Periprosthetic joint infection with streptococcus dysgalactiae subspecies equisimilis: Case report

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Human streptococci that belong to *Streptococcus dysgalactiae* (SD) have long been known as non-pathogenic for many years, but are now recognized as an important bacterial pathogen. They are classified under the name of beta-hemolytic groups C and G streptococci.[5]

*Streptococcus dysgalactiae* infections are mostly seen after animal contact or associated with the consumption of unpasteurized dairy food products. There are two subspecies of SD, subspecies dysgalactia (SDSD, an animal pathogen) and subspecies equisimilis (SDSE, a human pathogen). *Streptococcus dysgalactiae subspecies equisimilis* is a group C streptococci and primarily presents as skin and soft-tissue infections, including pyoderma, cellulitis, wound infections, abscesses, erysipelas, and necrotizing fasciitis.[2-5]

According to our knowledge, there is only one case report identifying SDSD in a periprosthetic joint infection (PJI), while there is no PJI case reported that is infected with SDSE in the English literature.[6]

**CASE REPORT**

A 77-year-old retired male patient referred to our clinic with complaint of swelling and redness in his left leg. He had the diagnosis of cellulitis at his leg and used cefazoline for two weeks. The cellulitis was healed clinically but then he was consulted to our clinic with swelling and hyperemia localized at his knee which was performed total knee arthroplasty (TKA) three years ago. Pain, swelling, hyperemia and limited range of motion were observed in the left knee during physical examination. He had a history of consumption of unpasteurized milk. There was no direct contact to pets or any other animals. A written informed consent was obtained from the patient.

Three aspirations were performed. One aspiration was positive. The antibiogram demonstrated antibiotic sensitivity against trimethoprim-sulfamethoxazole, ampicillin, linezolid, clindamycin, erythromycin, cefotaxime, ceftriaxone, vancomycin, levofloxacin and penicillin. Then, the final diagnosis was established based on the adaptation of the Musculoskeletal Infection Society (MSIS) criteria (Table I).[7]

1. One positive culture
2. High C-reactive protein (CRP) (11.21 mg/L)
& erythrocyte sedimentation rate (ESR) (57 mm/h)
3. More than 90% polymorphonuclear neutrophils (PMN%) in synovial fluid
4. Leukocyte esterase strip test showed 2+

The SD was identified in the first sample (Figure 1). The aspiration sample was inoculated to aerobic and anaerobic blood culture tubes (Becton-Dickinson, New Jersey, USA). Also, some of the sample was reserved for gram staining and direct inoculation to blood agar (Salubris Inc. Pharmaceuticals, Istanbul, Turkey), eosin methylene blue agar (Salubris) and chocolate agar (Salubris). The samples were incubated in 37°C for 18-24 hours in carbon dioxide medium. In gram staining, gram-positive cocci were seen. The blood culture tubes demonstrated signals for cultivation by day six. The cultivated bacteria were identified as SD using conventional and automatized systems (Vitek® 2 Compact, BioMérieux, Craponne, France). The antibiogram for the identified species were determined using the Clinical and Laboratory Standards Institute (CLSI) criteria. For further identification and confirmation, Vitek® MS (MALDI-TOF) (BioMérieux) was used and the subspecies were identified as SDSE.

The case was considered as hematogenous spread of infection and as suggested in PJI consensus meeting, thorough debridement of the synovial sheath was performed and polyethylene insert was adapted. Postoperative antibiotic treatment was continued with third-generation cephalosporin for two weeks and then stopped when the infection symptoms were relieved, the CRP and the ESR levels dropped down and presented a plateau (Figure 2). The patient was followed-up for 11 months without recurrent infection.

DISCUSSION

The PJI is the most devastating complication of total joint arthroplasty. The incidence of PJI is around 1% in TKA and the most common agent is Staphylococcus aureus. Streptococci are responsible for 5 to 7% of deep infection cases after total joint arthroplasty. Most of the streptococci are group A, alpha or beta hemolytic type. According to our research, there is only one arthroplasty case infected with SD[6] and this case is the only SDSE PJI reported in the literature.[9]

<table>
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<th>Major criteria</th>
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<td>1. Two positive periprosthetic cultures with phenotypically identical organisms, or</td>
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<td>2. A sinus tract communicating with the joint</td>
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<table>
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<th>Minor criteria</th>
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<tr>
<td>1. Elevated serum erythrocyte sedimentation rate (ESR) and serum C-reactive protein (CRP) concentration,</td>
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<tr>
<td>2. Elevated synovial leukocyte count,</td>
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<td>3. Elevated synovial polymorphonuclear neutrophils (PMN%),</td>
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<td>4. Presence of purulence in the affected joint,</td>
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<td>5. Isolation of a microorganism in one culture of periprosthetic tissue or fluid, or</td>
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<tr>
<td>6. Greater than five neutrophils per high-power field in five high-power fields observed from histologic analysis of periprosthetic tissue at 400 magnification.</td>
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**Diagnosis can be established when one major criterion exits or four of the minor criteria exist.**

**TABLE I**

Musculoskeletal Infection Society criteria for periprosthetic joint infection

**FIGURE 1.** Patient had total knee arthroplasty history at his infected knee. *Streptococcus dysgalactiae* was determined by culture.
Streptococcus dysgalactiae subspecies equisimilis has generally been recovered from the pharynges of carriers and from those with exudative pharyngitis and tonsillitis. There are a limited number of papers reporting SDSE infections in elderly population with an increasing incidence in this age group.\(^{[10]}\)

Group G streptococcal cellulitis, septic arthritis at or near sites of parenteral injection, with bacteremia and subsequent hematogenous complications have been reported in intravenous (IV) drug users. In our case, the PJI was followed by a cellulitis at a near site. However, there was no history of IV drug use.

Streptococcus dysgalactiae subspecies equisimilis isolates remain almost uniformly susceptible to beta-lactam agents. To avoid delayed or poor responses of infections because of failure of penicillin/cephalosporins, the addition of an aminoglycoside should be considered for serious infections.\(^{[11]}\) In patients with penicillin allergy, and in type I allergic patients, vancomycin and clindamycin is a sensible alternative.\(^{[11]}\)

Although the group C streptococcus PJI is extremely rare, the incidence of infection is increasing in elderly population. The group C streptococci should be kept in mind as an infecting agent particularly in elderly patients.

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