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**Metastatic infection caused by hypervirulent *Klebsiella pneumonia* and co-infection with *Cryptococcus* meningitis: A case report**

Shi YF *et al*. Hypervirulent *K. pneumonia* complicated with *Cryptococcus* meningitis

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**Author contributions:** Shi YF and Wu BQ analyzed the data, designed the report, and drafted the manuscript; Wang YK collected the data, reviewed the literature, and contributed to manuscript drafting; Wang YH and Liu H made contributions to collect the patient’s clinical data; Shi XH collected the data and revised the manuscript; Li XJ performed the microbiological analyses; Wu BQ was responsible for the revision of the manuscript for important content; All authors approved the final version.

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**Abstract**

***BACKGROUND***

*Klebsiella pneumoniae* (*K. pneumoniae*)used to affect mainly people with compromised immunity or weakened by other infections, but recent emergence of hypervirulent strains has increased infections even in healthy individuals. These infections include liver abscess, pneumonia, bacteremia, meningitis, necrotizing fasciitis, and endophthalmitis. Although metastatic infection by hypervirulent *K. pneumoniae* (hvKP) is increasingly recognized, co-infection with *Cryptococcus neoformans* (*C. neoformans*)meningitis in immunocompetent hosts is rare but fatal. So, it is necessary to determine the risk factors, complications, and comorbidity of this disease.

***CASE SUMMARY***

This report describes a 58-year-old man with hvKP pulmonary abscess, bacteremia, and meningitis, accompanied by fatal *Cryptococcus* meningitis. This patient presented with fever for 1 wk and drowsiness for 3 d. Laboratory findings revealed pulmonary abscess and bacteremia of *K. pneumoniae*. He was given intravenous antibiotic therapy, and the infection was under control for about 1 wk. However, his condition deteriorated rapidly because of metastatic purulent meningitis. Although hvKP and *C. neoformans* were isolated and confirmed, the patient died of spontaneous respiratory and cardiac arrest caused by cerebral hernia.

***CONCLUSION***

HvKP has emerged as a cause of metastatic infections in immunocompetent hosts. polymicrobial co-infections should be taken into consideration when metastatic infection is present.

**Key words:** Hypervirulent *Klebsiella pneumoniae*; Metastatic infection; *Cryptococcus neoformans* meningitis; Comorbidity; Case report

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**Core tip:** Hypervirulent *Klebsiella pneumoniae* (hvKP) has emerged as a cause of metastatic infections in immunocompetent hosts and presents mainly as a monomicrobial infection. We present a rare case of metastatic infection caused by hvKP and co-infection with *Cryptococcus neoformans* meningitis. The patient received antibiotics that were sensitive to pathogens in time, and the infection was under control for 1 wk. However, his condition deteriorated rapidly because of metastatic purulent meningitis, and he died of spontaneous respiratory and cardiac arrest caused by cerebral hernia. This case highlights the risk of complications and polymicrobial co-infections in hvKP infected patients. Timely ventricle drainage is strongly recommended for polymicrobial co-infected meningitis.

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**INTRODUCTION**

*Klebsiella pneumoniae* (*K. pneumoniae*)is a Gram-negative, nonmotile, capsulated, aerobic bacterium. It is widely found in nature, as well as in hospital settings, causing community-acquired and nosocomial infections. Invasive infections resulting from *K. pneumoniae* dissemination include liver abscess, pneumonia, bacteremia, meningitis, necrotizing fasciitis, endophthalmitis, and even sepsis[1]. In the past decade, more attention has been given to cases of hypervirulent *K. pneumoniae* (hvKP) infection in eastern countries. The death rate from hvKP infection has been as high as 60% as a result of antibiotic-resistant strains, despite the use of broad-spectrum antibiotics such as carbapenems[2,3].

Metastatic infection caused by hvKP is generally community-acquired and presents mainly as a monomicrobial disease. Here, we present and discuss a rare case of metastatic infection due to hvKP with pulmonary abscess, bacteremia, purulent meningitis, and co-infection with fatal *Cryptococcus* meningitis. To the best of our knowledge, this is the first case of such a metastatic infection caused by hvKP and *C. neoformans*, which should raise concern among clinicians for the risk of such infection.

