, coupled with ESBL, AmpC β-lactamase [4]. The current study focused on the acquisition of carbapenem resistance determinants in *K. pneumoniae* isolated from one inpatient, with emphasis on determining appropriate antimicrobial course of treatment.

Results

Clinical characteristics of the patient

This case was a 62-year-old male patient admitted to the department of neurosurgery in the First Affiliated Hospital of Wenzhou Medical University in October 2015 with headache. A diagnosis of cerebral aneurysm with subarachnoid hemorrhage was made, with a secondary pulmonary infection occurring after surgery. Antimicrobial treatment was started with 2 g of cefoperazone/sulbactam every 8 h (q8h), 0.5 g levofloxacin two times a day (bid.). On day 26, the fosfomycin-resistant *K. pneumoniae* FK-2624 was isolated from a sputum sample (Table 1). After that, due to pneumonia, the patient was switched to tigecycline via vein infusion (ivgtt.) with a loading dose of 100 mg. On Day 50, chest CT suggested atelectasis, which required a tracheotomy. On day 52, *Pseudomonas aeruginosa* was detected in purulent sputum, and the patient was switched to tobramycin with an ivgtt. Loading dose of 8 g q12h, and tobramycin with an ivgtt. Loading dose of 80 mg q12h due to susceptible to tobramycin (Table S3). On day 71, isolate FK-2723 was identified from the patient’s sputum sample and microbiological characterization found FK-2723 to be resistant to ceftriaxone, ceftazidime, levofloxacin, ciprofloxaxin, tobramycin, gentamicin and fosfomycin. This isolate was found to be susceptible to carbapenems, colistin and Tigecycline. On day 99, the patient was started on a 14-day course of combination treatment with imipenem and fosfomycin. On Day 113, FK-2820 was isolated and shown to be carbapenem-resistant but was susceptible to tigecycline (Table 1). Tigecycline treatment was initiated on Day 113 and after another 33 days hospitalization, the patient was transferred to a superior hospital in poor health, graded 6 by Glasgow coma scale (GCS) criterion (Fig. 1).

Antimicrobial susceptibility testing, characterization of resistance genes and virulence factors

The antibiotic resistance profiles, determined microbiologically, are shown in Table 1. The initial two isolates were carbapenem-sensitive. In the case of FK-2820, MICs for imipenem and ertapenem were 8 and 16 μg/mL, respectively. Although FK-2820 remained clinically sensitive to meropenem, the MIC of meropenem had increased four times, from 0.25 μg/mL to 1 μg/mL.

Initially, a phenotypic method was used for the detection of carbapenemase production. The mCIM analysis and EDTA synergy test are biochemical assay for carbapenemase activity, and the results were negative for FK-2820, suggesting that it does not express a genuine carbapenemase (Table 1). In order to independently test this result, to determine whether or not FK-2820 has a gene encoding a carbapenemase that would explain the observed carbapenem-resistant,
| Isolates codes | MIC (μg/ml) | Relative expression of Virulence factors |
|---------------|------------|-----------------------------------------|
| FK-2624       | 0.5        | K57, ybtA, ICEKp6, mrkD, entB, iroN, rmpA-6, wzi16 |
| FK-2723       | > 64       | K57, ybtA, ICEKp6, mrkD, entB, iroN, rmpA-6, wzi16 |
| FK-2820       | 8 > 64     | K57, ybtA, ICEKp6, mrkD, entB, iroN, rmpA-6, wzi16 |

Note:
1) MHT = Modified Hodge test, EDTA = EDTA synergy test
2) CRO = Ceftriaxone, CAZ = Ceftazidime, CTX = Cefotaxime, IPM = Imipenem, MEM = Meropenem, ETP = Ertapenem, LEV = Levofloxacin, TOB = Tobramycin, GEN = Gentamicin, AMK = Amikacin, FOS = Fosfomycin, COL = Colistin, TGC = Tigecycline
whole genome sequencing (WGS) was undertaken directed at the detection of mechanism in FK-2820 that distinguish it from FK-2723.

WGS was used to search for the presence of genes encoding virulence factors. Of the virulence factors found in this study, ybtA, ICEKp6, mrkD, entB, iroN, rmpA2–6, wzi16 and capsular serotype K57 were found in the three isolates (Table 1). These virulence genes are associated with high-affinity iron chelators or siderophores, iron uptake, biofilm formation and infection. Additionally, FK-2624 carries rmpA-2 which is not found in the subsequent isolates and is commonly carried on virulence plasmids such as KpVP-1 [8]. This raised the hypothesis that the initial isolate FK-2624 carried a virulence plasmid that had been lost from the subsequent isolates.

