Hypervirulent Klebsiella pneumoniae

The Harvard community has made this article openly available. Please share how this access benefits you. Your story matters

Citation

Published Version
doi:10.1093/ofid/ofu028

Citable link
http://nrs.harvard.edu/urn-3:HUL.InstRepos:14351081

Terms of Use
This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA
Hypervirulent Klebsiella pneumoniae

Payal K. Patel,1 Thomas A. Russo,2,3,4 and Adolf W. Karchmer1

1Division of Infectious Diseases, Department of Internal Medicine, Beth Israel Deaconess Medical Center, Harvard Medical School, Boston, Massachusetts; 2Veterans Administration Western New York Healthcare System, Buffalo, New York; Departments of 3Medicine, and 4Microbiology and Immunology, University at Buffalo-State University of New York, Buffalo, New York

Hypervirulent strains of Klebsiella pneumoniae are associated with abscess formation, commonly hepatic, and metastatic spread, even in healthy patients. We describe a case of this clinical syndrome, genotypic and phenotypic features of the isolate, and briefly review epidemiology, clinical manifestations, and pathogenesis of this underappreciated syndrome.

Keywords. hypermucoviscous; hypervirulence; Klebsiella pneumoniae; virulence factors.

CASE REPORT

In November 2013, over the course of several weeks, a 74-year-old Vietnamese man with a history of hypertension and vitamin D deficiency developed subacute onset of abdominal pain, fevers, back pain, progressive weakness, and dyspnea on exertion. He was given empiric oseltamivir, which did not ameliorate his symptoms. He had previously been in a “re-education”/prison camp before moving to the United States in 1990. He last visited Ho Chi Minh City, Vietnam in 2011. At admission, he was febrile (103°F), tachycardic (103 beats per minute), dyspneic with conversation, and tender in the paramuscular area of the lumbar and thoracic spine. The white blood cell count was $13.7 \times 10^3$ cells/mm$^3$; aspartate aminotransferase was 113 IU/L; the alanine aminotransferase was 101 IU/L; total bilirubin was 2.3 mg/dL; and the inflammatory markers C-reactive protein and erythrocyte sedimentation rate were 242 mg/L and 94 mm/hour, respectively. Blood cultures positive for Klebsiella pneumoniae prompted initiation of ceftriaxone. Imaging revealed multiple lesions in the liver, an epidural abscess that had a ventral and dorsal component, and involved the cervical, thoracic, and lumbar spine with concern for osteomyelitis at L1-L3 and adjacent bilateral psoas abscesses (Figure 1). He had a normal ophthalmologic exam. The epidural abscess was evacuated surgically, and the largest hepatic abscess was drained percutaneously. Cultures from the liver, thoracic, and lumbar abscesses also grew K. pneumoniae. A string test, a semiquantitative phenotypic test, to assess for hypermucoviscosity by stretching a bacterial colony on an agar plate with an inoculation needle, was positive. This finding, in conjunction with the clinical syndrome, suggested that a hypervirulent isolate of K. pneumoniae was responsible. Genotypic analyses supported this contention. Polymerase chain reaction generated amplicons for iucA and iroN (biosynthetic genes for the siderophores aerobactin and salmochelin, respectively), terB (a tellurite resistance gene), and rmpA (a regulatory gene whose product mediates increased capsule production and hypermucoviscosity). These genes are present on large virulence plasmids that appear to be critical for the increased pathogenic potential of hypervirulent K. pneumoniae [1, 2]. The patient received 8 weeks of intravenous ceftriaxone; thereafter, spinal and abdominal imaging showed resolution of his liver, psoas, and epidural abscesses.

Epidemiology

In the mid-1980s, case reports from Taiwan described healthy patients with community-acquired K. pneumoniae liver abscesses and serious concomitant end-organ manifestations such as endophthalmitis and meningitis [3]. This pathotype has become known as “hypervirulent” to differentiate it from “classical” K. pneumoniae, which is commonly isolated from infected patients in Western countries [1]. In contrast to the usual healthcare-associated classical K. pneumoniae infections, hypervirulent K. pneumoniae can cause serious organ and life-threatening infections in younger, healthy individuals from the community. In the last decade, 813 cases of the invasive liver abscess syndromes associated with hypervirulent K. pneumoniae were reported [4]. This result is likely an underestimation due to underreporting, limited methods for differentiating hypervirulent from classical pathotypes, and unfamiliarity of this syndrome outside of Asia [4]. Klebsiella pneumoniae is now the most common cause of pyogenic liver abscess in Asia and possibly in North
Colonization rates of the serotypes K1 and K2, which are more likely to have the hypervirulence plasmid, are higher in Asia, likely leading to a higher prevalence of cases in the Pacific Rim, although there is still some question as to whether Asian hosts have an undetermined increased susceptibility to this infection [1, 7]. In addition to Asian ethnicity, some epidemiological studies have showed a higher incidence of disease at 55–60 years and a male dominance [4]. Diabetes mellitus has also been identified as a risk factor for this clinical syndrome [1, 4].

