

Trends in DTR, CR, ECR, and FQR in Four Common Gram-Negative Bacteria: A Retrospective Study from 2013 to 2021

Zhen Zhang¹, Lei Tian²

¹Department of Pharmacy, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei Province, People's Republic of China; ²Department of Clinical Laboratory, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei Province, People's Republic of China

Correspondence: Lei Tian, Department of Clinical Laboratory, Tongji Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, Hubei Province, People's Republic of China, Email iso15189@126.com

Purpose: The purpose of this study was to examine the prevalence of four important drug-resistance phenotypes: difficult-to-treat resistance (DTR), fluoroquinolone resistance (FQR), carbapenem resistance (CR), and extended-spectrum cephalosporin resistance (ECR).

Methods: DTR was defined as insensitivity to all the β -lactams and fluoroquinolones tested. We retrospectively analyzed the distribution characteristics of specific drug-resistant phenotypes of the main Gram-negative bacteria causing bloodstream infections (BSIs) in Tongji Hospital (Wuhan, China) between 2013 and 2021: *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, and *Acinetobacter baumannii*.

Results: FQR was the main antimicrobial resistance phenotype of *E. coli*, accounting for 59.45% (1117/1879, 95% confidence interval, 57.21%–61.65%); the detection rates for CR and DTR were low, accounting for 1.86% (35/1879, 1.34%–2.58%), and 1.81% (34/1879, 1.30%–2.52%), respectively. However, the detection rates for CR and DTR in *K. pneumoniae* were 38.83% (497/1280, 36.20%–41.53%) and 35.94% (460/1280, 33.35%–38.60%), respectively. In *P. aeruginosa*, the detection rates of the four drug-resistant phenotypes (DTR, CR, FQR, and ECR) were all < 30%, but conversely, for *A. baumannii*, the detection rates were all > 80%. The changes in the data from 2013 to 2021 showed upward trends ($z > 0$) for CR-*E. coli*, DTR-*E. coli*, FQR-*E. coli*, CR-*K. pneumoniae*, DTR-*K. pneumoniae*, FQR-*K. pneumoniae*, and ECR-*K. pneumoniae*, but downward trends ($z < 0$) for ECR-*E. coli*, CR-*A. baumannii*, DTR-*A. baumannii*, FQR-*A. baumannii*, ECR-*A. baumannii*, CR-*P. aeruginosa*, DTR-*P. aeruginosa*, FQR-*P. aeruginosa*, and ECR-*P. aeruginosa*.

Conclusion: DTR warrants further attention, especially in BSI-associated *K. pneumoniae* and *A. baumannii*, in which the detection rates were very high. Between 2013 and 2021 in this region, DTR-*E. coli* and CR-*E. coli* showed obvious upward trends, whereas DTR-*P. aeruginosa* and ECR-*P. aeruginosa* showed obvious downward trends.

Keywords: DTR, CR, ECR, FQR, BSI

Introduction

The invention of antibiotics is an important milestone in the development of modern medical science. However, the emergence of antimicrobial resistance (AMR) has seriously threatened human health. It is reported that about 23,000 and 25,000 people die each year from infections complicated by AMR in the United States and Europe, respectively.¹ Consequently, AMR is a very noteworthy focus of research throughout the world.

To better monitor AMR, in 2008, the European Centre for Disease Prevention and Control (ECDC) and the USA Centers for Disease Control and Prevention (CDC) jointly launched the concepts of multidrug resistance (MDR), extensive drug resistance (XDR), and pandrug resistance (PDR) to describe different drug resistance phenotypes. MDR was defined as simultaneous insensitivity to three types of antibiotics. XDR was defined as sensitivity to only one or two antibiotics. PDR was

defined as nonsusceptibility to all antibiotics tested.² Although these concepts have some epidemiological value, several experts suggest that they ignore the toxicity and efficiency of antibiotics, and that their correlations with clinical outcomes are questionable.³ In 2018, Kadri et al proposed a new concept with which to monitor AMR among Gram-negative bacteria, difficult-to-treat resistance (DTR), which was defined as intermediate or resistance to all first-line agents, including β -lactams and fluoroquinolones.^{4,5} However, there have been very few reports of DTR around the world outside the United States and South Korea. Therefore, we retrospectively analyzed the distribution of DTR, CR, ECR, and FQR in four common Gram-negative bacteria causing bloodstream infections (BSIs), *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, and *Acinetobacter baumannii*, in our hospital between 2013 and 2021, to provide some epidemiological references for decision making about clinical treatment and infection control.

