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Retrospective Study

Distribution and drug resistance of pathogenic bacteria in emergency patients

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Informed consent statement:

Patients were not required to give informed consent to the study because the analysis used anonymous clinical data.

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Abstract

BACKGROUND

Antibiotic resistance has become a global threat for human health, calling for rational use of antibiotics.

AIM

To analyze the distribution and drug resistance of the bacteria, providing the prerequisite for use of antibiotics in emergency patients.

METHODS

A total of 2048 emergency patients from 2013 to 2017 were enrolled. Their clinical examination specimens were collected, followed by isolation of bacteria. The bacterial identification and drug susceptibility testing were carried out.

RESULTS

A total of 3387 pathogens were isolated. The top six pathogens were *Acinetobacter baumannii* (660 strains), *Staphylococcus aureus* (436 strains), *Klebsiella pneumoniae* (347 strains), *Pseudomonas aeruginosa* (338 strains), *Escherichia coli* (237 strains), and *Candida albicans* (207 strains). The isolation rates of these pathogens decreased year by year except *Klebsiella pneumoniae*, which increased from 7.1% to 12.1%. *Acinetobacter baumannii* is a widely-resistant strain, with multiple resistances to imipenem, ciprofloxacin, minocycline and tigecycline. The *Staphylococcus aureus* had high resistance rates to levofloxacin, penicillin G, and tetracycline. But the susceptibility of it to vancomycin and tigecycline were 100%. *Klebsiella pneumoniae* had high resistance rates to imipenem, cefoperazone/sulbactam, amikacin, and ciprofloxacin, with the lowest resistance rate to tigecycline. The resistance rates of *Pseudomonas aeruginosa* to cefoperazone/sulbactam and imipenem were higher, with the resistance rate to amikacin below 10%. Besides, *Escherichia coli* had high

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resistance rates to ciprofloxacin and cefoperazone/sulbactam and low resistance rates to imipenem, amikacin, and tigecycline.

CONCLUSION

The pathogenic bacteria isolated from the emergency patients were mainly *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*. The detection rates of drug-resistant bacteria were high, with different bacteria having multiple drug resistances to commonly used antimicrobial agents, guiding the rational use of drugs and reducing the production of multidrug-resistant bacteria.

Key words: Distribution; Drug resistance; Bacteria; Emergency department

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Core tip: The purpose of this study is to analyze the distribution and drug resistance of the bacteria isolated from the emergency department specimens, providing the prerequisite for rational use of antibiotics in emergency patients. The top six pathogens were *Acinetobacter baumannii* (660 strains), *Staphylococcus aureus* (436 strains), *Klebsiella pneumoniae* (347 strains), *Pseudomonas aeruginosa* (338 strains), *Escherichia coli* (237 strains), and *Candida albicans* (207 strains). The detection rates of drug-resistant bacteria were high, with different bacteria having multiple drug resistances to commonly used antimicrobial agents, guiding the rational use of drugs and reducing the production of multidrug-resistant bacteria.

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INTRODUCTION

With the wide application of antibacterial drugs, the sensitivity of common bacteria to antibiotics is decreasing. Especially, the resistance to β -lactams, aminoglycosides, fluoroquinolones, and sulfonamides extremely plagues the effective anti-infective treatment and seriously threatens the lives of patients. At the same time, the emergence of new antibacterial drugs makes problem of drug abuse increasingly serious. It has become one of the most important medical problems worldwide. The recent discovery of superbacteria in India and other countries is a serious consequence of the abuse of antimicrobial drugs, but currently there is still no effective treatment that can replace the antibiotics to inhibit and exterminate the pathogens. Therefore, the rational use of antibiotics has become an important topic for clinical use^[1]. However, the premise of rational drug use is to understand the distribution of pathogens and the trends and characteristics of drug resistance.

Generally, patients in the emergency department have a relatively rapid disease progression. Without clear etiology and bacteriological basis, early and empirical use of antibiotics is required. In China, the emergency department of the central hospitals is an important department for the treatment and rescue of critically ill patients. It is difficult to avoid the body injury of patients caused by invasive treatment while rescuing patients, so it is also inevitable to be complicated with various hospital infectious diseases^[2]. Besides, most of the emergency departments are open management, with more patients' family members, more hospital staff flow, and more severe ward environmental pollution, further increasing the probability of hospital-acquired infection in critically ill patients in the emergency department^[3]. For patients with severe infections, a combination of multiple antibiotics is often required, which may lead to bacterial resistance. Due to the great difference in the resistance of different pathogenic bacteria to antibacterial drugs, it brings great inconvenience to clinical medication, followed by poor therapeutic effect^[4,5]. In this study, we analyzed the distribution characteristics and drug resistance to different antibiotics of the bacteria isolated from the emergency department specimens in our hospital from 2013 to 2017. This study is of great significance for the selection of drugs for initial

empirical treatment in the emergency department and the careful selection of antibiotics to reduce the generation of drug-resistant strains.

MATERIALS AND METHODS

Source of strain

This study was approved by the Ethics Committee of Peking University Third Hospital, China, and all patients provided written informed consent.