WGS was used to search for the presence of resistance genes, and these resistance markers were confirmed by PCR amplification using diagnostic primers, followed by DNA sequencing. Two resistance markers: oqxA, and fosA5, were identified in all three strains. In addition to these markers, FK-2624 also harbored bla$_{SHV-187}$ and bla$_{TEM-116}$ (Table 1). The bla$_{SHV-187}$ and bla$_{TEM-116}$ genes were not detected in the WGS data from FK-2723 and FK-2820. An initial hypothesis was that, like the virulence gene rmpA-2, they may have been carried on a plasmid found only in FK-2624.

The following antibiotic resistance markers: bla$_{DHA-1}$, bla$_{SHV-12}$, qnrB4, aac (6')-Ile, and EreA2, CatA2, SulI, and tetD, were identified in both FK-2723 and FK-2820. Sequence analysis suggested that the genes bla$_{DHA-1}$, qnrB4, aac (6')-Ile were co-harbored on a plasmid, and results of BLASTn analysis suggested that this plasmid displays a high level of sequence identity to plasmid pR47–309 (Genbank accession CP040697.1).

**Subtractive characterisation of the K. pneumoniae isolate genomes**

MLST analysis showed that the three strains belong to ST660, and are closely related (Fig. 2, Fig. S2, Table S4). An initial phylogenetic analysis of those few ST660 K. pneumoniae for which sequence data is available (Fig. 2b) showed that these closely related genomes from the NCBI database, collected in Beijing and in Sichuan (in 2012 and in 2015, respectively), are classified as Klebsiella pneumoniae subsp ozaena. The pairwise sequence identity was calculated to be > 98% across the genomes of the three strains collected in this study (Table S2), with no more than 410 single polymorphisms in the core genome (Table S2).

Alignment of the sequence data from the three isolates (Fig. 2c, Table S5) shows a short region of discrepancy in the length of the genome and dissimilarity of sequence characteristics, consistent with the presence of plasmids. FK-2624 has ~ 160 kb of distinguishing DNA, while FK-2723 has ~ 380 kb and FK-2820 has ~ 400 kb of DNA that might correspond to plasmids (Table S5). Differences in GC-content of those dissimilar regions compare to the rest of the alignment are consistent with this. These physical findings for the genome support the earlier observations concerning the differences in virulence and resistance genes in the three strains, and the putative plasmid in FK-2624 encompasses the genes encoding the TEM-116 beta-lactamase (bla$_{TEM-116}$) and
the virulence factors *iro-1* and *rmpA-2* specific to that isolate (Table 1).

Further analysis of the WGS data for FK-2327 and FK-2820 showed that the putative plasmid DNA harbors the genes *qnrB4*, *ereA2*, *catA2*, *sulI*, *tetD*. These genes confer resistance to aminoglycosides; fluoroquinolones, macrolides, phenicols, sulphonamides, and tetracyclines, respectively. Additionally, FK-2723 and FK-2820 plasmids carry the gene *bla*$_{DHA-1}$ encoding a beta-lactamase.

**OMP analysis**

Characterization of the outer membrane protein profile by SDS-PAGE showed that isolate FK-2820 did not express a full complement of porins (Fig. 3, Fig. S1). The relative expression levels for two of the three major protein species, namely OmpA and OmpK35 are similar for the three isolates and two other ST660 genomes from NCBI. The phylogenetic tree was generated using UPGMA clustering. A genetic similarity index scale is indicated by the vertical line. The phylogeny of the three isolates and two other ST660 genomes from NCBI. Labels in the nodes are bootstrap support. The scale represents the number of substitutions per site. The MAUVE alignment of the three genome sequence datasets. SNPs are represented by vertical red lines, colored boxes at the far end of the alignment correspond to putative plasmids acquired or lost from each of the subsequent isolated strains.

Fig. 2 PFG analysis and MLST of 3 *K. pneumoniae* isolates. a Relatedness was analyzed using Quantity One software (Bio-Rad Laboratories, USA). The phylogenetic tree was generated using UPGMA clustering. A genetic similarity index scale is indicated by the vertical line. b Phylogeny of the three isolates and two other ST660 genomes from NCBI. Labels in the nodes are bootstrap support. The scale represents the number of substitutions per site. c MAUVE alignment of the three genome sequence datasets. SNPs are represented by vertical red lines, colored boxes at the far end of the alignment correspond to putative plasmids acquired or lost from each of the subsequent isolated strains.