**CASE PRESENTATION**

***Chief complaints***

Fever for 1 wk and drowsiness for 3 d.

***History of present illness***

The patient, a 58-year-old man, presented with high fever (> 40.0 °C) for 1 wk before admission, with simultaneous fatigue, occasional cough without sputum, and swelling of the left upper limb. The patient was referred to Sun Yat-sen University Cancer Center (SYSUCC) at first because routine blood tests in his hometown showed white blood cell (WBC) count 34.9 × 109/L and hemoglobin level 67 g/L. In the next 6 d at SYSUCC, laboratory findings confirmed pulmonary abscess and bacteremia caused by *K. pneumoniae*. He was given antibiotic therapy of imipenem–cilastatin combined with voriconazole at normal dose. However, he still had high fever and gradually developed altered mental state, which presented as abepithymia and drowsiness. There was no headache, nausea, or vomiting. Based on a rapid increase in creatinine (from normal to 264 µmol/L) and bacteremia, he was diagnosed with acute renal insufficiency and sepsis and admitted to the Medical Intensive Care Unit (MICU) of the Third Affiliated Hospital of Sun Yat-Sen University.

***History of past illness***

The patient had been well prior to onset of the present illness. He had no chronic illness such as diabetes mellitus, immunodeficiency, or psychiatric or psychological disease. He had no tuberculosis or lung cancer.

***Personal and family history***

The patient was a non-smoker. He had no history of alcohol or illicit drug abuse. He resided in Guangdong, southern China. He had no recent travel, tick bites, or contact with sick people.

***Physical examination upon admission***

When the patient was admitted to MICU (day 1), his vital signs were as follows: temperature 36.4 °C, pulse rate 95 beats/min, respiratory rate 20 breaths/min, blood pressure 126/85 mmHg, and pulse oxygen saturation 98% on 5 L/min oxygen flow rate. Low respiratory tone and fine crackles in the right upper lung were noted on pulmonary auscultation. There was no audible murmur on cardiac auscultation. Tenderness and hepatosplenomegaly were not detected. Tenderness in his left upper limb was noted. Both lower limbs had mild edema. He had abepithymia but was still conscious, oriented with Glasgow Coma Scale score 15, and had no neck stiffness. No rash was observed.

***Laboratory findings***

Laboratory tests from SYSUCC revealed WBC count 27.0 × 109/L with an elevated neutrophil ratio of 95.4%. The concentration of C-reactive protein (CRP) and procalcitonin (PCT) were 233.17 mg/L and 1.7 ng/mL, respectively. Glucose concentration was 7.11 mmol/L. Full blood culture grew *K. pneumoniae* (Table 1)*.* Cryptococcal antigen was negative. Galacto Mannan test and β-D-Glucan test both were negative. Concentration of G-lipopolysaccharides was 0.13 EU/mL. Bone marrow smear and biopsy confirmed no malignant tumor cells. Chest computed tomography (CT) scan found a lump that could not be identified as a tumor or infection in the upper right lung. Head CT scan showed no acute findings. CT scan of the left upper limb revealed no abnormality.

Laboratory data upon admission in MICU demonstrated WBC count 26.95 × 109/L with an elevated neutrophil ratio of 94.6%. CRP was 63.9 mg/L, PCT was 10.8 ng/mL, and erythrocyte sedimentation rate was 126 mm/h. Electrolytes were within normal limits. The concentration of glucose was 7.0 mmol/L with hemoglobin A1c in the normal range. Albumin level was 28.4 g/L. Creatinine was 341 μmol/L. Bedside cardiac ultrasound found left ventricular ejection fraction of 0.63 and no pericardial effusion. No intra-abdominal pathology was observed by bedside ultrasound. Chest CT scan found a 25 mm × 27mm, round, thick-walled cavity in the right upper lobe (Figure 1A and B).

**FINAL DIAGNOSIS**

Based on the clinical data, the final diagnoses were confirmed as pulmonary abscess, bacteremia, metastatic purulent meningitis, and sepsis due to hvKP, with co-infection with *C. neoformans*.