A total of 2048 critically ill patients hospitalized in the emergency department of Peking University Third Hospital from January 2013 to December 2017 were enrolled. The clinical examination specimens of these patients were collected, including deep sputum retained after adequate gargling (secretions from deep respiratory tract in patients with tracheotomy or intubation), urine, localized secretions, and blood from the apical segment of the catheter in patients with central venous catheterization¹. Finally, the respiratory specimens accounted for 51.6% of all samples, blood specimens accounted for 23.3%, urine specimens accounted for 18.7%, sterile body fluid specimens accounted for 4.0 % and other specimens accounted for 2.4%. The bacteria were then isolated from these specimens. Only the first isolated strains were used for patients with replicate strains isolated. *Staphylococcus aureus* ATCC25923, *Enterococcus faecalis* ATCC29212, *Escherichia coli* ATCC25922, and *Pseudomonas aeruginosa* ATCC27853 were used as control strains.

Methods

Bacterial identification and drug susceptibility testing were carried out according to the routine methods of the National Clinical Laboratory Procedures for bacteria culture, isolation, and identification.

The isolation media including China-blue plate and blood agar plate were purchased from Oxoid, United Kingdom. The bacterial identification was then performed using VITEK 2 Compact automatic microorganism analysis system (bioMérieux, France) and BD-Bruker MALDI Biotyper microorganism mass spectrometry rapid identification system (BD, United Kingdom).

Machine-based and paper-based susceptibility tests were used for evaluating drug sensitivity. Among them, the machine-based susceptibility test applied VITEK 2 Compact automatic microbiological analysis system to determine the minimum inhibitory concentration (MIC) of commonly used antibacterial drugs; the paper-based susceptibility test used Mueller Hinton Agar plates (Oxoid, United Kingdom). Cefoperazone-sulbactam, imipenem, tigecycline, amikacin, ciprofloxacin, and minocycline were purchased from Oxoid, United Kingdom. According to the Clinical and Laboratory Standards Institute (CLSI) M100-S27 document, 2017 edition, the sensitivity of both tests was determined as sensitive (S), intermediate resistant (I), and resistant (R). The determination of susceptibility to cefoperazone-sulbactam referred to cefoperazone. The susceptibility to tigecycline of *Enterobacteriaceae* and *Acinetobacter* was determined based on the United States Food and Drug Administration standard: S (MIC ≤ 2 $\mu\text{g/mL}$), I (MIC ≤ 4 $\mu\text{g/mL}$), and R (MIC ≥ 8 $\mu\text{g/mL}$).

Statistical analysis

Data were analyzed using the R 3.5.1 software. The rate was expressed as a percentage. The Cochran-Armitage trend test (CATT) was used to analyze the change trend of drug resistance rates with time. $P < 0.05$ was considered to be statistically significant.

RESULTS

Bacterial distribution

A total of 3387 pathogens were isolated from various clinical specimens of 2048 critically ill patients, including 1805 strains of Gram-negative bacteria (53.29%), 1141 Gram-positive bacteria (33.69%), and 441 fungi (13.02%). The Gram-negative bacteria were mainly *Acinetobacter baumannii* (660 strains), *Klebsiella pneumoniae* (347 strains), *Pseudomonas aeruginosa* (338 strains), and *Escherichia coli* (237 strains). The Gram-positive bacteria mainly included *Staphylococcus aureus* (436 strains), *Staphylococcus epidermidis* (181 strains), *Staphylococcus haemolyticus* (161 strains), and *Enterococcus faecalis* (140 strains). Besides, the fungi were mainly *Candida albicans* (207 strains), followed by *Candida tropicalis* (95 strains). As a result, the top six bacteria isolated from emergency patients were *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*.

From 2013 to 2017, the isolation rates of common pathogens in the emergency department decreased, including *Acinetobacter baumannii* (25.5% to 17.4%), *Staphylococcus aureus* (21.0% to 7.1%), *Pseudomonas aeruginosa* (12.3% to 10.3%), and *Escherichia coli* (7.6% to 6.5%). However, the isolation rate of *Klebsiella pneumoniae* increased from 7.1% to 12.1% (Figure 1).

Drug resistance of Gram-negative bacteria

Although the isolation rate of *Acinetobacter baumannii* has been decreasing these year, it was still ranked first for many years and was a widely-resistant strain, whose resistance rates to imipenem and ciprofloxacin were above 90%. The drug with the lowest resistance rate of *Acinetobacter baumannii* to common clinical antibiotics was tigecycline, followed by minocycline, but the sensitivity to the two drugs has been also decreasing each year ($P < 0.01$).

The resistance rates of *Pseudomonas aeruginosa* to cefoperazone/sulbactam and imipenem were higher, with the resistance to cefoperazone/sulbactam increasing year over year ($P < 0.01$). But its resistance rate to amikacin was below 10%.

Klebsiella pneumoniae had higher resistance rates to imipenem, cefoperazone/sulbactam, amikacin, and ciprofloxacin, with the lowest resistance rate to tigecycline. The resistance rate to minocycline was also high but it decreased year over year ($P < 0.01$).

For *Escherichia coli*, the resistance to ciprofloxacin was high, and the resistance rate to cefoperazone/sulbactam showed an upward trend year after year ($P < 0.01$). Besides, the resistance rates to imipenem, amikacin, and tigecycline were low, with the resistance rate to tigecycline being zero for the last three years (Table 1).

Drug resistance of Gram-positive bacteria

Staphylococcus aureus is the top one of the Gram-positive bacteria isolated from clinical examination specimens of emergency department patients. As shown in Table 2, the resistance rates of *Staphylococcus aureus* to levofloxacin, penicillin G, and tetracycline were high, but the resistance rates to penicillin G and tetracycline showed a downward trend ($P < 0.01$). Moreover, the resistance rates to vancomycin and tigecycline were always zero.

