Neonatal meningitis due to *Streptococcus dysgalactiae* subspecies *equisimilis*: a case report and literature review

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**Abstract**

*Streptococcus dysgalactiae* subspp. *equisimilis* (SDSE) was firstly reported as a toxon among human streptococcal pathogens in the year 1996. Diseases caused by SDSE may vary from milder skin involvements including wound infection, erysipelas, and cellulitis to life-threatening clinical pictures as streptococcal toxic shock syndrome and necrotizing fasciitis. We describe a case of SDSE sepsis and meningitis in a 2-day-old newborn. He was referred to our intensive care unit (ICU) with fever, respiratory distress, seizures, peripheral cyanosis, and somnolence. SDSE grew out of the CSF and blood cultures obtained on admission to the ICU. SDSE cases reported in the literature are frequently older patients with an underlying disease. Also, SDSE may cause serious neonatal infections.

**Keywords**

*Streptococcus Dysgalactiae Subspecies Equisimilis; Neonate; Meningitis*

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Introduction
Neonatal sepsis is an informed serious disease, and Group B streptococci (GBS) is the most common cause of neonatal sepsis. The use of prophylactic antimicrobials has decreased the incidence of GBS-related disease [1]. SDSE was firstly reported as a human pathogen in 1996 [2]. This microorganism has Lancefield Group C or G antigens and rarely an antigen. This microorganism causes beta hemolysis and exerts streptokinetic activity on human plasminogen and proteolytic activity on human fibrin. SDSE can be found in the oropharynx, gastrointestinal tract, genitourinary tract and as normal skin flora. Diseases caused by SDSE vary from mild skin infections, such as wound infections, erysipelas, and cellulitis to life-threatening infections, such as streptococcal toxic shock syndrome and necrotizing fasciitis [3]. The most common cause of bacteremia is cellulitis [4]. Invasive SDSE infections are becoming increasingly more common worldwide [2]. The invasive form of SDSE infection is more common among older patients especially, in the presence of underlying disease and in cases where skin integrity is disrupted. We report here a case of neonatal sepsis and menigitis caused by SDSE occurring 2 days after birth. We also reviewed the literature on SDSE infections.

Case report
The neonate described here is a male delivered to a gravida 1 mother vaginally at 38-weeks gestation with a birth weight of 3800 gr. The woman had rupture of membranes 8 hours prior to delivery. The rupture of membranes was spontaneous. The Apgar scores at 1 and 5 minutes were 8 and 9, respectively. The neonate began vomiting, developed cyanosis, and somnolence. On admission to our NICU, he was treated with oxygen therapy, intravenous fluid and dobutamine (5 µg/kg/min) for hypotension (arterial blood pressure of 23/14 mm Hg). An arterial blood gas showed metabolic acidosis (pH: 7.28, HCO3: 20 mcg/mL, vancomycin 1mcg/mL). On the third day of therapy, the blood and CSF glucose ratio was 48/89. His mother did not allow us to obtain vaginal or anal swab specimens for antibiotic susceptibility testing. By 3rd weeks of treatment, the blood and CSF results were normal, and the child was clinically improved. The child’s immunity was evaluated and found to be normal. Six months later there were no residual neurological sequelae.

Discussion
SDSE is a microorganism with increasing clinical significance, which causes both invasive and non-invasive infections [5]. Invasive infection with SDSE is seen in older individuals with a suppressed immune system or disrupted skin integrity. Group C β-hemolytic streptococcus have been involved in several localized human infections including pharyngitis, pyodermitis, cellulitis, wound infections, abscesses, erysipelas and necrotizing fasciitis. Severe invasive infections often occur in predisposed hosts: in fact they are common in patients affected by underlying immunodeficiency predisposing diseases or conditions such as age (neonate or elderly), diabetes mellitus, HIV-1 disease, alcoholism and injection drug use, and also in patients with chronic cardiovascular diseases and those undergoing chemotherapy or affected by cancer [6]. We can say that our case is in the immunodeficiency category because it is a newborn. Yamaoka et al. reported a case of streptococcal toxic shock syndrome caused by SDSE in a 12-hour-old newborn without a previous history of premature rupture of membranes or meconium aspiration [2]. In the literature, two cases of early onset neonatal sepsis caused by group G streptococci were reported before identification of this microorganism. One of these cases had a history of meconium aspiration, and the other had prolonged rupture of membranes [7]. The demographic and clinical characteristics of a total of 3 cases cited in the literature are shown in Table 1. The reported neonates -including ours- were delivered vaginally and had no identifiable risk factors.