To understand the genetic basis for this loss of OmpK36 expression, we undertook direct sequencing of the *ompK36* gene. PCR-based evaluation was confirmed.

Fig. 3 OMP analysis from the *K. pneumoniae* strains. Extracts prepared from the three isolates (FK-2624, FK-2723 and FK-2820) and the control *K. pneumoniae* ATCC 13883 were subject to SDS-PAGE and staining with Coomassie blue. Lanes M, protein mass marker. ATCC 13883, an isolate with known expression of both OmpK35 and OmpK36.
by preliminary analysis of the WGS data, revealing a frameshift mutation caused by a substantial deletion of 239 nucleotides, changing the reading frame and truncating the sequence of OmpK36 in FK-2820. This partial protein in FK-2820 is incapable to assemble a \( \beta \)-barrel, and would be subject to quality control proteases such as DegP and BepA known to degrade other defective \( \beta \)-barrels [9, 10].

While the transcriptome evaluations did not mirror the protein-based evaluation, they did yield consistent results. Compared with the initial isolate \( K. \ pneumoniae \) FK-2624, expression of \( \text{ompK35} \) was similar in FK-2723 and FK-2820 (1.70 and 1.54, respectively). In the case of \( \text{ompK36} \) there was a ten-fold decrease in the steady-state level of the transcript that did not change the protein steady-state level in FK-2723 (Fig. 3, Fig. S1), but the decrease in \( \text{ompK36} \) transcript levels observed in FK-2820 was substantial and significant (0.007 relative to the initial isolate, \( P < 0.05 \)) (Table 1), consistent with the increased decay of at least some transcripts carrying premature stop codons [11].

**Discussion**

Bacterial antimicrobial resistance has emerged as one of the main concerns of public health and constitutes a major, global challenge of the near future [12]. As a key example, the emergence and global spread of CRKP poses clinical and therapeutic challenges, since they exhibit resistance to the majority of commonly used antibiotics [13]. Carbapenems are the class of \( \beta \)-lactam antibiotics with strong activity against many Gram-positive and Gram-negative species, making them key treatment options in infections resistant to other drugs. However, since the first report of KPC-2 strains of \( K. \ pneumoniae \) [14], various carbapenemase-producing isolates are being identified across the globe [15]. In addition to the carriage of carbapenemases such as KPC-2, porin alteration can also play a role in establishing CRKP phenotypes in strains which produce ESBLS or AmpC \( \beta \)-lactamases [4]. In the current study, resistance was monitored and characterized in three strains isolated from one patient, with the aim of our study being to investigate the mechanism of treatment-emergent carbapenem-resistance.

FK-2624 was isolated from the patient on day 26 and shown to be susceptible to most commonly used antimicrobial agents except fosfomycin. By contrast, the MDR strain FK-2723 isolated on day 71 only exhibited susceptibility to carbapenems, colistin and tigecycline. After imipenem treatment for 14 days, the isolate FK-2820 was found to be carbapenem-resistant. WGS and molecular epidemiology analysis revealed homology among FK-2624, FK-2723 and FK-2820, and the difference in resistance patterns between FK-2624 and the other two strains was due to acquisition of plasmids that conferred certain resistance phenotypes to FK-2723 and FK-2820. All of tested strains harbored the gene \( \text{fosA5} \), which confers high level resistance to fosfomycin [16]. In addition to these core genes, three resistance genes were inferred to be plasmid-borne: \( \text{bla}_{\text{DHA-1}}, \text{qnrB4} \) and \( \text{aac (6')-Ic} \), and the coexistence of multiple resistant genes in these two isolates may contribute to high-level resistance to the majority of clinically available antimicrobial agents.