Methods

Data Source

All pathogenic microorganisms were isolated from blood-culture-positive samples at Tongji Hospital (Wuhan, China) between 2013 and 2021. In this retrospective study, the four commonest Gram-negative bacteria causing BSIs at our center, *E. coli*, *K. pneumoniae*, *A. baumannii*, and *P. aeruginosa*, were analyzed. The identification and antimicrobial sensitivity testing of the strains were performed in the microbiology laboratory of Tongji Hospital. Automated blood culture instruments, including the BD BACTEC™ FX 400 (BD Co., NJ, USA) or the BacT/ALERT® 3D Microbial Detection System (bioMérieux, Lyon, France), were used in the microbiology laboratory. The strains were identified with an automatic identification system (VITEK® 2 Compact, bioMérieux) and/or MALDI Biotyper® sirius IVD System (Bruker, Antu, China). Antimicrobial susceptibility testing was performed in the final year with the disk diffusion method according to the Clinical and Laboratory Standards Institute (CLSI) guidelines. Resistance to carbapenems, including imipenem and meropenem, was screened with the disk diffusion method and confirmed with the E-test method.

Definition of Specific Antimicrobial Resistance Phenotypes

In this study, DTR was defined as resistance or intermediate resistance in vitro to all β -lactam categories, including carbapenems and fluoroquinolones.^{4,5} Carbapenem resistance (CR) was defined as resistance in vitro to imipenem and meropenem. Extended-spectrum cephalosporin resistance (ECR) was defined as resistance in vitro to ceftazidime, cefotaxime, and cefepime. Fluoroquinolone resistance (FQR) was defined as resistance in vitro to ciprofloxacin and levofloxacin.⁶

Statistical Analysis

Data from the drug sensitivity experiments were analyzed using the WHONET 5.6 software. When multiple specimens were collected from the same patient simultaneously, only the first isolate of a given species from that patient was analyzed according to CLSI M-39.⁷ The interpretation criteria for the antimicrobial susceptibility results were based on the CLSI 2020 guidelines.⁸ The changing trends in the detection rates of specific drug resistance phenotypes were analyzed with the Mann–Kendall test, with the R 4.0.2 program (R Core Team). $z < 0$ indicated a downward trend, whereas $z > 0$ indicated an upward trend. $P < 0.05$ indicated statistical significance.

Ethics Approval and Consent to Participate

The study protocol was approved by the Tongji Hospital Ethics Committee for Research in Health. The committee also approved a waiver for informed consent to participate in this study given its retrospective design. All patient data were anonymized before analysis.

Results

Detection Rates of Specific Antimicrobial Resistance Phenotypes

For *E. coli*, FQR was the main drug resistance phenotype, accounting for 59.45% (1117/1879, 95% confidence interval, 57.21%–61.65%) of phenotypes detected, followed by ECR, accounting for 31.08% (584/1879, 29.03%–33.21%), whereas CR and DTR accounted for 1.86% (35/1879, 1.34%–2.58%) and 1.81% (34/1879, 1.30%–2.52%) respectively. For *K. pneumoniae*, the

detection rates for ECR, FQR, CR, and DTR were 50% (640/1280, 47.26%–52.74%), 48.13% (616/1280, 45.40%–50.86%), 38.83% (497/1280, 36.20%–41.53%), and 35.94% (460/1280, 33.35%–38.60%), respectively. For *P. aeruginosa*, the detection rates for ECR, FQR, CR and DTR were all < 30%, at 16.76% (59/352, 13.22%–21.02%), 18.18% (64/352, 14.50%–22.55%), 26.99% (95/352, 22.62%–31.86%), and 11.36% (40/352, 8.46%–15.10%), respectively. For *A. baumannii*, the detection rates for ECR, FQR, CR, and DTR were all > 80%, at 85.52% (384/449, 81.97%–88.48%), 83.96% (377/449, 80.28%–87.07%), 84.41% (379/449, 80.76%–87.47%), and 83.30% (374/449, 79.57%–86.46%), respectively.

