A case of necrotizing fasciitis due to *Streptococcus pneumoniae* serotype 5 in Saskatchewan

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Necrotizing fasciitis due to *Streptococcus pneumoniae* is a rare and grave condition, and only a few cases have been reported. Suggested risk factors include minor trauma, systemic lupus erythematosus, immunosuppression secondary to medication, use of intramuscular anti-inflammatories and alcoholism. A fatal case of pneumococcal necrotizing fasciitis that occurred in a 51-year-old woman with a history of alcohol abuse and oral anti-inflammatory use is presented. Her condition was caused by a multi-etiology outbreak of community-acquired pneumonia, from which *S pneumoniae* serotype 5 was also isolated. The case description outlines the subtle presentation and rapid clinical progression of this condition. Because serotype 5 antigen is included in the polysaccharide 23-valent pneumococcal vaccine, the present case highlights the importance of pneumococcal immunization programs in Canada.

**Key Words:** Aboriginal; Canada; Necrotizing fasciitis; *Streptococcus pneumoniae*

**CASE PRESENTATION**

A 51-year-old Aboriginal woman from northern Saskatchewan presented to a local family medical clinic in early October 2006 with a three-day history of left knee pain. Her vital signs included a blood pressure of 114/70 mmHg, a heart rate of 100 beats/min and a respiratory rate of 20 breaths/min. Her temperature was not documented. On examination, her knee was warm and painful, and an effusion was noted. Approximately six weeks prior, she had fallen on her right knee while walking. This injury was complicated by hemarthrosis and effusion, requiring needle drainage on two occasions. In addition, she had a history of pain, swelling and erythema involving her shoulder joint. Her past history was significant for alcohol abuse and unstable social and housing conditions. The laboratory results showed the following – WBC 9.8×10⁹/L; Hb level 111 g/L; platelet count 171×10⁹/L; sodium level 131 mmol/L (normal 137 mmol/L to 145 mmol/L); potassium level 3.2 mmol/L (normal 3.6 mmol/L to 5.0 mmol/L); chloride level 91 mmol/L (normal 98 mmol/L to 107 mmol/L); sodium level 131 mmol/L (normal 137 mmol/L to 145 mmol/L); creatine kinase level 484 U/L (normal 30 U/L to 107 mmol/L); carbon dioxide 16 mmol/L (normal 22 mmol/L to 30 mmol/L); urea level 21.9 mmol/L (normal 2.5 mmol/L to 6.1 mmol/L); aspartate aminotransferase level 268 U/L (normal 8 U/L to 39 U/L); creatine kinase level 484 U/L (normal 30 U/L to 107 mmol/L); and granulocyte count 8.8×10⁹/L (normal 2×10⁹/L to 10×10⁹/L). A presumptive diagnosis of inflammatory arthritis was made, and she was given indomethacin 50 mg three times a day for her symptoms.

Two days later, the patient became progressively more confused, disoriented and unresponsive to questions. She was brought by ambulance to the local emergency department where her temperature was 38.8°C, pulse 98 beats/min, blood pressure 140/83 mmHg and respiratory rate 32 breaths/min. Her Glasgow coma scale score was 6. She was unresponsive to verbal commands but responsive to painful stimuli. Bruising was noted on both legs, and a large area of erythema was noted around the left knee. Her respiratory examination was unremarkable. Laboratory results showed the following – WBC count 3.7×10⁹/L; Hb level 111 g/L; platelet count 171×10⁹/L; sodium level 131 mmol/L (normal 137 mmol/L to 145 mmol/L); potassium level 3.2 mmol/L (normal 3.6 mmol/L to 5.0 mmol/L); chloride level 91 mmol/L (normal 98 mmol/L to 107 mmol/L); carbon dioxide 16 mmol/L (normal 22 mmol/L to 30 mmol/L); urea level 21.9 mmol/L (normal 2.5 mmol/L to 6.1 mmol/L); and creatine kinase level 484 U/L (normal 30 U/L to 107 mmol/L). Oral analgesics were given and the patient was admitted to the intensive care unit. A presumptive diagnosis of sepsis was made, and she was started on a combination of antibiotics.