Moreover, \( K. \ pneumoniae \) strains could produce different virulence factors, such as fimbrial adhesins and siderophores, which are important in the colonization and development of the infection. To date, the delineation of hypervirulent \( K. \ pneumoniae \) (hvKp) virulence genes remains incomplete, and it remains unclear which genes are needed for maximal virulence [17]. We hold that the good markers should be critical factors in conferring the hypervirulent phenotype. If such markers are lost, then the phenotype will no longer be hypervirulent. A recent study demonstrated that \( \text{iroB}, \text{iucA}, \text{peg-344}, \text{rmpA} \) and \( \text{rmpA2} \) were the most accurate molecular markers for defining hvKp [18]. In current study, none of the three strains carried the hypervirulent markers, but the virulence genes \( \text{ybtA}, \text{mrkD}, \text{entB} \) and \( \text{iroN} \) in the MDR background of these \( K. \ pneumoniae \) isolates may further exacerbate infections caused by these bacteria and hamper treatment. \( K. \ pneumoniae \) ST660 remains uncommon in clinical infections, with only two ST660 strains in the publicly accessible database (http://bigd.db.pasteur.fr/). The emergence of this further case warrants further surveillance for ST660.

Our sequencing results showed the acquisition of mutations leading to a reading-frame change and early termination of translation in the \( \text{ompK36} \) coding sequence. The truncated porins were not detected, which is expected given that only a complete polypeptide can form the mature \( \beta \)-barrel structure [9]. Our study provided interesting data on the mRNA responses of \( K. \ pneumoniae \) to carbapenem antibiotics. Expression of \( \text{ompK36} \) was substantially decreased in FK-2820, which was consistent with the absence of the corresponding protein band in the SDS-PAGE analysis. However, the difference between FK-2624 and FK-2723 expression of \( \text{ompK36} \) were not reflected in the protein steady-state levels. In other words, transcriptional responses in \( K. \ pneumoniae \) were not predictive for protein steady-state levels. Relatively little is known about OMP regulation in \( Klebsiella \), but expression of \( \text{ompK36} \) may be subject to negative regulation through factors such as \( \text{micF} \) and \( \text{ompK} \) [19]. These, in turn, could impact on susceptibility to carbapenems and the regulatory cascades that might buffer these effects would be worthy of future investigation. Our results demonstrated the
alteration of OmpK36 alone may confer a high level of resistance to ertapenem, consistent with previous report [20]. Thus, further studies are needed to understand the pathways of regulation for resistance-related genes in K. pneumoniae during antibiotic pressure. A recent study revealed that a 13-day in vitro carbapenem resistance induction showed loss of OmpK 36 in mutant isolate [21]. In current study, our data showed that a 14-day treatment in vivo of MDR K. pneumoniae infection was not particularly effective in clearing MDR K. pneumoniae from the sputum. Furthermore, treatment with imipenem could lead to a premature stop codon of the ompK36 gene in the carbapenem-susceptible MDR K. pneumoniae, leading to the development of carbapenem resistance. The report first described the remodeling of outer membrane porin from carbapenem-susceptible MDR K. pneumoniae to carbapenem-resistant K. pneumoniae from a genomic perspective in vivo. This will give more inspiration to clinical anti-infective treatment. To our knowledge, this study in China is the first description of a connection between imipenem treatment and alteration of OmpK36 in K. pneumoniae.

Conclusions
This study manifested a link between imipenem treatment of patients infected by MDR K. pneumoniae, and the sequential alteration of porin expression in isolates of K. pneumoniae collected during the 14 days treatment. The evidences the need for rational use of antibiotics during patient admission, for the understanding of the impact of treatment-mediated changes to drug-resistance phenotypes, and for effective surveillance to provide guidance on the utility of antimicrobial agents to treat K. pneumoniae infections.

Methods
Bacterial isolates
Between October 2015 to February 2016, K. pneumoniae strains FK2624, FK-2723 and FK-2820 were isolated from the sputum samples of a patient in the First Affiliated Hospital in Wenzhou, China. FK-2624 was the first strain of K. pneumoniae isolated from the patient. After that, multiple swabs for bacterial isolation were taken successively. FK-2624 was chosen to conduct this study as it represented the drug resistance profiles for five of the early isolates (Table S1). Thereafter, FK-2723 was the carbapenem-susceptible strain isolated from the patient immediately before carbapenem treatment was enacted. Subsequently, and after a 14-day treatment with the carbapenem, imipenem, FK-2820 was isolated. Species identification was performed using the Matrix-Assisted Laser Desorption Ionization Time-Of-Flight Mass Spectrometry (MALDITOF MS, Bruker Daltonics, US). After collection, isolates were stored in 30% glycerol at −80°C. All of the investigation protocols in this study were approved by the Ethics Committee of the First Affiliated Hospital of Wenzhou Medical University (ethical number 2019–75). Informed consent was waived because this retrospective study focused on the bacterial isolates and had no impact on interventions to the patient.