Trends in Specific Drug Resistance Phenotypes

In *E. coli*, the detection rate of FQR ranged from 50% to 70% from 2013 to 2021; the detection rate for ECR ranged from 20% to 40%; and the detection rates for CR and DTR were < 5% (Figure 1A). In *K. pneumoniae*, DTR, CR, ECR, and FQR increased significantly after 2015. The detection rate for DTR was < 3% in 2013 and 2014, but increased from 25% to 50% between 2015 and 2021. The detection rate for CR was < 10% in 2013 and 2014, but increased from 30% to 55% after 2015. The detection rate for FQR was < 15% in 2013 and 2014, but 40%–70% after 2015 (Figure 1B). In *A. baumannii*, the detection rates for DTR, CR, ECR, and FQR were all > 70% from 2013 to 2021 (Figure 1C). However, the detection rates for the four drug-resistant phenotypes of *P. aeruginosa* were ≤40%, and showed obvious downward trends after 2019 (Figure 1D). Statistical analysis with the Mann–Kendall test indicated that CR-*E. coli* and DTR-*E. coli*

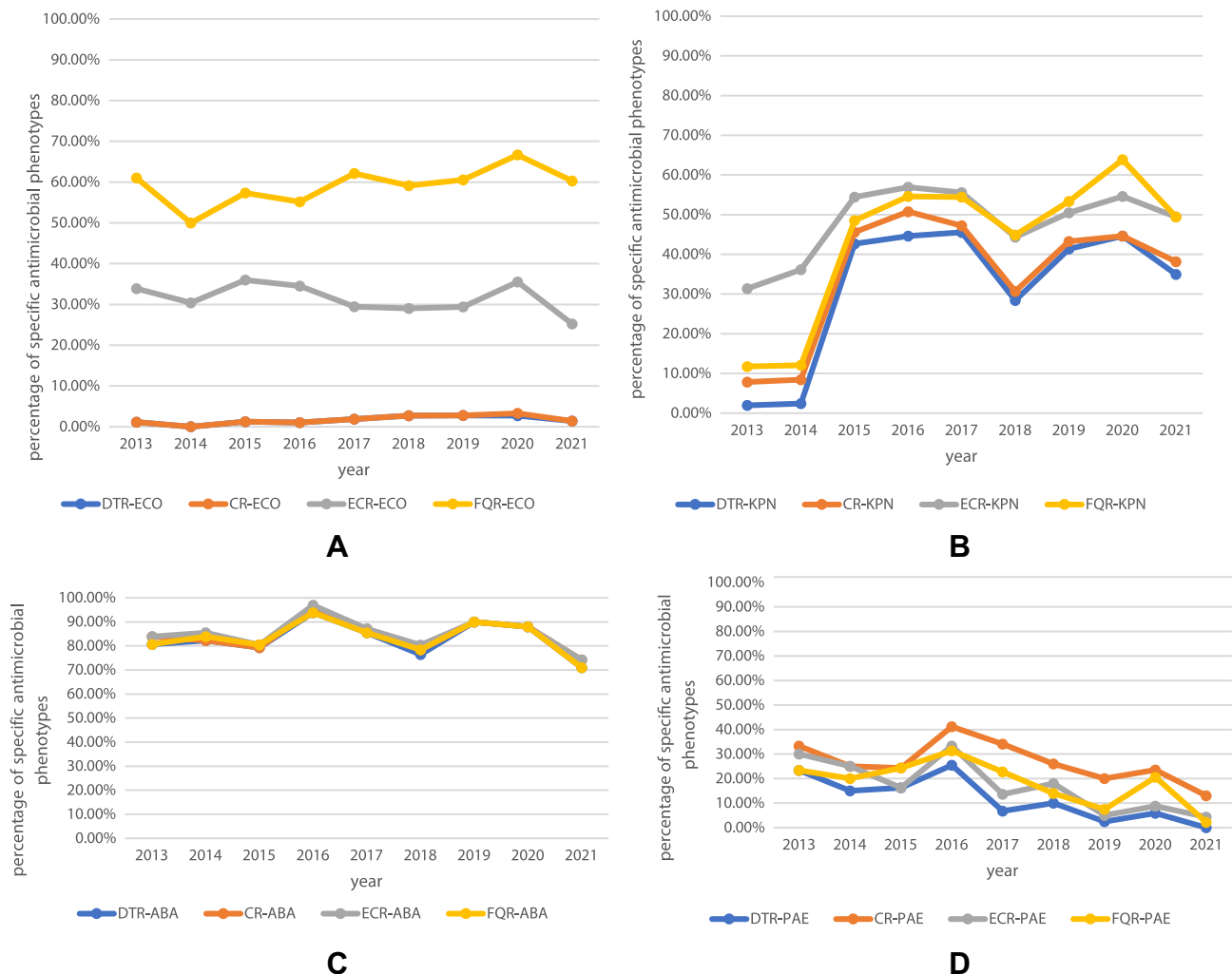


Figure 1 Trends of special antimicrobial resistance phenotypes from 2013 to 2021. ((A), *E. coli*, (B), *K. pneumoniae*, (C), *A. baumannii*, (D), *P. aeruginosa*).
Abbreviations: ECO, *E. coli*; KPN, *K. pneumoniae*; PAE, *P. aeruginosa*; ABA, *A. baumannii*.

Table 1 Results of Statistical Analysis on the Trend of Specific Drug Resistance Phenotypes

Specific Drug Resistance Phenotypes	Z	P	P<0.05
CR-ECO	2.1894	0.02857	+
DTR-ECO	1.9809	0.0476	+
ECR-ECO	-1.3553	0.1753	-
FQR-ECO	1.1468	0.2515	-
CR-KPN	0.52129	0.6022	-
DTR-KPN	1.0483	0.2945	-
ECR-KPN	0.7298	0.4655	-
FQR-KPN	1.5639	0.1179	-
CR-ABA	-0.31277	0.7545	-
DTR-ABA	-0.10426	0.917	-
ECR-ABA	-0.10426	0.917	-
FQR-ABA	-0.10426	0.917	-
