Sacroiliitis and osteomyelitis caused by serotype 3 Streptococcus pneumoniae in a previously healthy adult: a case report

Yoshiro Hadano1
Bin Chang2

1Department of Infectious Diseases, St. Mary’s Hospital, Kurume, Fukuoka, Japan; 2Department of Bacteriology I, National Institute of Infectious Diseases, Shinjuku-ku, Tokyo, Japan

Abstract: Streptococcus pneumoniae is a rare cause of pyogenic sacroiliitis and osteomyelitis. A case of pneumococcal sacroiliitis and osteomyelitis is reported herein. The patient was a previously healthy 54-year-old man with a 3-day history of fever and a 3-week history of increasing right-side thigh pain. Computed tomography showed bone destruction of the right sacroiliac joint with perifocal abscess formation in the right iliac muscle. The patient was diagnosed with pyogenic sacroiliitis and osteomyelitis with a right iliac muscle abscess. Mucoid serotype 3 S. pneumoniae was obtained by culture of abscess. The patient was successfully treated with an 8-week course of antibiotics.

Keywords: Streptococcus pneumoniae, pyogenic sacroiliitis, osteomyelitis, serotype 3, invasive pneumococcal diseases

Introduction

Streptococcus pneumoniae is commonly found in nasopharyngeal mucosa of healthy people and can cause various diseases, such as pneumonia and meningitis. The bacterium is a relatively uncommon cause of septic arthritis, including pyogenic sacroiliitis, which is a rare manifestation of invasive pneumococcal diseases (IPD). The prognosis of IPD is influenced by both bacterial factors, such as serotype, and host characteristics, such as age, chronic cardiovascular and pulmonary diseases, diabetes mellitus, alcoholism, smoking, and immunosuppressive states. Because serotype 3 S. pneumoniae have a thick capsule, it is thought that it is highly pathogenic. In this report, we describe a rare case of pyogenic sacroiliitis and osteomyelitis with a right iliac muscle abscess caused by serotype 3 S. pneumoniae in a healthy adult.

Case report

A previously healthy 54-year-old Japanese man was admitted to our hospital with a 3-week history of worsening right-side thigh pain and a 3-day history of fever. Four weeks prior to admission, the patient had a mild cough with sputum. His symptoms improved without any treatment. Three weeks prior to admission, he developed mild right thigh pain without trauma that did not interfere with his daily life. One week prior to admission, he came to a local clinic because of his right thigh pain, and loxoprofen was prescribed. His symptoms transiently improved with loxoprofen, but 3 days prior to admission, the patient had chills and his thigh pain worsened. He returned to the local clinic, and amoxicillin/clavulanate was prescribed. On the day of admission, he visited another community hospital and was transferred to our hospital.
hospital on the diagnosis of right iliac muscle abscess. He denied sore throat, cough, dyspnea, and chest pain. His past history was not remarkable. The patient had no previous medical history, no use of tobacco or illicit drug, and rarely drink alcohol. He denied recent animal exposure, recent travel, or any history of allergies. On physical examination, his blood pressure was 140/60 mmHg, pulse rate was 66 beats per minute, temperature was 38.0°C, and respiratory rate was 20 breaths per minute with oxygen saturation of 97% on room air. The results of the physical examination were unremarkable, except for right buttock tenderness and positive right psoas sign. Laboratory data obtained on admission revealed a white blood cell count of 19,070 μL with 94% neutrophils. Examination of the serum chemistry revealed the following results: blood urea nitrogen 23.9 mg/dL, creatinine 1.6 mg/dL, albumin 2.2 g/dL, total protein 6.6 g/dL, and C-reactive protein 26.0 mg/dL. Based on these findings, the patient was suspected of having right psoas abscess. CT of the pelvic area showed bone destruction of the right sacroiliac joint with perifocal abscess formation in the right iliac muscle (Figure 1A). A diagnostic CT-guided percutaneous drainage of the abscess was performed, and Gram staining of the pus revealed Gram-positive diplococcus bacteria and polymorphonuclear leukocytes (Figure 2).