Analysis of patients infected with *Klebsiella pneumoniae*

As mentioned above, the isolation rates of common pathogens decreased year after year except *Klebsiella pneumoniae*. We then further analyzed the clinical data of all patients infected with *Klebsiella pneumoniae* (Table 3). There were 347 patients with *Klebsiella pneumoniae* infection aged between 18 and 96 years, with an average age of 73.35 ± 14.60 years. Notably, we found that the percentage of patients with deep venous catheterization or retention catheterization was much higher than that of patients without both operations.

DISCUSSION

This study showed that the top six pathogens isolated from emergency patients in our hospital from 2013 to 2017 were *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*, consistent with other reports in the emergency department of the top three hospitals in the same area. Besides, the detection rates of drug-resistant bacteria were high, showing the severe situation of antibiotic resistance. Therefore, these findings provided a good basis for the early delivery of empirical medication for critically ill patients.

With the emergence of antibacterial drugs, the rate of bacterial resistance remains high^[6]. In this study, we found that *Staphylococcus aureus* had high resistance rates to levofloxacin, penicillin G, and tetracycline, but the resistance rates to vancomycin and tigecycline were always zero. Besides, five linezolid-insensitive strains of *Staphylococcus aureus* were discovered, consistent with the previous report^[7]. Therefore, vancomycin should be given priority in the clinical treatment of severe patients with *Staphylococcus aureus* infection. However, in case of adverse reactions, such as hyperpyrexia and hypersensitivity after vancomycin administration, tigecycline may be selected and targeted drugs may be given after drug sensitivity testing^[8,9]. Hussein *et al* indicated that a high MRSA prevalence was found amongst healthcare workers (HCWs) in Kurdistan Region, Iraq. Twenty-two point five percent of HCWs were *Staphylococcus aureus* carriers compared with 18.7% of non-HCWs. 61.0% of *S. aureus* strains isolated from HCWs were MRSA compared with 21.6% from non-HCWs. The mean working years of MRSA carriers was significantly higher than that of MRSA non-carriers. Basic infection control measures, a screening

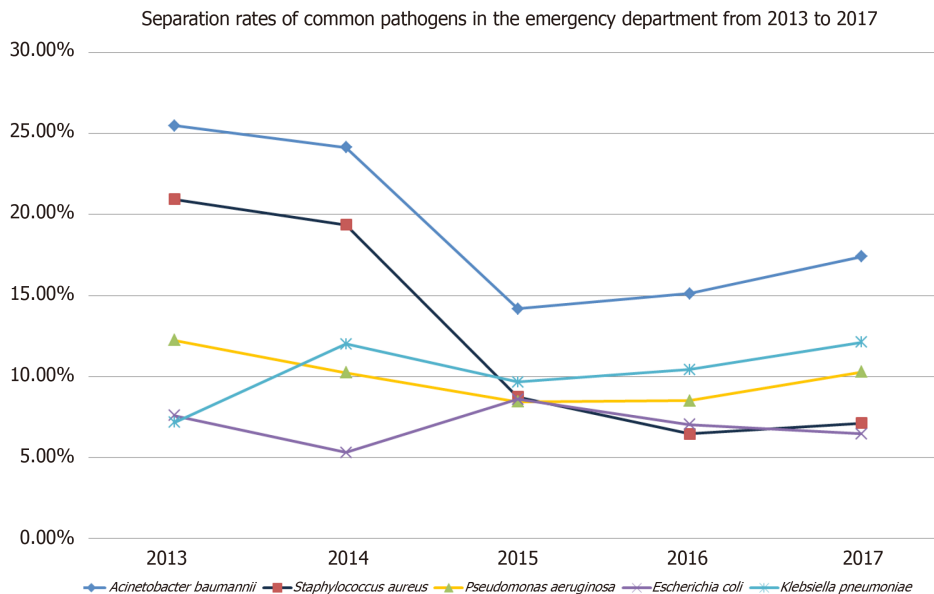


Figure 1 Separation rates of common pathogens in the emergency department from 2013 to 2017.

programme, and treatment of MRSA-positive HCWs can help as an effective measure to control MRSA infections^[10]. In addition, it was shown that among 13 patients infected with vancomycin-resistant *Enterococcus*, most patients had lower limb wound infection^[11]. Therefore, for patients with lower limb infection of vancomycin-resistant enterococcus, strict observation and active treatment are required to prevent multiple infections.

Acinetobacter baumannii is a common colonized pathogen in hospital, and it is also a relatively common pathogen of nosocomial acquired pneumonia^[12]. In this study, its isolation rate ranked first for many years, and it was an extensively drug-resistant strain. *Acinetobacter baumannii* in the emergency department is mostly due to nosocomial infection during hospitalization^[13,14], and there are few reports that *Acinetobacter baumannii* causes community infection.