CR-PAE	-1.7724	0.07633	-
DTR-PAE	-2.3979	0.01649	+
ECR-PAE	-2.3979	0.01649	+
FQR-PAE	-1.7724	0.07633	-

Abbreviations: DTR, Difficult to treat resistance; FQR, Fluoroquinolone resistance; CR, carbapenem resistance; ECR, Extended-spectrum cephalosporin resistant; ECO, *E. coli*; KPN, *K. pneumoniae*; ABA, *A. baumannii*; PAE, *P. aeruginosa*.

showed significant upward trends ($z > 0$, $P < 0.05$), whereas DTR-*P. aeruginosa* and ECR-*P. aeruginosa* showed significant downward trends ($z < 0$, $P < 0.05$; Table 1).

Discussion

Although MDR, XDR, and PDR are very important concepts in monitoring drug resistance, these classifications do not consider the toxicity or therapeutic efficacy of antimicrobial drugs themselves, and judging whether a strain is XDR or PDR requires that a very wide variety of antimicrobial drugs be tested. The new definition, DTR, focuses on the effect of antimicrobial resistance on treatment decisions and clinical outcomes.⁵ DTR is a very practical value indicator, and the emergence of DTR indicates that when deciding a clinical treatment, some antibacterial drugs that are less effective and/or more toxic should be considered. DTR is also a public health threat and contributes to the international spread of plasmids, antibiotic overuse, and the need for costly infection control measures in healthcare facilities.⁴

In this study, we found that DTR was not common in *E. coli*, but could not be ignored in *A. baumannii* and *K. pneumoniae*. According to the epidemic trends in 2013–2021, the specific drug resistance phenotypes of DTR and CR have increased significantly in *E. coli*, whereas DTR and ECR have clearly decreased in *P. aeruginosa*.