Antimicrobial susceptibility testing, detection of resistance genes and virulence factors
The modified carbapenem inactivation method (mCIM) and EDTA synergy test was carried out according to CLSI recommendations for phenotypic screening of carbapenemase producers [22, 23]. Antimicrobial susceptibility test was determined by the agar dilution method and interpreted according to the recommendations in the latest CLSI guidelines, including ceftazidime (0.015–64 g/mL), cefotaxime (0.015–64 g/mL), imipenem (0.015–16 g/mL), meropenem (0.015–16 g/mL), ertapenem (0.0035–16 g/mL), levofloxacin (0.0035–16 g/mL), ciprofloxacin (0.0035–16 g/mL), tobramycin (0.125–256 g/mL), gentamicin (0.125–256 g/mL), amikacin (0.25–128 g/mL) and fosfomycin (0.25–1024 g/mL). The minimum inhibitory concentrations (MICs) of colistin (0.015–16 g/mL) and tigecycline (0.125–16 g/mL) were determined by the broth microdilution method. The latest EUCAST breakpoints (available at http://www.eucast.org/clinical-breakpoints/) were used for colistin and tigecycline. All of the antibiotics were purchased from Becton Dickinson (Sparks, MD). E. coli ATCC 25922 was used as the quality control strain.

A panel of various, relevant antimicrobial resistance genes were amplified by PCR, and the positive amplicons were further confirmed by DNA sequencing. These resistance genes included those encoding the extended spectrum β-lactamase genes (blaCTX-M, blaper, blashV, blaTEM and blavEB) [24], carbapenemase genes (blages, blami/NCM-A, blasm, blakPC, blavIM, blalMP, blanDM and blaoXA-48) [25, 26], AmpC β-lactamase genes (blacMY, blashOXY, blasMOX, and bladHA) [27], plasmid-mediated quinolone resistance (PMQR) genes (qnrA, qnrB, qnrC, qnrD, qnrS, aac (6′)-Ib-cr) [28], qepAB multidrug efflux pump genes [29] and the aminoglycoside resistant genes (aac(6′)-Ib, aac(3′)-Ib, rmtB, armA, APH, ANT, rmtA, rmtB rmtC and rmtD) [30–32]. DNA sequences were characterized by sequence comparisons using BLAST (http://blast.ncbi.nlm.nih.gov/Blast.cgi) [33].

To check for the presence of genes that associated with virulence in K. pneumoniae [34–36], virulence profiles were established by PCR, namely capsular serotype K1, K2, K5, K20, K54, K57, hypermucoviscosity
phenotype (magA), allantoin metabolism (allS), regulator of mucoid phenotype A (rmpA), iron system capture (iroN), adhesion type 3 fimbriae (mrkD), iron transport and phosphotransferase function (kfu), siderophore (entB) and siderophore yersiniabactin (ybtA).

Whole-genome sequencing (WGS)
Genomic DNA of K. pneumoniae FK-2624, FK-2723 and FK-2820 were purified using the Bioflux DNA purification kit (Bioflux BSC12S1, Beijing) as recommended by the manufacturer. A total amount of 1 μg DNA per sample was used as input material for the DNA sample preparations. Sequencing libraries were generated using NEBNext Ultra™ DNA Library Prep Kit for Illumina (NEB E7645S, USA) following manufacturer’s recommendations and index codes were added to attribute sequences to each sample. At last, PCR products were purified (AMPure XP system A63880, Beckman, USA) and libraries were analyzed for size distribution by Agilent2100 Bioanalyzer and quantified using real-time PCR. The genomes were sequenced by Illumina NovaSeq PE150.

Sequence reads for each isolate were assembled individually. All good quality paired reads were assembled using the SOAP denovo [37] into a number of contigs. After assembly, the contigs were scaffolded using MeDuSa [38] and ordered of scaffolds was done using MAUVE version 0.25 [39] using as a reference a complete close genome detected using MagiBlast. Annotated was performed using the National Center for Biotechnology Information’s Prokaryotic Genome Annotation Pipeline [40]. To identify potential antibiotic resistance genes in the genomic sequence of the isolates, sequence alignment of the protein sequences of antibiotic resistance genes in the Center for Genomic Epidemiology (http://www.genomicepidemiology.org/). Virulence genes were identified using the virulence factor database (http://www.mgc.ac.cn/VFs/) and Pathogen-Finder (https://cge.cbs.dtu.dk/services/PathogenFinder/).