Klebsiella pneumoniae ranked second in the number of Gram-negative strains isolated in the emergency department. Recently, it has been reported that the drug resistance rate of carbapenems resistant enterobacteriaceae (CRE) shows a steady increasing trend, among which *Klebsiella* bacteria account for the largest proportion^[15]. In this study, *Klebsiella pneumoniae* showed higher resistance to imipenem, cefoperazone/sulbactam, amikacin, and ciprofloxacin. The resistance of *Klebsiella pneumoniae* to carbapenem is mainly due to the production of carbapenemases, and few antibacterial drugs are available for the strains resistant to carbapenem, making clinical treatment difficult. A study in southern Europe showed a higher incidence of inappropriate empirical treatment for the multidrug-resistant *Klebsiella pneumoniae* bloodstream infection, resulting in a more than two-fold increase in patient mortality^[16]. However, it has also been demonstrated that patients who received carbapenem monotherapy or combination therapy within the first five days after blood culture positive for β -lactamase-producing *Klebsiella pneumoniae* infection had a significantly lower mortality than those who received non-carbapenem antibiotics^[17]. Therefore, carbapenems are still recommended as the treatment of choice for patients with severe infections, and compound agents containing β -lactamase inhibitors can be considered for patients with mild to moderate infections. Besides, due to the increasing reports of carbapenem-resistant *Klebsiella pneumoniae* in recent years, high doses of carbapenem can be given for treatment^[18], and tigecycline can also be selected, owing to its good *in vitro* antibacterial activity^[19].

The positive rate of β -lactamase-producing *Escherichia coli* from emergency sources should be low, but our study revealed that *Escherichia coli* had high resistance to ciprofloxacin and cefoperazone/sulbactam, indicating that its resistance to quinolones and cephalosporins is still very prominent. In addition, the resistance rate of *Escherichia coli* to imipenem was low, which was consistent with the previous study^[20].

As mentioned above, *Klebsiella pneumoniae* has been the only bacteria with the increasing detection rate among the most common bacteria at the emergency department for the past five years, and the resistance to antibiotics such as meropenem and imipenem was also gradually increasing. These findings were in agreement with previous study^[21], bringing great difficulties for clinical anti-infective

Table 1 Analysis of resistance rates of Gram-negative bacteria

Gram-negative bacteria	Drugs	2013		2014		2015		2016		2017		CATT	P value
		R	I/S	R	I/S	R	I/S	R	I/S	R	I/S		
<i>Acinetobacter baumannii</i>	Cefoperazone/sulbactam	84	97	91	86	79	15	80	23	80	25	47.72	< 0.01
	Imipenem	174	7	170	7	89	5	97	6	102	3	0.05	0.82
	Tigecycline	7	174	6	171	0	94	10	93	12	93	11.13	< 0.01
	Amikacin	91	90	89	89	-	-	-	-	-	-	0.00	1.00
	Ciprofloxacin	171	10	167	10	89	5	98	5	103	2	1.46	0.23
	Minocycline	116	65	70	107	17	77	29	74	15	90	79.13	< 0.01
<i>Pseudomonas aeruginosa</i>	Cefoperazone/sulbactam	18	69	17	58	7	49	19	39	27	35	10.17	< 0.01
	Imipenem	29	58	39	36	28	28	28	30	24	38	0.31	0.58
	Tigecycline	-	-	-	-	-	-	-	-	-	-	-	-
	Amikacin	5	82	5	70	2	54	5	53	2	60	0.06	0.80
	Ciprofloxacin	19	68	20	55	16	40	15	44	10	52	0.47	0.50
	Minocycline	-	-	-	-	-	-	-	-	-	-	-	-
<i>Klebsiella pneumoniae</i>	Cefoperazone/sulbactam	12	39	58	30	23	41	33	38	40	33	1.94	0.16
	Imipenem	5	46	54	34	25	39	32	39	34	39	3.91	0.05
	Tigecycline	-	-	6	82	5	59	12	59	7	66	1.48	0.22
	Amikacin	1	50	36	52	19	45	28	43	25	48	6.86	< 0.01
	Ciprofloxacin	20	31	58	30	35	29	35	36	40	33	0.04	0.85
	Minocycline	23	28	31	57	19	45	13	58	12	61	17.53	< 0.01
<i>Escherichia coli</i>	Cefoperazone/sulbactam	3	51	2	37	12	45	4	44	13	26	11.30	< 0.01
	Imipenem	0	54	4	35	3	54	0	48	5	34	3.21	0.07
	Tigecycline	-	-	3	36	0	57	0	48	0	39		
	Amikacin	3	51	1	38	1	56	1	47	7	32	3.20	0.07
	Ciprofloxacin	35	19	25	14	34	23	28	20	27	12	0.00	0.98
	Minocycline	19	35	7	32	4	53	9	39	4	35	8.41	< 0.01

S: Sensitive; I: Intermediate resistant; R: Resistant.

treatment. Therefore, we further analyzed the clinical data of all patients infected with *Klebsiella pneumoniae* and found that the proportion of patients with deep venous catheterization or retention catheterization was much higher than that of patients without both operations, indicating that indwelling deep venous catheter or urinary catheter is an independent risk factor for bloodstream infection with *Klebsiella pneumoniae*^[22]. Long-term indwelling urethral catheter causes bloodstream infection likely due to the colonization of *Klebsiella pneumoniae* in damaged urethral mucosa during intubation and its regular release into the blood^[23]. Whereas the central venous catheter provides a direct way for bacteria to invade into the bloodstream, inducing the catheter-related bloodstream infection. Further, because the catheter is left in the blood vessel, the pathogenic bacteria in the blood can easily attach the front end of the catheter to gradually form a biofilm that is difficult to remove, which becomes a secondary infection source and aggravates the severity of infection. Therefore, in patients with clinically diagnosed bloodstream infections, when anti-infective drug therapy fails, the possibility of catheter-associated bacteremia should be considered. The ureter or central venous catheter should be removed timely. Other reports show that the use of antibiotics, especially cephalosporins and quinolones, is a risk factor for *Klebsiella pneumoniae* bloodstream infection^[23,24]. Besides, the total amount and days of antibiotic use were also found to be significantly associated with the development of *Klebsiella pneumoniae* infection^[25]. Hence, short-course antibiotics may be useful in reducing drug-resistant bacteria^[26].