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Saskatoon, Saskatchewan

Tan, Division of Infectious Diseases, Royal University Hospital, pneumonia in the absence of respiratory symptoms. Figure by Dr Ben necrotizing fasciitis, toxic shock syndrome and community-acquired pneumonia in children younger than 10 years of age. These IPD cases caused outbreaks of invasive pneumococcal disease (IPD) in Alberta, British Columbia, Manitoba and Saskatchewan. Urban outbreaks in British Columbia and Alberta have involved inner-city, adult populations with risk factors for IPD – alcoholism, illicit drug use, history of hepatitis B or C and living in a housing shelter (5, 6). Most individuals have severe disease – pneumonia with sepsis, with many requiring intensive care unit admission (5, 6). Canadian Aboriginal people have a higher risk of IPD (7), and a higher risk of serotype 5 disease (OR 3.0, 95% CI 1.3 to 6.6) (5).

S. pneumoniae serotype 5 is a common pneumococcal serotype in Africa and India (4). Since 2005, this strain has been identified with increasing frequency in Canada and has caused outbreaks of invasive pneumococcal disease (IPD) in Alberta, British Columbia, Manitoba and Saskatchewan. Urban outbreaks in British Columbia and Alberta have involved inner-city, adult populations with risk factors for IPD – alcoholism, illicit drug use, history of hepatitis B or C and living in a housing shelter (5, 6). Most individuals have severe disease – pneumonia with sepsis, with many requiring intensive care unit admission (5, 6). Canadian Aboriginal people have a higher risk of IPD (7), and a higher risk of serotype 5 disease (OR 3.0, 95% CI 1.3 to 6.6) (5).

During October 2006, nine cases of IPD due to serotype 5. S. pneumoniae were identified in an isolated community (population 3500) in northern Saskatchewan. Four IPD cases were in children younger than 10 years of age. These IPD cases formed part of a larger outbreak of multi-etiology community-acquired pneumonia, which included other agents such as Chlamydia pneumoniae, Mycoplasma pneumoniae and/or respiratory syncytial virus. The present case of NF is part of this outbreak.

S. pneumoniae is a rare case of NF, with approximately 12 cases documented in the literature (8-15). These case reports suggest the following risk factors for the development of pneumococcal NF – a history of diabetes, systemic lupus erythematosus, immunosuppression, alcohol use, coronary artery disease and administration of intramuscular nonsteroidal anti-inflammatory drugs (NSAIDs) (8,9,11-14). Only two cases of pneumococcal NF have been reported in healthy individuals who sustained minor trauma. One of these individuals used topical anti-inflammatory medication over the site of her injury.

150 × 110 g/L to 160 g/L); platelet count 144 × 10^9/L on admission showed the following – WBC count 3.2 × 10^9/L; Hb level 99 g/L (normal 110 g/L to 160 g/L); platelet count 144 × 10^9/L (normal 150 × 10^9/L to 400 × 10^9/L); creatine kinase level 894 U/L (normal 30 U/L to 200 U/L); creatine kinase isoenzyme – MB level 24 U/L (normal 0 U/L to 15 U/L); alkaline phosphatase level 53 U/L (normal 30 U/L to 110 U/L); alanine aminotransferase level 186 U/L (normal 10 U/L to 40 U/L); gamma glutamyl transferase level 80 U/L (normal 10 U/L to 35 U/L); sodium level 137 mmol/L (normal 135 mmol/L to 146 mmol/L); potassium level 3.3 mmol/L (normal 3.5 mmol/L to 5.1 mmol/L); chloride level 106 mmol/L (normal 100 mmol/L to 110 mmol/L); carbon dioxide 18 mmol/L (normal 22 mmol/L to 31 mmol/L; creatinine level 287 μmol/L (normal 45 μmol/L to 110 μmol/L); urea level 22.1 mmol/L (normal 3.7 mmol/L to 7.0 mmol/L); international normalized ratio 1.1 (normal 0.8 to 1.2) and partial thromboplastin time 29 s (normal 26 s to 36 s). Her chest x-ray demonstrated the presence of a large, right middle lobe infiltrate (Figure 1). Her knee aspiration yielded purulent fluid.