**TREATMENT**

The patient received imipenem–cilastatin, tigecycline combined with voriconazole (Figure 2), together with supportive treatment. The patient continued to have high fever (maximum 39.3 °C), which finally began to relieve 4 d after treatment. After 11 d of treatment, his WBC count dropped continuously to 8.2 × 109/L, with a normal neutrophil predominance of 78%, as well as decrease of CRP and PCT (Figures 3 and 4). Creatinine dropped from 341 to 57 µmol/L. The lung abscess was lessened upon CT scan (Figure 1). Two sets of peripheral blood cultures were ordered during episodes of fever, but all results were negative. The patient was transferred to the Respiratory Department (a general ward) on day 11. However, 3 d later, the patient had projectile vomiting and altered mental state. Physical examination demonstrated intermittent somnolence and neck stiffness. Head CT scan and lumbar puncture were arranged. CT images showed marked dilatation of the bilateral and third ventricles caused by hydrocephalus (Figure 5). Lumbar puncture showed intracranial pressure of 120 mmH2O, and cerebrospinal fluid (CSF) was transparent and colorless. CSF was collected for routine examination, biochemistry, Gram staining, and bacterial culture. CSF cytology and biochemistry showed 1560 WBCs/µL with 90% neutrophils, 18 red blood cells/µL, protein 2.03 g/L, glucose 0.03 mmol/L, and chloride ions 110.3 mmol/L. It was noticeable that *C. neoformans* was found by CSF staining, with a count of 85566/mL.

The patient was transferred to the Neurology Department on day 16. He was intubated and transferred to the neurologic intensive care unit because of unstable vital signs on the same night. Considering the high risk of surgery, the patient’s family refused bilateral ventricular drainage that was proposed by a neurosurgeon. The patient was given anti-infective therapy of meropenem combined with 5-fluorocytosine and fluconazole. On day 18, the patient’s vital signs and homeostasis deteriorated rapidly. He died of cerebral hernia caused by co-infection with purulent meningitis. *K. pneumoniae* was subsequently isolated from CSF (Table 2), closely followed by isolation from sputum collected by bronchoscopy (Table 3). Both were string test positive, which reflected the hypermucoviscous phenotype and were considered as hypervirulent strains.

**OUTCOME AND FOLLOW-UP**

The infection was under control for 1 wk, but his condition deteriorated rapidly because of co-infective purulent meningitis. Unfortunately, the patient died of spontaneous respiratory and cardiac arrest caused by cerebral hernia.

**DISCUSSION**

A number of serious infections caused by *K. pneumoniae* that may lack response to anti-infective therapy have been increasingly recorded over the past decade. These strains acquire extra genetic traits and become hypervirulent or antibiotic resistant[4]. Capsular polysaccharide is the most important virulence factor of *K. pneumoniae*, and others include fimbriae, lipopolysaccharide, determinants of iron acquisition, outer membrane proteins, and nitrogen source utilization[5]. Some underlying risk factors such as diabetes mellitus may damage host defense by causing inhibition of phagocytosis and bactericidal activity, which contributes to the initiation and spread of infection[6]. Bacterial virulence is still the primary and most important factor. Since the 1980s, hvKP from the community has attracted increased attention due to its ability to cause metastatic and aggressive infections called “invasive klebsiella syndrome” in healthy hosts[7-9]. There is an urgent need to highlight clinical awareness and management of hvKP infection.

Over the past 10 years, hvKP has become more pervasive with a varied geographic distribution and more likely to cause serious infection in China. Some strains present with both hypervirulence and antibiotic resistance[10,11]. Compared with classic *K. pneumoniae*, hvKP is more effective at capsule production, forming a high mucous phenotype and causing severe invasive multi organ infection[9]. However, there is no internationally agreed definition for hvKP and its virulence level. Eight capsular serotypes, such as K1 and K2, have been associated with hypervirulent strains[12,13]. Besides, mucoid phenotype A (rmpA) and mucoviscosity-associated gene A (magA) are two major genes linked with hypervirulence. Hypermucoviscosity correlates with higher virulence gene content[14], which supports the value of string tests. The string test, with a viscous string > 5 mm in length, reflects the hypermucoviscous phenotype, and is still the most common laboratory-based method presently available[8]. The string test for *K. pneumoniae* in the present case was positive. This hypervirulent strain caused pulmonary abscess, bacteremia, and metastatic purulent meningitis.