Additionally, Kleborate (version 0.3.0) was used to identify resistant and virulent genes with their corresponding subtypes [41, 42]. The nucleotide sequence of the genome of FK-2624, FK-2723 and FK-2820 have been submitted to GenBank with accession no. VIGL000000000, VIGM000000000 and VIGK000000000, respectively. Alignments with the three assembled genomes were done using MAUVE version 0.25 [39]. Sequences that were different between isolates were tested for different GC content, using window-aact form GLIMMER v3.02 [43]. Putative plasmids were assembled and the resulting plasmid assemblies were annotated with Prokka (v1.12) [44]. Graphical representation was generated using the R packages.

Roary (version 3.12.0) was used to align the assembled genomes and two additional ST660 MLST genomes found in NCBI (GCA_002853165 and GCA_002205995.1) and extract the core genome [45]. The core genome was used to generate a phylogenetical tree using RaxML (version 8.2.9), with a GTR + GAMMA model, and a bootstrapping of 1000 replicates [46].

MLST determination of genetic relatedness of isolates
Pairwise comparisons method, pulsed-field gel electrophoresis (PFGE) and multi-locus sequencing typing (MLST) were used to establish relatedness between same-patient isolates.

The sequences of the three complete genomes of K. pneumoniae isolates were independently used in sequence alignment. The sequence reads were mapped to the reference genomes using the Bowtie2 software, which is good for mapping short sequence reads to medium-sized and large genomes. The alignment of clean data of 3 isolates with reference K. pneumoniae MGH 78578 (MDR bacterium isolated from a patient [47], accession number CP000647) was performed with the default settings of programs. Finally, the alignment percentage (supplementary material Table S2) was showed by Bowtie2, revealed a high degree of genetic conservation was observed between the three K. pneumoniae strains.

PFGE was carried out on our strains according to the method described previously with minor modification [48]. Genomic DNA was extracted from the K. pneumoniae isolates, followed by restriction enzyme Xba I (Takara 1093A, Japan) digestion for 2 h. PFGE was performed using a CHEF-Mapper XA PFGE system (Bio-Rad, USA) for 18 h with a switch time 6–36 s. Then DNA fingerprints were revealed by GelRed staining. The banding patterns were visualized by GelDoc XR gel imaging system (Bio-Rad, USA) and cluster analysis of similarity values of the PFGE profiles were finally performed by Quantity One program (Bio-Rad Laboratories, USA). The sequence types (STs) of FK-2624, FK-2723 and FK-2820 were determined by MLST. Seven housekeeping genes (gapA, infB, mdh, pgI, phoE, rpoB and tonB) were amplified and sequenced according to Diancourt et al. [49]. Alleles and sequence types were assigned by the MLST database (http://www.pasteur.fr/mlst/Kpneumoniae.html).

Outer membrane protein isolation and SDS-PAGE
FK-2624, FK-2723 and FK-2820 were cultured in Muller Hinton broth with shaking overnight at 37 °C. Cells were isolated by centrifugation (4200 rpm for 15 min), the cell pellets were washed with 10 mM Tris-HCl, 5 mM MgCl2 (pH 7.3), and the cells then lysed by sonication as described [50]. The cell lysate supernatants were treated
with 2% solution of sodium lauroylsarcosinate for 30 min at room temperature, centrifuged 30 min at 17000 rpm and the pellets containing outer membrane proteins (OMPs) were suspended in 10 mM Tris-HCl, 5 mM MgCl2 (pH 7.3).

Samples containing the OMPs were mixed with loading buffer (TaKaRa 9173, Japan) and denatured by heating at 100 °C for 3 min. Proteins were then separated by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) using 12% polyacrylamide separation gel. Bands corresponding to the major protein species were visualized by staining the gels with 0.2% Coomassie brilliant blue (Solarbio C8430, China) in 10% acetic acid and 45% methanol. *K. pneumoniae* ATCC 13883 (an isolate with known expression of OmpK35 and OmpK36) served as a control strain for OMP profiling.