In our monitoring data, FQR was the most frequent phenotype in *E. coli*, with a prevalence rate of 59.45%, followed by ECR, with a prevalence rate of 31.08%. The drug resistance of *E. coli* is consistent with the monitoring data reported in the United States and South Korea, but the prevalence of FQR is significantly higher than that reported in the United States and South Korea (27.3% and 44.1% respectively).^{4,5} The prevalence rate of ECR was 31.08%, which is similar the 33.1% reported in South Korea, but significantly higher than the 6.4% reported in the United States.^{4,5} The prevalence rates of *E. coli*-DTR and *E. coli*-CR in our study are consistent with those in the United States (0.04% and 0.1%, respectively) and South Korea (1.5% and 0%, respectively).^{4,5} The resistance rates of *E. coli* to fluoroquinolones and cephalosporins in our region were higher than those reported in the United States and South Korea, which may be attributable to the extensive use of fluoroquinolones and cephalosporins in our region. Research at a teaching hospital in China showed that from 2010 to 2016, the outpatient prescription rate for fluoroquinolones decreased from 19.38% to 13.21%, and that for inpatients decreased from 64.34% to 34.65%.⁹ Between 2011 and 2016, the resistance rate of *E. coli* to levofloxacin decreased from 61.3% to 53.9% and that to ciprofloxacin decreased from 64.3% to 58.2%.⁹ Therefore, the reduction in *E. coli* resistance to fluoroquinolones correlated positively with the reduced use of antimicrobial agents. Analogously, the high resistance of *E. coli* to carbapenems may be related to the overuse of these antimicrobials. A study

of the effects of an antimicrobial restriction system to curb the inappropriate use of carbapenem indicated that antimicrobial restriction systems can increase the appropriate prescription of carbapenem antibiotics and effectively reduce the production of carbapenem-resistant bacterial strains.¹⁰

In the present study, the prevalence rates of FQR, ECR, CR, and DTR in *K. pneumoniae* were similar, at 48.13%, 50.00%, 38.83%, and 35.94%, respectively, but were significantly higher than those in the United States (9.7%, 8.9%, 2.1%, and 1.7%, respectively) and South Korea (19.2%, 21.2%, 2.1%, and 1.4%, respectively).^{4,5} Carbapenemases, extended-spectrum β -lactamases, and AmpC β -lactamases are the main mechanisms of *K. pneumoniae* resistance to a variety of antibiotics. The high prevalence of CR is mainly responsible for the high prevalence of DTR. CR is prevalent in many countries, but the resistance mechanism of CR differs in different regions. In China, *bla*_{KPC-2} is the main carbapenemase producer, whereas *bla*_{KPC-16} is most widely distributed in Japan and *bla*_{KPC-15} in Singapore.¹¹ IMP-11- and IMP-6- producing CR-*K. pneumoniae* are most commonly reported in Japan.¹² *bla*_{OX-48} is mainly distributed in Turkey, but was recently also reported in China, the United States, and several European countries (Spain, France, Germany, and Switzerland).^{11,12} In a previous study of the CR genotypes in our hospital, *bla*_{kpc-2} mainly colonized the infected group of patients, accounting for 72.4% of infections, whereas *bla*_{NDM-1} mainly colonized the noninfected patient group, accounting for 42.2% of infections.¹³ The high prevalence of CR-*K. pneumoniae* in our hospital may be related to the simultaneous expression of multidrug-resistance genes. The mechanism of resistance to carbapenems in *K. pneumoniae* in this region remains to be clarified.

In this study, the prevalence rates for FQR, ECR, CR, and DTR in *A. baumannii* were all > 80%, and were significantly higher than those reported in the United States (49.5%, 55.4%, 29.4%, and 18.3%, respectively) and South Korea (65.5%, 70.0%, 63.4%, and 59.4%, respectively).^{4,5} However, the drug resistance of *P. aeruginosa* showed a gradual downward trend. The detection rates of FQR, ECR, CR, and DTR were 18.18%, 16.76%, 26.99%, and 11.36%, respectively, whereas the corresponding data reported from the United States were 18.5%, 9.1%, 10.4%, and 2.3%, respectively, and those from South Korea were 33.0%, 28.1%, 28.1%, and 9.0%, respectively.^{4,5} The high drug resistance rates for *A. baumannii* may be related to the simultaneous expression of multiple drug resistance mechanisms, such as a change in the antibiotic target, the expression of an efflux pump, the production of various enzymes that hydrolyze antibiotics, and so on.¹⁴ A study by Wei Feng indicated that the mutational inactivation of OprD was the main factor responsible for CR-*P. aeruginosa*.¹⁵ The molecular mechanisms of drug resistance in *A. baumannii* and *P. aeruginosa* in this region warrant further research.