After an initial work up in the emergency room, his primary diagnosis was bacterial sacroiliitis and osteomyelitis with right iliac muscle abscess. Transthoracic echocardiography was performed, and the results showed no evidence of an infection. The patient was started on intravenous cefmetazole at a dose of 2 g every 12 hours. No bacterium was detected from his blood culture, but the pus culture grew S. pneumoniae. Additional examination revealed the following results: 90 mm/h erythrocyte sedimentation rate, 56 mg/dL IgM (within normal range 33–183 mg/dL), 1450 mg/dL IgG (within normal range 861–1,747 mg/dL), 148 mg/dL C3 (within normal range 73–138 mg/dL), 40 mg/dL C4 (within normal range 11–31 mg/dL), and 70 mg/dL CH50 (within normal range 25–48 mg/dL). Human immunodeficiency virus antibody, M protein, and Bence Jones protein were negative. CT of the abdomen showed no evidence of any abdominal finding including spleen. Serum protein fraction came back with an acute inflammation pattern without M protein. Mucoid S. pneumoniae was obtained by culturing of abscess sample. Serotyping of S. pneumoniae was performed by the capsular swelling method using pneumococcal antiserum (Statens Serum Institute, Copenhagen, Denmark) and the stain was determined to be serotype 3. Susceptibility of S. pneumoniae to antibiotics was analyzed using...
the broth microdilution method according to a protocol of the Clinical and Laboratory Standards Institute. Minimal inhibitory concentration breakpoints were defined according to the Clinical and Laboratory Standards Institute (M100) criteria. The S. pneumoniae strain exhibited susceptibility to penicillin, ampicillin, ceftriaxone, meropenem, tosufloxacin, vancomycin, and trimethoprim/sulfamethoxazole, but was resistant to clindamycin (Table 1). On day 4, magnetic resonance imaging of the pelvic area revealed increased signal intensity of the right sacroiliac joint and surrounding bone, and diffuse thickening and effusion surrounding the muscle (Figure 1B). On day 5, we attempted to perform de-escalation. The patient was switched to treatment with ampicillin at a dose of 2 g every 6 hours on the basis of the susceptibility test because his renal function had improved. He was given levofloxacin 500 mg orally because of the drug eruption due to ampicillin on day 25. The patient was successfully treated with an 8-week course of antibiotics. No evidence of a relapse of the infection was noted at the 6-month follow-up.

**Ethics statement**

Written informed consent has been obtained from the patient to obtain their results and to publish the case details with accompanying images.

**Discussion**

Pyogenic sacroiliitis is a rare infectious diseases, representing 1%–2% of all cases of septic arthritis, most frequently in children and young adults. The risk factors for pyogenic sacroiliitis include being intravenous drug user, having infections in other organs systems, such as urinary tract infection, and pregnancy. According to previous reports, adult patients are more likely to have atypical presentation,
concurrent infections, and local complication. Delay in the diagnosis can lead to additional complication such as osteomyelitis, abscess formation, and bacteremia. Based on previous studies, bacteremia is documented in 71% of adults with pneumococcal septic arthritis. In this case, we believe that the cause of the sacroiliitis, osteomyelitis, and iliopsoas abscess was the secondary infection as a result of primary bacteremia. Although the results of the blood cultures were negative, this patient had chills, and amoxicillin/clavulanate were prescribed before admission to our hospital. Treatment options include antibiotic therapy alone, CT-guided drainage, or surgical drainage. The duration of antimicrobial treatment is at least 4–8 weeks. A review of pneumococcal septic arthritis showed 2.8% of adult patients with pneumococcal septic arthritis have sacroiliac involvement. In addition, three cases of pneumococcal sacroiliacitis in adult have been reported in the literature (Table 2). These cases include a 31-year-old woman who presented with a fever, depressed level of consciousness, and right buttock and thigh pain, which was later proven to be caused due to *S. pneumoniae* by blood cultures; a 47-year-old woman who presented with a 4-week history of left-sided lower back pain which was proven to be caused due to *S. pneumoniae* by blood cultures; and a 62-year-old who presented with a 3-day history of right buttock pain which was proven to be caused due to *S. pneumoniae* by blood cultures, muscle biopsy, and surgical drainage. In the present case, the patient did not have an immunodeficiency or chronic illness. Interestingly, all of the patients had no risk factor for IPD. *S. pneumoniae* is covered by a capsule, which is a very important factor for its pathogenicity. In the case presented herein, serotype 3 *S. pneumoniae* was isolated from the lesion and determined as the causative pathogen. Serotype 3 *S. pneumoniae* strains always have a characteristically thicker capsule, show mucoid formation on blood agar plates, possess greater virulence, and have a higher mortality rate when compared to other serotype strains. In the present case, the treatment duration was 8 weeks with no evidence of a relapse of the infection during the 6-month follow-up. We believe an adequate treatment duration leads to successful treatment, even in the case of serotype 3 infection.

**Conclusion**

In conclusion, we reported a case of serotype 3 pneumococcal sacroiliitis and osteomyelitis with right iliac muscle abscess in a healthy adult. It is important to keep in mind that bacterial sacroiliitis and osteomyelitis due to *S. pneumoniae* are rare, but can occur even in a previously healthy adult. Delay in the diagnosis can lead to additional complication such as osteomyelitis, abscess formation, and bacteremia.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**