### Analysis of ompK35 and ompk36 genes

The coding sequences of the *ompK35* and *ompK36* genes for the representative isolates were amplified and sequenced using primers listed in Table 2. Amplification was carried out with the following thermal cycling conditions: 5 min at 94 °C and 35 cycles of amplification consisting of 30 s at 94 °C, 30 s at 58 °C, and 1 min at 72 °C, with 10 min at 72 °C for the final extension. DNA fragments were analyzed by electrophoresis in a 1% agarose gel at 120 V for 20 min, stained with GelRed (Biotium 41,003, USA). DNA sequence analysis compared with those of *K. pneumoniae* NTUH-2044 (NCBI accession number AP 006725) was conducted.

Isolates were examined by real-time RT-PCR to measure the steady-state levels of the transcripts as an indication of the expression of *ompK35* and *ompK36*. For RNA isolation, *K. pneumoniae* isolates were grown in fresh LB medium at 37 °C overnight. Total RNA was extracted from 3 mL of culture using a RNeasy Mini Kit (Qiagen 74, 524, Germany) according to the manufacturer's instructions. The extracted RNA samples were stored at −80 °C. Purified RNA was reverse transcribed into cDNA for RT-PCR analysis using a cDNA synthesis kit (TaKaRa 6210A, Japan) according to the manufacturer’s instructions. Gene expression levels were measured with RT-PCR using TB Green® Premix Ex Taq™ (TaKaRa RR420Q, Japan). Primers were designed on the basis of the nucleotide sequence in GenBank (Table 2). Expression of each gene was normalized to that of a housekeeping gene (*rpoB*), the primers used for RT-PCR were *rpoB*-F (5′-AAGGCGAATC CAGCTTGTTTCAGC-3′) and *rpoB*-R (5′-TGACGTTG CAGTTTCGCACCCCATCA-3′). The relative expression of *ompK35* and *ompk36* was then calibrated against the corresponding expression by *K. pneumoniae* ATCC 13883. For quality-controlling strain, relative expression was equal to 1.0. All experiments were performed in triplicate and mean of the Ct values were used for analysis. Gene expression levels were calculated using 2−ΔΔCt method [51].

### Statistical analysis

The statistical correlation of the expression of *ompk35* and *ompk36* was calculated by Student’s *t* test. SPSS (version 17; IBM, USA, IL) was used for statistical analysis. A *P* value lower than 0.05 was considered statistically significant.

### Supplementary information

Supplementary information accompanies this paper at https://doi.org/10.1186/s12866-020-01898-1.

### Additional file 1: Table S1.

*Klebsiella pneumoniae* isolated during hospitalization.

### Additional file 2: Table S2.

Genomic DNA comparison between the tested strains, SNPs and (% identity).

### Additional file 3: Table S3.

MICs of *Pseudomonas aeruginosa*.

### Additional file 4: Table S4.

The allelic profile of ST 660.

### Additional file 5: Table S5.

Genomic DNA assigned to plasmids.

### Additional file 6: Figure S1.

 SDS-PAGE analysis of OMPs from representative strains. 

### Additional file 7: Figure S2.

Pulsed-field gel electrophoresis analysis. 

### Abbreviations

MDR: Multidrug-resistant; MALDI-TOF MS: Matrix-assisted laser desorption/ionization time of flight mass spectrometry; MIC: Minimum inhibitory concentration; PCR: Polymerase chain reaction; WGS: Whole-genome sequencing; SDS-PAGE: Sodium dodecyl sulfate polyacrylamide gel electrophoresis; ESBL: Extended-spectrum β-Lactamases; CRKP: Carbapenem-resistant *K. pneumoniae*; mCM: Modified carbapenem inactivation method; PFGE: Pulsed-field gel electrophoresis; MLST: Multi-locus sequencing typing; qRT-PCR: Quantitative reverse transcription polymerase chain reaction.
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Authors' contributions
TZ, JC, and XT contributed to the design of the experiments. XT, XZ, YZ, RF, LP-I and CW performed the experiment. XT, XZ, and RF wrote the initial draft of the manuscript. TZ, JC, XT, GW, JL, YZ, LP-I, JD, TL and HL contributed to the acquisition, analysis, interpretation of the data included in this manuscript. TZ, JC, XT, RF, JL, CW, TL and HL revised the manuscript. All authors approve of the final manuscript being submitted and agree to be accountable for the work detailed in the submitted manuscript.

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Availability of data and materials
The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The work was approved by the Ethics Committee of the First Affiliated Hospital of Wenzhou Medical University. The committee ruled that consent to participate was waived (ethical number 2019-Hospital of Wenzhou Medical University). The committee ruled that consent to participate was waived (ethical number 2019–Hospital of Wenzhou Medical University).

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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