**Clinical Manifestations**

It is becoming increasingly clear that the initially described pyogenic liver abscess in the absence of biliary tract disease represents just one of many primary infections due to this organism. Other presentations include pneumonia, endophthalmitis, meningitis, nonhepatic abscess at a variety of sites, and necrotizing fasciitis [1]. These patients are commonly bacteremic. Metastatic spread from a site of infection is a defining, and potentially devastating, characteristic. Of 512 cases reviewed in Taiwan, the country with the highest prevalence of this syndrome, 15% of patients developed metastatic infection [4]. Other studies have shown up to 13% of patients with hypervirulent *K. pneumoniae* liver abscess developed concomitant central nervous system manifestations such as meningitis or endophthalmitis [8]. Metastatic infection most commonly involves the lung, causing pneumonia or empyema [1, 4]. Other sites of metastatic infection include bone and the genitourinary system, and there are case reports of endocarditis and a Lemierre syndrome variant thought to be secondary to hypervirulent *K. pneumoniae* [1].

**Virulence**

Phenotypically, hypervirulent *K. pneumoniae* is more resistant to complement and neutrophil-mediated bactericidal activity in vitro and virulent in vivo than classical strains [9]. Virulence is associated with the acquisition of a 200- to 220-kilobase plasmid that contains *rmpA* and siderophore biosynthetic genes. Increased capsule production, which is mediated at least in part by *RmpA*, enhances virulence in animal infection models. In contrast to classical *K. pneumoniae*, the hypervirulent isolates secrete greater amounts of siderophores that mediate iron acquisition from the host. This phenotype enhances growth and survival in human ascites ex vivo [10]. Other potential virulence factors for hypervirulent *K. pneumoniae* have been recently reviewed [1]. However, significant knowledge gaps exist, and undoubtedly numerous additional factors yet to be described will prove to be critical in the pathogenesis of infection. It remains unclear what features of hypervirulent *K. pneumoniae* enable metastatic spread. Furthermore, although intestinal colonization is almost certainly a prerequisite for disease, the portal of entry leading to infection and the mechanism by which this occurs is also unknown.

**DISCUSSION**

Our case demonstrates some classic features of infection due to hypervirulent *K. pneumoniae*. The infection was community-acquired and in a previously healthy patient. Metastatic spread, one of the defining features, was observed in the epidural space, spine, and psoas muscles. This clinical syndrome connotes important implications for management. Unfortunately, a rapid and reliable test to identify hypervirulent *K. pneumoniae* is still lacking. Although the string test is widely considered to be reliable for distinguishing hypervirulent from classical *K. pneumoniae* strains, its sensitivity is not optimal and a significant minority of classical *K. pneumoniae* strains are positive [11, 12]. These performance issues are especially problematic in regions of lower hypervirulent *K. pneumoniae* prevalence, such as the United States. Nonetheless, when suspected, a prompt ophthalmological exam should be performed because of the rapid course and high degree of morbidity associated with hypervirulent *K. pneumoniae* endophthalmitis [4, 8]. Vigilance for other metastatic manifestations should also be emphasized because there may be a need for source control as demonstrated by this case. Although diabetes mellitus has been identified as a risk factor, many patients with the syndrome have no

Figure 1. Magnetic resonance imaging scan performed upon admission demonstrated increased signal at the L1-L3 disc level and in the adjacent vertebral bodies concerning for discitis or osteomyelitis. A large anterior epidural abscess extends down from at least T10. Canal narrowing, which is highlighted, appears to be most severe at L4-L5 from disc bulge and epidural abscess.
underlying immunodeficiency or comorbidities. Strict glycemic control in diabetic patients was shown to decrease metastatic complications in one study [13].

Over the last few decades, mortality rates have declined with dual approach of percutaneous drainage and antibiotic therapy [4, 14]. However, percutaneous drainage can be challenging with hypervirulent *K. pneumoniae* due to its hypermucoviscous nature. Unlike other etiologies of hepatic abscess, hypervirulent *K. pneumoniae* liver abscess is typically monomicrobial [6]. Antimicrobial susceptibility patterns of hypervirulent *K. pneumoniae* remain largely pan-sensitive for now, usually only resistant to ampicillin [8]. Classical *K. pneumoniae*, however, has increasingly acquired genes for extended spectrum β-lactamases and carbapenemases.

Treatment of these infections with a third-generation cephalosporin is preferred, although there is some debate as to whether first-generation cephalosporins could be equally effective [4, 6, 14]. Duration of therapy has not been established in controlled trials. In general, a period of 2–6 weeks has been used, with a lengthier duration used for larger abscesses that resolve slowly or in the presence of metastatic disease that requires a more prolonged duration [4]. A conservative approach would be to treat until radiographic evidence of infection is resolved. However, even with this approach, anecdotal reports of recurrences have been described [1].

In summary, infection caused by hypervirulent *K. pneumoniae* is associated with notable morbidity and potential mortality. To provide optimal care and minimize these untoward consequences requires prompt recognition of this pathogen. This distinction would facilitate attention to key aspects of treatment such as control of hyperglycemia in patients with diabetes mellitus and aggressively looking for metastatic spread of infection. Although progress has been made in understanding the virulence factors driving this clinical syndrome, rapid diagnostic tests and an increased understanding of the epidemiology of this unique pathogen are still needed. These needs are becoming more pressing with recent disconcerting reports describing acquisition of multidrug resistance by hypervirulent *K. pneumoniae* isolates [15].

**Acknowledgments**

**Financial support.** This work was supported in part by the National Institutes of Health Grant 1R21AI088318-01A1 (to T. A. R.).

**Potential conflicts of interest.** All authors: No reported conflicts. All authors have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest.

**References**