There is a certain course of treatment with antibiotics, which should be administered on time once adopted to maintain sufficient concentration of drugs in patients to reduce the generation of drug-resistant strains^[27,28]. Some recent studies offer some hope for tackling bacterial resistance^[29]. Bacteriophages have many advantages over antibiotics in their use to treat and prevent infection by drug-resistant bacteria. Their therapeutic effects are significantly different from those of antibiotics, making them still sensitive to multidrug-resistant bacteria^[30]. Therefore, phages are currently seen as a potential effective treatment for many multidrug-

Table 2 Analysis of resistance rates of Gram-positive bacteria

Gram-positive bacteria	Drugs	2013		2014		2015		2016		2017		CATT	P value
		R	I/S	R	I/S	R	I/S	R	I/S	R	I/S		
<i>Staphylococcus aureus</i>	Vancomycin	0	149	0	142	0	58	0	44	0	43	-	-
	Levofloxacin	131	18	130	12	38	20	25	19	21	22	53.68	< 0.01
	Linezolid	1	148	4	138	0	58	0	44	0	43	0.75	0.39
	Tigecycline	0	149	0	142	0	58	0	44	0	43	-	-
	PenicillinG	146	3	140	2	55	3	41	3	41	2	3.45	0.06
	Tetracycline	123	26	131	11	35	23	26	18	25	18	28.27	< 0.01

resistant bacteria^[31]. However, at the present stage, it is still of great significance to strengthen the pathogenic examination and monitoring in the emergency department and understand the distribution and drug resistance trends of the prevalent strains, so as to guide the rational use of drugs, reduce the production of multidrug-resistant bacteria, reduce the hospital infection rate and improve the success rate of patient treatment.

In summary, the pathogenic bacteria isolated from the emergency department were mainly *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*, with high detection rates of drug-resistant bacteria. When critically ill patients are admitted to the emergency department, initial antibiotic treatment should be selected empirically according to the distribution characteristics of bacteria in this area while bacteriological examination should be conducted on the clinical samples as soon as possible, and the later drug regimen should be adjusted timely according to the results of pathogen culture and drug sensitivity. For patients with extremely serious infections and life risk at any time, multi-drug regimens can be considered to achieve early control of the disease.

Table 3 Analysis of clinical data of patients with *Klebsiella pneumoniae* infection (*n* = 347)

Influence factors		Cases	Percentage (%)	Influence factors		Cases	Percent age (%)
Gender	Male	175	53.19	Deep vein catheterization	Yes	224	68.09
	Female	154	46.81		No	105	31.91
Age (yr)	≥ 60	277	84.19	Mechanical ventilation	Yes	93	28.27
	< 60	52	15.81		No	236	71.73
Tracheotomy	Yes	9	2.74	Retention catheterization	Yes	241	73.25
	No	320	97.26		No	88	26.75

ARTICLE HIGHLIGHTS

Research background

Antibiotic resistance has become a global threat for human health, calling for rational use of antibiotics.

Research motivation

The premise of rational drug use is to understand the distribution of pathogens and the trends and characteristics of drug resistance.

Research objectives

In this study, we analyzed the distribution characteristics and drug resistance to different antibiotics of the bacteria isolated from the emergency department specimens in our hospital from 2013 to 2017. This study is of great significance for the selection of drugs for initial empirical treatment in the emergency department and the careful selection of antibiotics to reduce the generation of drug-resistant strains.

Research methods

The isolation media including China-blue plate and blood agar plate were purchased from Oxoid, United Kingdom. The bacterial identification was then performed using VITEK 2 Compact automatic microorganism analysis system (bioMérieux, France) and BD-Bruker MALDI Biotyper microorganism mass spectrometry rapid identification system (BD, United Kingdom). Data were analyzed using the R 3.5.1 software. The rate was expressed as a percentage. The Cochran-Armitage trend test was used to analyze the change trend of drug resistance rates with time. $P < 0.05$ was considered to be statistically significant.

Research results

The top six bacteria isolated from emergency patients were *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*. From 2013 to 2017, the isolation rates of common pathogens in the emergency department decreased, including *Acinetobacter baumannii* (25.5% to 17.4%), *Staphylococcus aureus* (21.0% to 7.1%), *Pseudomonas aeruginosa* (12.3% to 10.3. %), and *Escherichia coli* (7.6% to 6.5%). However, the isolation rate of *Klebsiella pneumoniae* increased from 7.1% to 12.1%. The drug with the lowest resistance rate of *Acinetobacter baumannii* to common clinical antibiotics was tigecycline, followed by minocycline, but the sensitivity to the two drugs has been also decreasing each year ($P < 0.01$). The resistance rates of *Pseudomonas aeruginosa* to cefoperazone/sulbactam and imipenem were higher, with the resistance to cefoperazone/sulbactam increasing year over year ($P < 0.01$). But its resistance rate to amikacin was below 10%. *Klebsiella pneumoniae* had higher resistance rates to imipenem, cefoperazone/sulbactam, amikacin, and ciprofloxacin, with the lowest resistance rate to tigecycline. The resistance rate to minocycline was also high but it decreased year over year ($P < 0.01$). For *Escherichia coli*, the resistance to ciprofloxacin was high, and the resistance rate to cefoperazone/sulbactam showed an upward trend year after year ($P < 0.01$). the resistance rates of *Staphylococcus aureus* to levofloxacin, penicillin G, and tetracycline were high, but the resistance rates to penicillin G and tetracycline showed a downward trend ($P < 0.01$).