*K. pneumoniae* remains one of the most prevalent pathogens in bacterial meningitis[15,16]. *C. neoformans* is the most common cause of fungal meningitis in immunocompetent hosts[17]. Meningitis caused by mixed infection of *K. pneumoniae* and *C. neoformans* is rare. Despite the use of potent antibiotics, the mortality of *K. pneumoniae* meningitis remains high. In such cases, invasive treatment, such as lateral ventricular drainage or irrigation, is recommended to reduce intracranial pressure and relieve inflammatory reaction in the brain tissue[18], especially for those cases with fatal polymicrobial co-infection[19].

In the present case, the middle-aged male patient neither became infected in hospital, nor did he have putative risk factors such as history of diabetes mellitus or hypoimmunity. Until to his death, the clinical data strongly supported the diagnosis of a severe outbreak of sepsis caused by *K. pneumoniae*, which caused metastatic infection accompanied by *C. neoformans* meningitis. Besides, the patient initially complained of swelling of the left upper limb, which reminded us of complicated skin and soft tissue infections. However, there is no evidence to confirm that, and the swelling was relieved during hospitalization. *K. pneumoniae* was not isolated again until the patient’s death, nor was *C. neoformans* identified. The *K. pneumoniae* strain was not further investigated genetically to confirm the capsular serotype or the virulence genes. A positive string test indicated a hypervirulent strain, which explained the aggressive progression of infection. Considering the difficulties for identification of potential pathogens, which maybe polymicrobial, high-throughput approaches such as biomarkers have been increasingly used in scientific research and clinical studies in recent years[20].

Early recognition of meningitis is not always easy, particularly when the patient presents with nonspecific symptoms. Detailed history of illness, systematic examination, and multidisciplinary consultation are required. For example, for cases with obviously altered mental state, clinicians should consider the possibility of cryptococcal meningitis[21]. Besides anti-infective therapy, timely cerebroventricular drainage is equally or even more important when purulent meningitis is confirmed to improve clinical outcomes[19].

In conclusion, hvKP has emerged as a not uncommon pathogen that could cause metastatic and aggressive infections. Meningitis should be taken into consideration when a metastatic infection is present. Timely anti-infection therapy and ventricle drainage are strongly recommended, especially for polymicrobial co-infection cases.

At last, although it is not possible to explore thoroughly such a complicated infection because of the limitations of case reports, it is our duty to report and accumulate such cases.

**CONCLUSION**

HvKP has emerged as a cause of metastatic infections in immunocompetent hosts. Polymicrobial co-infections should be taken into consideration when metastatic infection is present.

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**Table 1 Resistance profiles and minimum inhibitory concentrations for *Klebsiella pneumoniae* from blood culture**

|  |  |  |
| --- | --- | --- |
| **Antibiotic** | **Drug sensitivity** | **MIC** |
| Ampicillin | R | ≥ 32 |
| Cefazolin | S | ≤ 4 |
| Ceftazidime | S | ≤ 1 |
| Ceftriaxone | S | ≤ 1 |
| Cefepime | S | ≤ 1 |
| Aztreonam | S | ≤ 1 |
| Imipenem  | S | ≤ 1 |
| Ertapenem | S | ≤ 0.5 |
| Amoxicillin/clavulanate | R | UD |
| Piperacillin/tazobactam | S | ≤ 4 |
| Ciprofloxacin | S | ≤ 0.25 |
| Levofloxacin | S | ≤ 0.25 |
| Tobramycin | S | ≤ 1 |
| Gentamicin | S | ≤ 1 |
| Amikacin | S | ≤ 2 |
| Cefotiam | S | UD |
| Cefotetan | S | ≤ 4 |
| Latamoxef | S | UD |
| Cefoperazone/sulbactam | S | ≤ 4 |
| Ampicillin/sulbactam | S | 4 |
| SMZCo | S | ≤ 20 |
| Macrodantin | I | 64 |
| Moxifloxacin | S | ≤ 0.25 |
| ESBL detection | Negative |

Results from Sun Yat-Sen University Cancer Center. Reported day before admission to Medical Intensive Care Unit of The Third Affiliated Hospital of Sun Yat-Sen University. ESBL: Extended-spectrum β-lactamase; I: Intermediate; MIC: Minimum inhibitory concentration; R: Resistant; S: Sensitive; UD: Undetected; SMZCo: Compound sulfamethoxazole.

