Case Report

*Streptococcus agalactiae* vaginitis: nonhemolytic variant on the Liofilchem® Chromatic StreptoB

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**Abstract:** *Streptococcus agalactiae* (group B Streptococcus, GBS) vaginal pathogenicity is not uniformly acknowledged throughout the literature; accordingly, in women, genital itching and burning, along with leukorrhea are commonly and almost exclusively referred to bacterial vaginosis, candidiasis and trichomoniasis. Conversely, GBS virulence for vagina was recognized in the past, as the organism has been observed to potentially cause local inflammation and discharge, as well as lactobacilli rarefaction. We depict here a case where a nonhemolytic (γ-hemolytic) GBS strain was found to be the etiologic agent of vaginal infection. Such uncommon *S. agalactiae* phenotypes are hard to be recognized and may be therefore responsible for misdiagnosing and underestimation of GBS vaginitis prevalence; here, we had the support of the Liofilchem® Chromatic StreptoB medium, that successfully detected such an atypical variant.

**Keywords:** *Streptococcus agalactiae*, group B antigen, GBS, vaginitis, leukorrhea

**Case report**

*Streptococcus agalactiae* (group B Streptococcus, GBS) is responsible for serious infections in newborns, such as sepsicemia, meningitis, and pneumonia. Also, it may cause diverse diseases in the adult population, including skin, soft tissue, urinary tract and osteoarticular pathologies, pneumonia, peritonitis, meningitis, endocarditis, pyometra and bacteremia (without a focus). Nevertheless, GBS pathogenicity for vagina is not uniformly acknowledged; hence, the organism is not usually included, unfortunately, in the etiologic diagnosis of vaginitis. Conversely, it is mostly dismissed as a saprophyte or a transient colonizer despite the isolation as a single organism from vaginal exudates of symptomatic women [3, 5].

A 23-year old patient came to our attention due to genital pruritus and burning, white discharge and mucosal erythema (the latter was documented at speculum examination). Exudate observation under a microscope showed numerous polymorphonuclear cells and Gram positive cocci (bacteria were even inside leukocytes), along with lactobacilli rarefaction, and absence of ‘clue cells’, *Trichomonas vaginalis* trophozoites and fungal images (blastospires, hyphae, pseudohyphae). pH value was >4.5 (that is alkaline), with a negative amine test. Both the clinical and the microbiological pictures indicated therefore an ongoing vaginal infection due to coccoid organisms, and with underlying rarefaction of the saprophyte microflora.

When cultivating the woman’s exudate, a massive growth of nonhemolytic (γ-hemolytic), Gram positive, catalase- and coagulase-negative cocci was obtained on the bioMérieux Columbia agar (plus 5% sheep blood) (Figure 1); culture was pure, as no other bacterial phenotypes could be observed on none of blood medium, mannitol salt and MacConKey agar; also, Sabouraud plates did not show any fungal growth (after concomitant incubation both at 36±1°C and 25±1°C, for 4 days). Such an isolate was unexpected, and initially labeled as belonging to the normal microflora.
Nevertheless, given it was a pure culture, we carefully studied it, and surprisingly found it possessed the Lancefield group B antigen (latex test by Liofilchem®, Roseto degli Abruzzi, Italy). Also, after 24 h incubation, the Chromatic Strepto B medium (Liofilchem®) showed green-blue colonies (Figure 2) that are described (in the product package insert) to be formed by GBS (enterococci produce instead purple colonies). In agreement, the Vitek2 system (bio-Mérieux, Marcy l’Etoile, France) provided identification as S. agalactiae (99% certainty). The isolate was finally deposited into our laboratory collection as S. agalactiae strain Sa29.

The patient made a full recovery after 10-day topical lactobacilli (1 capsule/die) plus 6-day oral amoxicillin (1 gr, twice a day). Post-therapy cultures performed after two weeks from stopping treatment showed restoration of a normal microflora as well as GBS eradication.

Particularly, it was also suggested that the β-lactam be assumed by her partner, too.

We believe readers may capture a few interesting messages from this brief communication. First, S. agalactiae pathogenicity for vagina is poorly acknowledged, then the organism is mostly labeled as an innocent by-stander despite the isolation as a pure culture from leukorrhea patients [1, 3, 5, 6]. Nevertheless, GBS has been observed to provoke cytokine release in the vaginal tissue, genital inflammation as well as decrease of the lactobacillary microflora; it is therefore a potential cause of vulvovaginal burning, discharge (usually clear/white-coloured), and fiery red labia, even with fissures [1].

Secondly, nonhemolytic GBS isolates are presumed to be uncommon (5-8% of all isolates) [6]; indeed, we believe their prevalence might be underestimated due to misdiagnosing. The case we presented, moreover, further highlights that such phenotypical variants may show the same pathogenicity as the more commonly encountered β-hemolytic S. agalactiae strains. Particularly, then, when γ-hemolytic streptococci are found to massively grow from symptomatic women, and in the absence of any other detectable vaginal pathogen, they should always be screened for the group B antigen. Otherwise, their recognition on a visual basis is exceedingly hard and, unfortunately, chromogenic media may fail in identifying such phenotypes [6]. We achieved a good result with the Liofilchem® Chromatic Strepto B. However, no further observation has been published to our knowledge about this medium performance, so a wider number of β- and γ-hemolytic GBS strains must be tested in order to define this product’s properties and limits.

Nowadays, vaginal inflammation is mostly considered to be related to Candida and T. vaginatis [2, 4]; nonetheless, we would like to finally emphasize that GBS may represent an underestimated agent of vaginitis. Recognition of γ-hemolytic variants may shed more light on S. agalactiae genital infection prevalence; in this context, a reliable etiologic diagnosis, achieved through both standard and chromogenic media (provided that these may detect nonhemolytic phenotypes), is indispensable to avoid improper treatments and chronification of female discomfort.

Disclosure of conflict of interest

None to declare.

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References