Research conclusions

The pathogenic bacteria isolated from the emergency department were mainly *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*, with high detection rates of drug-resistant bacteria. When critically ill patients are admitted to the emergency department, initial antibiotic treatment should be selected empirically according to the distribution characteristics of bacteria in this area while bacteriological examination should be conducted on the clinical samples as soon as possible, and the later drug regimen should be adjusted timely according to the results of pathogen culture and drug sensitivity. For patients with extremely serious infections and life risk at any time, multi-drug regimens can be considered to achieve early control of the disease. These findings provided a good basis for the early delivery of empirical medication for critically ill patients.

There is a certain course of treatment with antibiotics, which should be administered on time once adopted to maintain sufficient concentration of drugs in patients to reduce the generation of drug-resistant strains. Some recent studies offer some hope for tackling bacterial resistance. Bacteriophages have many advantages over antibiotics in their use to treat and prevent infection by drug-resistant bacteria. Their therapeutic effects are significantly different from those of antibiotics, making them still sensitive to multidrug-resistant bacteria. A total of 2048 critically ill patients were enrolled. The clinical examination specimens of these patients were collected, including deep sputum retained after adequate gargling (secretions from deep respiratory tract in patients with tracheotomy or intubation), urine, localized secretions, and blood from the apical segment of the catheter in patients with central venous catheterization. As mentioned above, *Klebsiella pneumoniae* has been the only bacteria with the increasing detection rate among the most common bacteria at the emergency department for the past five years, and the resistance to antibiotics such as meropenem and imipenem was also gradually increasing. The pathogenic bacteria isolated from the emergency department were mainly *Acinetobacter baumannii*, *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Candida albicans*, with high detection rates of drug-resistant bacteria. Phages are currently seen as a potential effective treatment for many multidrug-resistant bacteria. However, at the present stage, it is still of great significance to strengthen the pathogenic examination and monitoring in the emergency department and understand the distribution and drug resistance trends of the prevalent strains, so as to guide the rational use of drugs, reduce the production of multidrug-resistant bacteria, reduce the hospital infection rate and improve the success rate of patient treatment.

Research perspectives

There may be bias in data collection of retrospective studies. The future research direction is the rational use of antibiotics.

REFERENCES

- 1 Christensen A, Scheel O, Urwitz K, Bergh K. Outbreak of methicillin-resistant *Staphylococcus aureus* in a Norwegian hospital. *Scand J Infect Dis* 2001; **33**: 663-666 [PMID: 11669223 DOI: 10.1080/00365540110026944]
- 2 Leroy S, Marc E, Bavoux F, Tréluyer JM, Gendrel D, Bréart G, Pons G, Chalumeau M. Hospitalization for severe bacterial infections in children after exposure to NSAIDs: a prospective adverse drug reaction reporting study. *Clin Drug Investig* 2010; **30**: 179-185 [PMID: 20155990 DOI: 10.2165/11532890-000000000-00000]
- 3 Ostrowsky B, Steinberg JT, Farr B, Sohn AH, Sinkowitz-Cochran RL, Jarvis WR. Reality check: should we try to detect and isolate vancomycin-resistant enterococci patients? *Infect Control Hosp Epidemiol* 2001; **22**: 116-119 [PMID: 11232874 DOI: 10.1086/501874]
- 4 Pallin DJ, Camargo CA, Schuur JD. Skin infections and antibiotic stewardship: analysis of emergency department prescribing practices, 2007-2010. *West J Emerg Med* 2014; **15**: 282-289 [PMID: 24868305 DOI: 10.5811/westjem.2013.8.18040]
- 5 Lin YB, Gardiner MF. Fingernail-induced corneal abrasions: case series from an ophthalmology emergency department. *Cornea* 2014; **33**: 691-695 [PMID: 24831196 DOI: 10.1097/ICO.0000000000000133]
- 6 Magalhães M, Doherty C, Govan JR, Vandamme P. Polyclonal outbreak of *Burkholderia cepacia* complex bacteraemia in haemodialysis patients. *J Hosp Infect* 2003; **54**: 120-123 [PMID: 12818585 DOI: 10.1016/S0195-6701(03)00118-X]
- 7 Mendes RE, Flamm RK, Hogan PA, Ross JE, Jones RN. Summary of linezolid activity and resistance mechanisms detected during the 2012 LEADER surveillance program for the United States. *Antimicrob Agents Chemother* 2014; **58**: 1243-1247 [PMID: 24323470 DOI: 10.1128/AAC.02112-13]
- 8 Vandijck DM, Blot SI, Decruyenaere JM. Update on the management of infection in patients with severe sepsis. *Dimens Crit Care Nurs* 2008; **27**: 244-248 [PMID: 18953190 DOI: 10.1097/01.DCC.0000338868.31917.0e]
- 9 Díaz A, Barria P, Niederman M, Restrepo MI, Dreyse J, Fuentes G, Couble B, Saldias F. Etiology of community-acquired pneumonia in hospitalized patients in Chile: the increasing prevalence of respiratory viruses among classic pathogens. *Chest* 2007; **131**: 779-787 [PMID: 17356093 DOI: 10.1378/chest.06-1800]
- 10 Miyaji R, Asano K, Matsubara S. Induction of Axial Chirality in 8-Arylquinolines through Halogenation Reactions Using Bifunctional Organocatalysts. *Chemistry* 2017; **23**: 9996-10000 [PMID: 28429870 DOI: 10.1002/chem.201701707]
- 11 Öksüz L, Gürlü N. [Serotype distribution and antibiotic resistance of *Streptococcus pneumoniae* strains isolated from the adult patients in a Turkish university hospital]. *Mikrobiyol Bul* 2017; **51**: 195-208 [PMID: 28929957 DOI: 10.5578/mb.48638]
- 12 Falcone M, Russo A, Giannella M, Cangemi R, Scarpellini MG, Bertazzoni G, Alarcón JM, Taliani G, Palange P, Farcomeni A, Vestri A, Bouza E, Violi F, Venditti M. Individualizing risk of multidrug-resistant pathogens in community-onset pneumonia. *PLoS One* 2015; **10**: e0119528 [PMID: 25860142 DOI: 10.1371/journal.pone.0119528]
- 13 Chan J, Wong J, Saginur R, Forster AJ, van Walraven C. Epidemiology and outcomes of bloodstream infections in patients discharged from the emergency department. *CJEM* 2015; **17**: 27-37 [PMID: 25781381 DOI: 10.2310/8000.2013.131349]
- 14 Arbex MA, Pereira LA, Carvalho-Oliveira R, Saldiva PH, Braga AL. The effect of air pollution on pneumonia-related emergency department visits in a region of extensive sugar cane plantations: a 30-month time-series study. *J Epidemiol Community Health* 2014; **68**: 669-674 [PMID: 24782416 DOI: 10.1136/jech-2013-203709]
- 15 Xu Y, Gu B, Huang M, Liu H, Xu T, Xia W, Wang T. Epidemiology of carbapenem resistant Enterobacteriaceae (CRE) during 2000-2012 in Asia. *J Thorac Dis* 2015; **7**: 376-385 [PMID: 25922715]