**Table 2 Resistance profiles and minimum inhibitory concentrations for *Klebsiella pneumoniae* from cerebrospinal fluid culture**

|  |  |  |
| --- | --- | --- |
| **Antibiotics** | **Drug sensitivity** | **MIC** |
| Ampicillin | R | 16 |
| Piperacillin | S | ≤ 16 |
| Cefazolin | S | ≤ 8 |
| Cefoxitin | S | ≤ 8 |
| Cefuroxime | S | ≤ 4 |
| Cefotaxime | S | ≤ 2 |
| Ceftazidime | S | ≤ 1 |
| Ceftriaxone | S | ≤ 8 |
| Cefepime | S | ≤ 8 |
| Aztreonam | S | ≤ 8 |
| Imipenem | S | ≤ 1 |
| Meropenem | S | ≤ 1 |
| Ertapenem | S | ≤ 2 |
| Amoxicillin/clavulanate | S | ≤ 8/4 |
| Piperacillin/tazobactam | S | ≤ 16 |
| Ticarcillin/clavulanate | S | ≤ 16 |
| Ciprofloxacin | S | ≤ 1 |
| Levofloxacin | S | ≤ 2 |
| Tobramycin | S | ≤ 4 |
| Gentamicin | S | ≤ 4 |
| Amikacin | S | ≤ 8 |
| Tetracycline | S | ≤ 4 |
| Trimethoprim/sulfanilamide | S | ≤ 2/38 |
| Cefotaxime/clavulanate | S | ≤ 0.5 |
| Ceftazidime/ clavulanate | S | ≤ 0.25 |

Results from the Third Affiliated Hospital of Sun Yat-Sen University. Report date: 1 d after death. MIC: Minimum inhibitory concentration; R: Resistant; S: Sensitive.

**Table 3 Resistance profiles and minimum inhibitory concentrations for *Klebsiella pneumoniae* from sputum culture**

|  |  |  |
| --- | --- | --- |
| **Antibiotic** | **Drug sensitivity** | **MIC** |
| Ampicillin | R | > 32 |
| Cefazolin | S | 2 |
| Cefoxitin | S | ≤ 2 |
| Cefuroxime | S | 4 |
| Cefotaxime | S | ≤ 0.5 |
| Ceftazidime | S | ≤ 1 |
| Cefepime | S | ≤ 1 |
| Imipenem | S | ≤ 0.5 |
| Piperacillin/tazobactam | I | 32 |

Results from the Third Affiliated Hospital of Sun Yat-Sen University. Report date: 3 d after death. MIC: Minimum inhibitory concentration; R: Resistant; S: Sensitive; I: Intermediate.



**Figure 1 CT tomography scan at admission and 10 d later.** A, B: Axial CT images: lung window setting (A) and mediastinal window setting (B) (slice thickness 3 mm) showed a 25 mm × 27 mm, round, thick-walled cavity in the right upper lobe. C, D: Axial CT images: Lung window setting (C), and mediastinal window setting (D) (slice thickness 3 mm) showed that the thick-walled cavity had been absorbed to a 21 mm × 21 mm, thin-walled cavity after treatment for 10 d. CT: Computed tomography.



**Figure 2 Body temperature and antibiotic use during hospitalization.** Antibiotic use: imipenem-cilastatin sodium 2 g every 8 hr (days 1-11) and 1 g every 6 hr (days 12-16), tigecycline 0.1 g every 12 hr (days 4-16), voriconazole 0.2 g every 12 hr (days 1-11 and 12-16), meropenem 2 g every 8 hr + 5-fluorocytosine 1.5 g every 6 hr + fluconazole 0.4 g every 12 hr (days 16-18).



**Figure 3 Change in white blood cell count and neutrophil predominance during hospitalization.**



**Figure 4 Change in C-creative protein and procalcitonin during hospitalization.**



**Figure 5 Axial unenhanced computed tomography scan of the head.** A: Axial unenhanced computed tomography scan of the head showed marked dilatation of the occipital horn of the lateral ventricles and third ventricles; B: Marked dilatation of the bilateral ventricle.