

Available online at www.sciencedirect.com

ScienceDirect

journal homepage: www.e-jmii.com

CORRESPONDENCE

Moraxella osloensis bacteremia complicating with severe pneumonia in a patient with lung cancer

Dear Editor,

Moraxella osloensis is an aerobic, Gram-negative coccobacillus (formerly classified as *Moraxella nonliquefaciens*). The clinical significance and appropriate therapy for patients with *M. osloensis* bacteremia are not well studied. Here, we report a case of a male patient with lung cancer who presented with *M. osloensis* bacteremia complicating with pneumonia. The species was initially identified as *Comamonas testosteroni* using a commercially automated system, and was reidentified as *M. osloensis* using Bruker Biotyper matrix-assisted laser desorption ionization—at the time of flight mass spectrometry (MALDI-TOF MS; Bruker Daltonics, GmbH, Bremen, Germany) and partial 16S ribosomal DNA gene sequencing analysis.

The patient received a diagnosis of lung cancer of the right upper lobe 4 years ago. He suffered from fever and cough for 2 days before admission, and a chest radiograph showed infiltrations of the right lower lobe of the lung (Figure 1). He used a bilateral positive airway pressure ventilator for pneumonia with acute respiratory failure. The laboratory data revealed the results of white blood cells 9070 cells/mL with 83% neutrophils, hemoglobin 11.9 mg/dL, platelets 242,000 cells/mL, serum glucose 254 mg/dL, creatinine 2.34 mg/dL, blood urea nitrogen 18 mg/dL, and C-reactive protein 19.8 mg/dL. The patient received empirical therapy with intravenous cefoperazone/sulbactam 2 g/2 g every 12 hours initially. Four days later, the results of blood culture were identified as *C. testosteroni* using the Phoenix 100 ID/AST system (Becton Dickinson, Sparks, MD, USA). The isolate was presumptively identified as *M. osloensis* (score value 1.848) using Bruker Biotyper MALDI-TOF (Bruker Daltonics). The result of Gram stain of the colonies showed coccobacilli. Sequencing analysis of partial 16S ribosomal DNA gene of the isolate was performed with two primers: 8FPL (5'-AGAGTTTGATCCTGGCTCAG-3') and 1492 (5'-GGTTACCTG

TTACGACTT-3'). The sequences obtained were compared with published sequences in the GenBank database, using the BLASTN algorithm (<http://www.ncbi.nlm.nih.gov/blast>). The isolate was identified as *M. osloensis* (accession number KC456542.1) with maximal identity of 100% (858/858 bp).

The antimicrobial susceptibility test showed that the isolate was resistant to cefazolin, cephalexin, penicillin, ampicillin, and amoxicillin/clavulanate, but susceptible to levofloxacin, ciprofloxacin, ceftazidime, cefepime, cefoperazone/sulbactam, amikacin, gentamicin, and carbapenems (doripenem, imipenem, and meropenem) using the Phoenix 100 ID/AST system (Becton Dickinson). The patient was discharged on hospital Day 21 to complete a 14-day course of cefoperazone/sulbactam therapy.

Among the *Moraxella* species, *M. catarrhalis* is the most common isolate of the respiratory tract. *M. osloensis* has been isolated from the respiratory tract, blood, cerebrospinal fluid, and urine.^{1,3} Clinical reports of infections caused by *M. osloensis* are rare.¹ *M. osloensis* has been implicated in community-acquired pharyngitis, septic arthritis, meningitis, ophthalmitis, and otitis media.^{1–5} Hospital-acquired central venous catheter infections in an elderly woman receiving chronic parenteral nutrition have also been reported.^{1,6} Our patient with *M. osloensis* bacteremia complicating with pneumonia is different from previous case reports. *M. osloensis* is often misidentified as *C. testosterone*,² and molecular methods may be required to confirm the species identification. The appropriate treatment of bacteremia due to *M. osloensis* has not been well studied. In past decades, *M. osloensis* has been found to be sensitive to penicillin, first-generation cephalosporins, and aminoglycosides.⁵ Although, our patient had been resistant to penicillin, amoxicillin/clavulanate, and cefazolin. However, the prognosis for patients with *M. osloensis* infections is generally good.^{1–5} Further studies are still

<http://dx.doi.org/10.1016/j.jmii.2015.03.005>

1684-1182/Copyright © 2015, Taiwan Society of Microbiology. Published by Elsevier Taiwan LLC. All rights reserved.