- DOI: [10.3978/j.issn.2072-1439.2014.12.33](https://doi.org/10.3978/j.issn.2072-1439.2014.12.33)]
- 16 **Girometti N**, Lewis RE, Giannella M, Ambretti S, Bartoletti M, Tedeschi S, Tumietto F, Cristini F, Trapani F, Gaibani P, Viale P. *Klebsiella pneumoniae* bloodstream infection: epidemiology and impact of inappropriate empirical therapy. *Medicine (Baltimore)* 2014; **93**: 298-309 [PMID: [25398065](https://pubmed.ncbi.nlm.nih.gov/25398065/) DOI: [10.1097/MD.0000000000000111](https://doi.org/10.1097/MD.0000000000000111)]
 - 17 **Patterson DL**, Ko WC, Von Gottberg A, Mohapatra S, Casellas JM, Goossens H, Mulazimoglu L, Trenholme G, Klugman KP, Bonomo RA, Rice LB, Wagener MM, McCormack JG, Yu VL. Antibiotic therapy for *Klebsiella pneumoniae* bacteremia: implications of production of extended-spectrum beta-lactamases. *Clin Infect Dis* 2004; **39**: 31-37 [PMID: [15206050](https://pubmed.ncbi.nlm.nih.gov/15206050/) DOI: [10.1086/420816](https://doi.org/10.1086/420816)]
 - 18 **Giannella M**, Trecarichi EM, Giacobbe DR, De Rosa FG, Bassetti M, Bartoloni A, Bartoletti M, Losito AR, Del Bono V, Corcione S, Tedeschi S, Raffaelli F, Saffioti C, Spanu T, Rossolini GM, Marchese A, Ambretti S, Cauda R, Viscoli C, Lewis RE, Viale P, Tumbarello M; Italian Study Group on Resistant Infections of the Società Italiana Terapia Antinfettiva (ISGRI-SITA). Effect of combination therapy containing a high-dose carbapenem on mortality in patients with carbapenem-resistant *Klebsiella pneumoniae* bloodstream infection. *Int J Antimicrob Agents* 2018; **51**: 244-248 [PMID: [28842283](https://pubmed.ncbi.nlm.nih.gov/28842283/) DOI: [10.1016/j.ijantimicag.2017.08.019](https://doi.org/10.1016/j.ijantimicag.2017.08.019)]
 - 19 **Geng TT**, Xu X, Huang M. High-dose tigecycline for the treatment of nosocomial carbapenem-resistant *Klebsiella pneumoniae* bloodstream infections: A retrospective cohort study. *Medicine (Baltimore)* 2018; **97**: e9961 [PMID: [29465589](https://pubmed.ncbi.nlm.nih.gov/29465589/) DOI: [10.1097/MD.0000000000000996](https://doi.org/10.1097/MD.0000000000000996)]
 - 20 **Bou-Antoun S**, Davies J, Guy R, Johnson AP, Sheridan EA, Hope RJ. Descriptive epidemiology of *Escherichia coli* bacteraemia in England, April 2012 to March 2014. *Euro Surveill* 2016; **21** [PMID: [27608263](https://pubmed.ncbi.nlm.nih.gov/27608263/) DOI: [10.2807/1560-7917.ES.2016.21.35.30329](https://doi.org/10.2807/1560-7917.ES.2016.21.35.30329)]
 - 21 **Tian L**, Tan R, Chen Y, Sun J, Liu J, Qu H, Wang X. Epidemiology of *Klebsiella pneumoniae* bloodstream infections in a teaching hospital: factors related to the carbapenem resistance and patient mortality. *Antimicrob Resist Infect Control* 2016; **5**: 48 [PMID: [27891222](https://pubmed.ncbi.nlm.nih.gov/27891222/) DOI: [10.1186/s13756-016-0145-0](https://doi.org/10.1186/s13756-016-0145-0)]
 - 22 **Bilavsky E**, Temkin E, Lerman Y, Rabinovich A, Salomon J, Lawrence C, Rossini A, Salvia A, Samso JV, Fierro J, Paul M, Hart J, Gniadkowski M, Hochman M, Kazma M, Klein A, Adler A, Schwaber MJ, Carmeli Y; MOSAR WP5 study team. Risk factors for colonization with extended-spectrum beta-lactamase-producing enterobacteriaceae on admission to rehabilitation centres. *Clin Microbiol Infect* 2014; **20**: O804-O810 [PMID: [24674024](https://pubmed.ncbi.nlm.nih.gov/24674024/) DOI: [10.1111/1469-0691.12633](https://doi.org/10.1111/1469-0691.12633)]