Clinical experts have been investigating new antimicrobial agents for the treatment of CR-*A. baumannii* infections. A study by Wasan Katip indicated that a nonloading dose of colistin methane sulfonate increased the opportunity to achieve a favorable outcome, but also increased nephrotoxicity in patients.¹⁶ In another study, Wasan Katip demonstrated that the addition of meropenem to colistin reduced the 30-day mortality rate, achieved greater clinical and microbiological responses, and did not increase the nephrotoxicity compared with colistin monotherapy.¹⁷ However, colistin plus vancomycin showed no significant differences in the 30-day mortality, clinical response, or microbiological response compared with colistin alone in treating infections of CR-*A. baumannii*.¹⁸

The present study had several limitations. First, the use of a study design in which a tertiary hospital was used as the reference, rather than a geographical design that included all bacterial data from a given population, greatly limited the representativeness of the data. The fact that no information was available on the number of blood cultures taken per year during the study period, on the positivity rates of the blood cultures, or on the patient population of the hospital also limits the generalizability of our findings. Second, the mechanisms of resistance were not investigated in this study, including the resistance to cephalosporins, fluoroquinolones, and carbapenems. We look forward to examining these mechanisms of drug resistance in subsequent studies.

Conclusions

The drug resistance monitoring data from a single center from 2013 to 2021 showed that CR-*E. coli*, DTR-*E. coli*, FQR-*E. coli*, CR-*K. pneumoniae*, DTR-*K. pneumoniae*, ECR-*K. pneumoniae*, and FQR-*K. pneumoniae* showed upward trends, whereas ECR-*E. coli*, CR-*A. baumannii*, DTR-*A. baumannii*, ECR-*A. baumannii*, FQR-*A. baumannii*, DTR-*P. aeruginosa*, CR-*P. aeruginosa*, ECR-*P. aeruginosa*, and FQR-*P. aeruginosa* showed downward trends. Among these trends, the

upward trends in CR-*E. coli* and DTR-*E. coli* and the downward trends in DTR-*P. aeruginosa* and ECR-*P. aeruginosa* were significant.

Abbreviations

AMR, antimicrobial resistance; BSI, Bacterial bloodstream infection; MDR, Multidrug resistance; XDR, extensive drug resistance; PDR, pandrug-resistance; CDC, Center for Disease Control and Prevention; ECDC, European Centre for Disease Prevention and Control; DTR, Difficult to treat resistance; FQR, Fluoroquinolone resistance; CR, carbapenem resistance; ECR, Extended-spectrum cephalosporin resistant.

Funding

There is no funding to report.

Disclosure

The authors report no conflicts of interest in this work.