A presumptive diagnosis of necrotizing fasciitis (NF) of the left knee with toxic shock syndrome and pneumonia was made. Intravenous ceftriaxone (2 g every 12 h) and clindamycin (900 mg every 6 h) were administered.

She was immediately taken to the operating room for debridement of the leg, but due to the extent of involvement, she required left knee arthrothomy, followed by hip disarticulation and exploratory laparotomy. A frozen section of the iliac fascia revealed microscopic necrosis, but no bacteria were noted. Due to her worsening condition, a second laparotomy with anticipated debridement of the posterior abdomen and iliofascia was considered, but could not be performed because of her family's decision to withdraw care. The patient died on the second day of hospitalization.

Gram stain of pus from her knee revealed Gram-positive cocci. Streptococcus pneumoniae (serotype 5) was isolated from knee fluid, and blood and tissue cultures. The isolate was sensitive to penicillin, cefotaxime, clindamycin, fluoroquinolones and vancomycin, but was resistant to co-trimoxazole.

DISCUSSION

S. pneumoniae is a Gram-positive diplococcus. There are 90 known serotypes, of which the top 10 account for over 60% of infections worldwide (1, 2). Pneumococci can cause conjunctivitis, otitis media, community-acquired pneumonia and invasive infections such as sepsis, meningitis and soft tissue infections, among others. Transmission is primarily from person to person through respiratory droplets. People at both ends of the age spectrum are at a high risk of pneumococcal disease. Other risk factors include crowding, exposure to smoke, congenital or acquired immune deficiency, asplenia and a history of cochlear implants (3).

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which resulted in pruritus, scratching and subsequent excoriations. Whether the topical NSAID facilitated the progression of infection is unknown. NSAIDs are potent inhibitors of granulocyte function. In this case, the patient had used oral NSAIDs, which may have contributed to the accelerated course of illness; alternatively, they may have masked the symptoms, allowing her illness to progress (16).

Two different presentations of *S. pneumoniae* NF have been identified – cellulitis of the upper body (face, neck and torso), usually accompanied by autoimmune (systemic lupus erythematosus) or hematological disorders, and fasciitis, predominantly involving the limbs and usually in association with coexisting diabetes mellitus or chronic alcohol use (14). While an antecedent history of trauma can be present in 75% of NF cases due to group A streptococcus, this feature is usually absent in NF due to *S. pneumoniae* (14). Pneumococcal serotypes implicated in NF include 6A, 9, 9V, 10A and 14. Our case report is the first documentation of NF due to serotype 5.

Although *S. pneumoniae* serotype 5 is not included in the seven-valent conjugate vaccine, it is included in the polysaccharide 23-valent vaccine (PPV23). Individuals eligible for the publicly funded PPV23 may vary by province, but in general, the National Advisory Committee on Immunization (3) recommends PPV23 for the following groups – adults 65 years and older, and individuals five to 64 years of age with chronic cardiorespiratory disease, cirrhosis, alcoholism, chronic renal disease, nephritic syndrome, diabetes mellitus, chronic cerebrospinal fluid leak, sickle cell disease, functional or anatomical asplenia, HIV infection and other conditions associated with immunosuppression. Given the history of alcohol abuse in the 51-year-old woman, the present case represents a missed opportunity for vaccination. Individuals in this high-risk group are difficult to reach for routine immunization programs. We have thus identified a fatal case of *S. pneumoniae* serotype 5 representing a group of people whose risk for disease also decreases their chances of accessing vaccination. Additional approaches to increasing PPV23 immunization coverage in this high-risk, hard-to-access group need to be considered.

## CONCLUSION

The present case highlights the difficulty in early diagnosis of NF due to the subtle signs at illness onset and the absence of leukocytosis. The use of oral anti-inflammatory drugs commonly prescribed for noninfectious arthritides may contribute to the severity of infection or mask the symptoms. While PPV23 has limited efficacy against prevention of invasive disease in people with diabetes and alcoholism (17), the final outcome might have been altered if this individual had been immunized. Finally, the emergence of *S. pneumoniae* serotype 5 in Canada may be accompanied by severe disease presentations in individuals at high risk of pneumococcal infections.

### ACKNOWLEDGEMENTS

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### REFERENCES