In the literature, screening of pregnant women was recommended as for the presence of GBS to prevent the development of early-onset neonatal sepsis [8]. In our case, the mother did not undergo screening during pregnancy. Analysis of the emm gene for the amino acid sequence situated and the N-terminal end of the M-protein in patients with SDSE has been performed in epidemiological studies related to epidemics caused by invasive and non-invasive microorganisms [2]. The presence of the emm I gene in children is associated with invasive infection. SDSE cases reported in the literature are frequently older patients with an underlying disease. Mortality rates have been high among patients with SDSE. In patients with bacteremia due to SDSE, mortality rates are reported to be 15-18% [4]. In addition to this reported case of SDSE in a neonate at our hospital, only 2 other cases among neonates have been reported to survive. The prognosis of neonates with SDSE infection is better than adults. This may be because neonates usually do not have any underlying disease but adults with SDSE often do, and neonates may be diagnosed earlier while they are still in the hospital at birth. In conclusion, SDSE may cause serious neonatal infections.
Table 1. Demographic and clinical characteristics of Streptococcus dysgalactiae subspecies equisimilis infection in neonatal cases.

<table>
<thead>
<tr>
<th>References</th>
<th>Age</th>
<th>Gender</th>
<th>Delivery method</th>
<th>Birth weight (gr)</th>
<th>Clinical syndrome</th>
<th>Maternal screening</th>
<th>Underlying condition(s)</th>
<th>Therapy</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yamaoka et al. 2010</td>
<td>&lt;5</td>
<td>Male</td>
<td>Vaginal delivery</td>
<td>2894</td>
<td>Streptococcal toxic shock syndrome, septicemia, meningitis</td>
<td>Positive</td>
<td>No</td>
<td>Ampicillin</td>
<td>Complete resolution</td>
</tr>
<tr>
<td>Carstensen et al. 1988</td>
<td>&lt;5</td>
<td>Male</td>
<td>Vaginal delivery</td>
<td>4000</td>
<td>Septicemia</td>
<td>Positive</td>
<td>Meconium aspiration</td>
<td>Penicillin, Gentamicin</td>
<td>Complete resolution</td>
</tr>
<tr>
<td>Carstensen et al. 1988</td>
<td>&lt;5</td>
<td>Female</td>
<td>Vaginal delivery</td>
<td>3570</td>
<td>Septicemia</td>
<td>Result unshared</td>
<td>Rupture of membranes</td>
<td>Penicillin, Gentamicin</td>
<td>Complete resolution</td>
</tr>
<tr>
<td>Present case</td>
<td>&lt;5</td>
<td>Male</td>
<td>Vaginal delivery</td>
<td>3800</td>
<td>Septicemia, meningitis</td>
<td>Could not be performed</td>
<td>None</td>
<td>Ampicillin, Vancomycin</td>
<td>Complete resolution</td>
</tr>
</tbody>
</table>

Scientific Responsibility Statement
The authors declare that they are responsible for the article’s scientific content including study design, data collection, analysis and interpretation, writing, some of the main line, or all of the preparation and scientific review of the contents and approval of the final version of the article.

Animal and human rights statement
All procedures performed in this study were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. No animal or human studies were carried out by the authors for this article.

Conflict of interest
None of the authors received any type of financial support that could be considered potential conflict of interest regarding the manuscript or its submission.

References

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