Streptococcus pneumoniae as a Cause of Salpingitis

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ABSTRACT

Background: A case of pneumococcal septicemia associated with laparoscopically documented acute salpingitis is reported.
Case: Gram-stained cul-de-sac pus revealed gram-positive encapsulated diplococci.
Conclusion: This case coupled with reanalysis of prior genital tract involvement in nonpregnant individuals argues that Streptococcus pneumoniae can mimic gonococcal diseases.

KEY WORDS
Pneumococcal septicemia, acute salpingitis, etiology

Streptococcus pneumoniae is not a normal constituent of vaginal bacterial flora12; however, given the opportunity, S. pneumoniae can be a significant pathogen for the female genital tract.2–6

Classically, involvement with the female genital tract has occurred in conjunction with primary pulmonary disease, engenderment of bacteriuriaemia, and subsequent metastatic involvement.7,8 More recently, there is the growing perception that S. pneumoniae, when present within the vaginal flora at the time of parturition or rupture of the fetal membranes, has the ability to ascend and infect the amniotic sac and secondarily the fetus.2–6

Genital tract disease due to S. pneumoniae has been reported in nonpregnant females. Isolated cases of spontaneous pneumococcal peritonitis have been described.9,10 Hadfield et al.11 reported a case of a 46-year-old woman with bilateral tubo-ovarian masses. Biopsy specimens from both tubes and from the wall of the abscesses demonstrated gram-positive, lancet-shaped diplococci which were documented to be S. pneumoniae by immunoperoxidase staining. Rahav et al.12 reported a case of postmenopausal pneumococcal tubo-ovarian abscess from which S. pneumoniae was recovered.

The purpose of this paper is to report a case of pneumococcal salpingitis documented by laparoscopy and comment on its significance in terms of the pathogenetic spectrum attributable to S. pneumoniae as it involves the female genital tract.

CASE REPORT

Initially, the patient (age 24) was transferred from jail to the emergency room because of severe abdominal cramping. The diagnosis of systemic lupus erythematosus had been previously documented, resulting in the patient's being placed on prednisone, 5 mg/day. The pelvic examination performed at that time was normal and she was diagnosed as having viral gastroenteritis. Two days later, because of a suicidal gesture involving excess self-administration of the antidepressant medication fluoxetine hydrochloride (Prozac), the patient was brought back to the hospital and admitted. At that time, she presented as a well-nourished, well-developed black female in no acute distress; however, vital signs included a temperature elevation of 39.8°C, pulse 116, respiration 16, and blood pressure 104/60. Pertinent physical findings included areas of spotty hyperpigmentation of the
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forehead and malar area, forearms, hands, and feet consistent with discoid lupus. An abdominal examination was within normal limits. The pelvic examination was deferred. The chest X-ray, sinus films, and a computed tomography (CT) of the head were all normal. Laboratory evaluation revealed a white blood cell (WBC) count of 2,100 with a marked shift to the left, hemoglobin 9.8 g, hematocrit 29%, and platelets 90,000. Her electrolytes, liver function test, and arterial blood gases were within normal limits. Her drug screen was positive for acetaminophen. A fluorescent antinuclear antibody test (FNA) was positive with an antinuclear antibody (ANA) titer of 1/1,280 and an antinative DNA titer of 1/80. The C3, C4, and C5 fractions of complement were low. The patient was placed under observation, but she continued to run a febrile course. Blood cultures, urine cultures, and throat cultures done at the time of admission were negative.

On the 3rd hospital day, the patient developed a sudden onset of severe cramping lower quadrant pain. An abdominal examination revealed diffuse direct and rebound tenderness. Cervical motion tenderness was demonstrated on pelvic examination. Cultures for Neisseria gonorrhoeae, Chlamydia trachomatis, and wet mount preparations for Trichomonas vaginalis were negative. Serological testing for human immunodeficiency virus (HIV)-1 and -2 was negative. The venereal disease research laboratory (VDRL) was nonreactive. Consultation was obtained from the Department of Obstetrics and Gynecology. At the time of this examination, the patient had a temperature elevation of 40.0°C, pulse 127, respiration 20, and blood pressure 118/50. Physical examination revealed four-quadrant tenderness to direct palpation and rebound tenderness. The patient refused a pelvic examination. All blood cultures drawn grew out S. pneumoniae. A pelvic ultrasound revealed a 5 × 5 × 2-cm complex mass with fluid collection in the cul-de-sac. Consent was obtained and laparoscopy performed under general anesthesia. Preoperatively, the patient was placed on ampicillin/sulbactam (Unasyn) 1.5 g q 6 h for 3 doses. At the time of surgery, the fallopian tubes were noted to be hyperemic and swollen. Purulent material could be expressed from both fallopian tubes. Approximately 300–400 cc of pus was present in the pelvis. A gram-stained smear of this material revealed the presence of gram-positive, well-encapsulated, lancet-shaped diplococci which subsequently demonstrated a positive Quelling reaction. No organism was isolated; however, the culture techniques were inappropriate for the recovery of S. pneumoniae. The purulent material was drained from the cul-de-sac. An incidental appendectomy was performed. The pathology report revealed a normal appendix. Postoperatively, the patient continued to run a fever until 2 days after the removal of the Jackson-Pratt (JP) drains.

Discussion

Involvement of the female genital tract as a metastatic process secondary to maternal septicemia due to S. pneumoniae is a relatively well-documented phenomenon in the preantibiotic era. The occurrence of cases of chorioamnionitis and/or perinatal septicemia in the absence of pulmonary involvement indicated the possibility of contiguous spread from the vaginal/cervical reservoir and subsequent involvement of the female upper genital tract. The fallopian tubes in the 2 cases of spontaneous peritonitis due to S. pneumoniae in females were described as being swollen and hyperemic with pus emanating from the ends. What was described is not a specific disease entity (spontaneous peritonitis) but more probably a progressive consequence of salpingitis.

Incrimination of S. pneumoniae as an etiological agent in cases of acute salpingitis is probably an underreported phenomenon. While S. pneumoniae will grow on 5–7% sheep blood agar culture when incubated in a CO2 environment, the recovery of alpha hemolytic streptococci is usually not worked up any further and is often reported as “mixed vaginal flora.” Recovery of alpha hemolytic streptococci from patients with acute salpingitis needs to be microbiologically evaluated to exclude the possibility that these isolates are S. pneumoniae.

The case in question coupled with those previously identified as being “spontaneous peritonitis” with isolation of S. pneumoniae from women with tubo-ovarian complexes indicates that S. pneumoniae can function as a primary pathogen for the fallopian tubes and that the entire pathogenic spectrum previously attributable to N. gonorrhoeae can also be mimicked by S. pneumoniae.

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