Please cite this article in press as: Lee W-S, et al., *Moraxella osloensis* bacteremia complicating with severe pneumonia in a patient with lung cancer, Journal of Microbiology, Immunology and Infection (2015), <http://dx.doi.org/10.1016/j.jmii.2015.03.005>

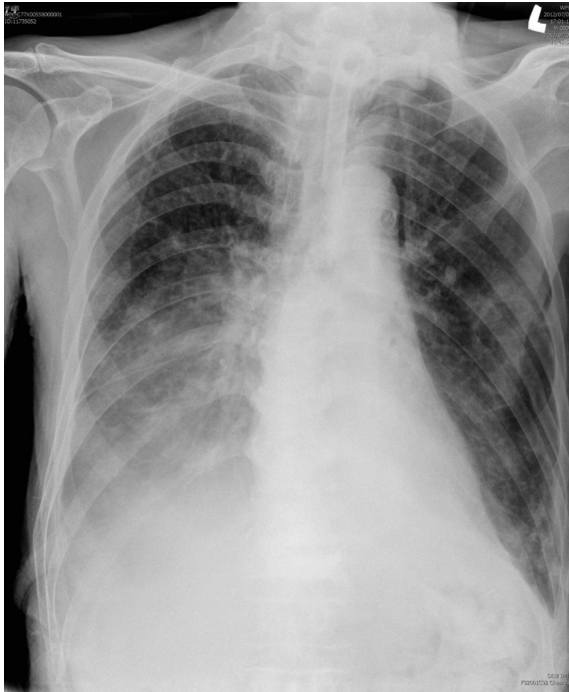


Figure 1. Chest radiograph revealing the infiltration patch of the right lower lobe of the lung.

required to evaluate the clinical manifestations, diagnosis, and antibiotic resistance of infections due to *M. osloensis*.

Conflicts of interest

All contributing authors declare no conflicts of interest.

References

1. Shah SS, Ruth A, Coffin SE. Infection due to *Moraxella osloensis*: case report and review of the literature. *Clin Infect Dis* 2000;30:179–81.
2. Tsui TL, Tsao SM, Liu KS, Chen TY, Wang YL, Teng YH, et al. *Comamonas testosteroni* infection in Taiwan: reported two cases and literature review. *J Microbiol Immunol Infect* 2011;44:67–71.
3. Berrocal AM, Scott IU, Miller D, Flynn Jr HW. Endophthalmitis caused by *Moraxella* species. *Am J Ophthalmol* 2001;132:788–90.

4. Han XY, Tarrand JJ. *Moraxella osloensis* blood and catheter infections during anticancer chemotherapy: clinical and microbiologic studies of 10 cases. *Am J Clin Pathol* 2004;121:581–7.
5. Hsu SF, Lin YT, Chen TL, Sui LK, Hsueh PR, Huang ST, et al. Antimicrobial resistance of *Moraxella catarrhalis* isolates in Taiwan. *J Microbiol Immunol Infect* 2012;45:134–40.
6. Tseng SH, Chien LJ, Chang FY. National action plan to eliminate central line-associated bloodstream infections in Taiwan. *J Microbiol Immunol Infect* 2014;47:265–7.

Wen-Sen Lee

Division of Infectious Diseases, Department of Internal Medicine, Wan Fang Medical Center, Taipei Medical University, Taipei, Taiwan

Department of Internal Medicine, School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan

Po-Ren Hsueh

Department of Laboratory Medicine, National Taiwan University Hospital, Taipei, Taiwan

Department of Internal Medicine, College of Medicine, National Taiwan University Hospital, Taipei, Taiwan

Fang-Lan Yu

Department of Laboratory Medicine, Wan Fang Medical Center, Taipei Medical University, Taipei, Taiwan

Department of Internal Medicine, School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan

Fu-Lun Chen

Tai-Chin Hsieh

Tsong-Yih Ou*

Division of Infectious Diseases, Department of Internal Medicine, Wan Fang Medical Center, Taipei Medical University, Taipei, Taiwan

Department of Internal Medicine, School of Medicine, College of Medicine, Taipei Medical University, Taipei, Taiwan

*Corresponding author. Number 111, Section 3, Hsing Long Road, Taipei 116, Taiwan.

E-mail address: 89425@wanfang.gov.tw (T.-Y. Ou)

18 December 2014

Available online ■ ■ ■