References

1. Marston HD, Dixon DM, Knisely JM, Palmore TN, Fauci AS. Antimicrobial Resistance. *JAMA*. 2016;316(11):1193–1204. doi:10.1001/jama.2016.11764
2. Magiorakos AP, Srinivasan A, Carey RB, et al. Multidrug-resistant, extensively drug-resistant and pandrug-resistant bacteria: an international expert proposal for interim standard definitions for acquired resistance. *Clin Microbiol Infect*. 2012;18(3):268–281. doi:10.1111/j.1469-0691.2011.03570.x
3. Bassetti M, Vena A, Sepulcri C, Giacobbe DR, Peghin M. Treatment of Bloodstream Infections Due to Gram-Negative Bacteria with Difficult-to-Treat Resistance. *Antibiotics*. 2020;9(9):314. doi:10.3390/antibiotics9090632
4. Kadri SS, Adjemian J, Lai YL, et al. Difficult-to-Treat Resistance in Gram-negative Bacteremia at 173 US Hospitals: retrospective Cohort Analysis of Prevalence, Predictors, and Outcome of Resistance to All First-line Agents. *Clin Infect Dis*. 2018;67(12):1803–1814. doi:10.1093/cid/ciy378
5. Huh K, Chung DR, Ha YE, et al. Impact of Difficult-to-Treat Resistance in Gram-negative Bacteremia on Mortality: retrospective Analysis of Nationwide Surveillance Data. *Clin Infect Dis*. 2020;71(9):e487–e96. doi:10.1093/cid/ciaa084
6. Zhang Z, Sun Z, Tian L. Antimicrobial Resistance Among Pathogens Causing Bloodstream Infections: a Multicenter Surveillance Report Over 20 Years (1998–2017). *Infect Drug Resist*. 2022;15:249–260. doi:10.2147/IDR.S344875
7. Clinical and Laboratory Standards Institute. *Analysis and Presentation of Cumulative Antimicrobial Susceptibility Test Data, M39-A4*. Wayne, PA: Clin Lab Stand Institute; 2014.
8. Clinical and Laboratory Standards Institute. *Performance Standards for Antimicrobial Susceptibility Testing, Thirtieth Informational Supplement, M100-S30*. Wayne, PA: Clin Lab Stand Institute; 2020.
9. Wang H, Wang H, Yu X, et al. Impact of antimicrobial stewardship managed by clinical pharmacists on antibiotic use and drug resistance in a Chinese hospital, 2010–2016: a retrospective observational study. *BMJ Open*. 2019;9(8):e026072. doi:10.1136/bmjopen-2018-026072
10. Wanla W, Katip W, Supakul S, et al. Effects of an antimicrobial restriction system on appropriate carbapenem use in a hospital without infectious diseases consultation. *Int J Gen Med*. 2017;10:443–449. doi:10.2147/IJGM.S145133
11. Li J, Huang Z, Tang M, et al. Clonal Dissemination of Multiple Carbapenemase Genes in Carbapenem-Resistant Enterobacterales Mediated by Multiple Plasmids in China. *Infect Drug Resist*. 2021;14:3287–3295. doi:10.2147/IDR.S327273
12. Chen HY, Jean SS, Lee YL, et al. Carbapenem-Resistant Enterobacterales in Long-Term Care Facilities: a Global and Narrative Review. *Front Cell Infect Microbiol*. 2021;11:601968. doi:10.3389/fcimb.2021.601968
13. Lin Q, Wang Y, Yu J, et al. Bacterial characteristics of carbapenem-resistant Enterobacteriaceae (CRE) colonized strains and their correlation with subsequent infection. *BMC Infect Dis*. 2021;21(1):638. doi:10.1186/s12879-021-06315-0
14. Kyriakidis I, Vasileiou E, Pana ZD, Tragiannidis A. Acinetobacter baumannii Antibiotic Resistance Mechanisms. *Pathogens*. 2021;10(3):2154. doi:10.3390/pathogens10030373
15. Feng W, Huang Q, Wang Y, et al. Changes in the resistance and epidemiological characteristics of Pseudomonas aeruginosa during a ten-year period. *J Microbiol Immunol Infect*. 2021;54(2):261–266. doi:10.1016/j.jmii.2019.08.017
16. Katip W, Uitrakul S, Oberdorfer P. Clinical Efficacy and Nephrotoxicity of the Loading Dose Colistin for the Treatment of Carbapenem-Resistant Acinetobacter baumannii in Critically Ill Patients. *Pharmaceutics*. 2021;14(1):365. doi:10.3390/pharmaceutics14010031
17. Katip W, Uitrakul S, Oberdorfer P. A Comparison of Colistin versus Colistin Plus Meropenem for the Treatment of Carbapenem-Resistant Acinetobacter baumannii in Critically Ill Patients: a Propensity Score-Matched Analysis. *Antibiotics*. 2020;9(10):871. doi:10.3390/antibiotics9100647
18. Katip W, Oberdorfer P. Clinical Efficacy and Nephrotoxicity of Colistin Alone versus Colistin Plus Vancomycin in Critically Ill Patients Infected with Carbapenem-Resistant Acinetobacter baumannii: a Propensity Score-Matched Analysis. *Pharmaceutics*. 2021;13(2):87. doi:10.3390/pharmaceutics13020162

Infection and Drug Resistance

Dovepress

Publish your work in this journal

Infection and Drug Resistance is an international, peer-reviewed open-access journal that focuses on the optimal treatment of infection (bacterial, fungal and viral) and the development and institution of preventive strategies to minimize the development and spread of resistance. The journal is specifically concerned with the epidemiology of antibiotic resistance and the mechanisms of resistance development and diffusion in both hospitals and the community. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Submit your manuscript here: <https://www.dovepress.com/infection-and-drug-resistance-journal>