 - 23 **Freeman JT**, Nimmo J, Gregory E, Tiong A, De Almeida M, McAuliffe GN, Roberts SA. Predictors of hospital surface contamination with Extended-spectrum β -lactamase-producing *Escherichia coli* and *Klebsiella pneumoniae*: patient and organism factors. *Antimicrob Resist Infect Control* 2014; **3**: 5 [PMID: [24491119](https://pubmed.ncbi.nlm.nih.gov/24491119/) DOI: [10.1186/2047-2994-3-5](https://doi.org/10.1186/2047-2994-3-5)]
 - 24 **Wener KM**, Schechner V, Gold HS, Wright SB, Carmeli Y. Treatment with fluoroquinolones or with beta-lactam-beta-lactamase inhibitor combinations is a risk factor for isolation of extended-spectrum-beta-lactamase-producing *Klebsiella* species in hospitalized patients. *Antimicrob Agents Chemother* 2010; **54**: 2010-2016 [PMID: [20211888](https://pubmed.ncbi.nlm.nih.gov/20211888/) DOI: [10.1128/AAC.01131-09](https://doi.org/10.1128/AAC.01131-09)]
 - 25 **Shanthi M**, Sekar U. Extended spectrum beta lactamase producing *Escherichia coli* and *Klebsiella pneumoniae*: risk factors for infection and impact of resistance on outcomes. *J Assoc Physicians India* 2010; **58** Suppl: 41-44 [PMID: [21568008](https://pubmed.ncbi.nlm.nih.gov/21568008/)]
 - 26 **Qadir MI**. Review: phage therapy: a modern tool to control bacterial infections. *Pak J Pharm Sci* 2015; **28**: 265-270 [PMID: [25553704](https://pubmed.ncbi.nlm.nih.gov/25553704/)]
 - 27 **Angus DC**. Management of sepsis: a 47-year-old woman with an indwelling intravenous catheter and sepsis. *JAMA* 2011; **305**: 1469-1477 [PMID: [21467273](https://pubmed.ncbi.nlm.nih.gov/21467273/) DOI: [10.1001/jama.2011.438](https://doi.org/10.1001/jama.2011.438)]
 - 28 **Safdar N**, Maki DG. The pathogenesis of catheter-related bloodstream infection with noncuffed short-term central venous catheters. *Intensive Care Med* 2004; **30**: 62-67 [PMID: [14647886](https://pubmed.ncbi.nlm.nih.gov/14647886/) DOI: [10.1007/s00134-003-2045-z](https://doi.org/10.1007/s00134-003-2045-z)]
 - 29 **Viertel TM**, Ritter K, Horz HP. Viruses versus bacteria-novel approaches to phage therapy as a tool against multidrug-resistant pathogens. *J Antimicrob Chemother* 2014; **69**: 2326-2336 [PMID: [24872344](https://pubmed.ncbi.nlm.nih.gov/24872344/) DOI: [10.1093/jac/dku173](https://doi.org/10.1093/jac/dku173)]
 - 30 **Chanishvili N**. Bacteriophages as Therapeutic and Prophylactic Means: Summary of the Soviet and Post Soviet Experiences. *Curr Drug Deliv* 2016; **13**: 309-323 [PMID: [27090515](https://pubmed.ncbi.nlm.nih.gov/27090515/)]
 - 31 **Macía-Rodríguez C**, Alende-Castro V, Vázquez-Ledo L, Novo-Veleiro I, González-Quintela A. Skin and soft-tissue infections: Factors associated with mortality and re-admissions. *Enferm Infecc Microbiol Clin* 2017; **35**: 76-81 [PMID: [27061974](https://pubmed.ncbi.nlm.nih.gov/27061974/) DOI: [10.1016/j.eimc.2016.02.030](https://doi.org/10.1016/j.eimc.2016.02.030)]



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