Microbiological Profile of Vaginosis among Women of Reproductive Age Group Attending a Tertiary Care Hospital

RENU MATHEW, SUDHAKSHINA.R, M.KALYANI, S.JAYAKUMAR, BINESH LAL, SHAMEEM BANU

ABSTRACT
Background and Objective: Although Nugent’s criterion is considered as the gold standard, routinely a combination of various methods is used for the diagnosis of bacterial vaginosis. In the present study, we compared the culture, Spiegel’s criteria and Amsel’s criteria with the Nugent’s method for the diagnosis of bacterial vaginosis.

Materials and Methods: Five hundred and twenty seven women who attended the Government Maternity Hospital and a tertiary care centre in south India for antenatal care or for any other complaint formed the study population. The diagnosis of bacterial vaginosis was done by culture and Amsel’s, Nugent’s and Spiegel’s criteria. The positive predictive value, the negative predictive value and the sensitivity and specificity of these methods in comparison with Nugent’s criteria, considering it as the gold standard, were calculated. The statistical analysis was done by using the Chi Square test or the Fisher’s exact test as was appropriate.

Results: In comparison with Nugent’s criteria, the positive predictive value, the negative predictive value and the sensitivity and specificity of Amsel’s criteria were found to be 80.4%, 94.8%, 78% and 95.6% and those of Spiegel’s criteria were found to be 77.5%, 100%, 100% and 93.2%. The culture was 51% sensitive and 88.7% specific, the positive predictive value was 85.5% and the negative predictive value was 58%. We diagnosed 100 (19%) cases of bacterial vaginosis by Nugent’s method, 129 (24%) cases by Spiegel’s method, 97 (18%) cases by Amsel’s criteria and 88 (16.7%) cases by culture.

Conclusion: Amsel’s and Spiegel’s criteria were comparable with Nugent’s criteria for the diagnosis of bacterial vaginosis. Culture was the least sensitive method for the diagnosis of bacterial vaginosis.

KEY MESSAGE
- Amsel’s and Spiegel’s criteria were comparable with Nugent’s criteria for the diagnosis of bacterial vaginosis.
- Culture was the least sensitive method for the diagnosis of bacterial vaginosis.

INTRODUCTION
Bacterial vaginosis is a common clinical condition in women of the reproductive age group [1]. It represents a unique and complex change in the flora of the vagina, which is characterized by a reduction in the prevalence and the numbers of lactobacilli and an increase in the concentration of Gardnerella vaginalis and the resident anaerobic bacteria [1]. Most of the women are asymptomatic, but some women with bacterial vaginosis have a foul smelling, thin, homogeneous, frothy, vaginal discharge [1],[2]. In addition to being a nuisance infection, bacterial vaginosis can lead to a variety of obstetric and gynaecological complications such as preterm birth and pelvic inflammatory disease (PID) [1],[2]. As it is just an overgrowth of the normal flora of the vagina without inflammation, there is no single best method for the diagnosis of bacterial vaginosis [1],[2]. Most often, multiple criteria are used for the diagnosis of bacterial vaginosis. One of the methods of diagnosis is the Amsel’s composite criteria which includes clinical diagnosis and a few simple laboratory tests [3].

Bacterial vaginosis can also be diagnosed by Spiegel’s and Nugent’s criteria [4]. Both the criteria are based on the evaluation of the normal flora in gram stained smears of vaginal discharge. In the present study, we compared the culture, Amsel’s criteria and Spiegel’s criteria with Nugent’s criteria, while considering it as gold standard for diagnosis of bacterial vaginosis [5],[6].

MATERIALS AND METHODS
Five hundred and twenty seven women who attended two hospitals in south India for antenatal care and the insertion or the removal of an intrauterine contraceptive device, with complaints of discharge, abdominal pain or any other complaint formed the study population. The study had the approval of the Institutional Ethics Committee.

Married women in the age group of 21–35 years and women with or without vaginal discharge complaints were included. Women who were menstruating at the time of the specimen collection and women who were on medication for any bacterial, fungal, parasitic or viral infections for up to one month prior to the specimen collection were excluded. A detailed clinical history of each woman was taken and their vaginal swabs were collected. The vaginal swabs were used for gram staining, for the determination of the pH of the vagina,
Whiff's test and culture. The diagnosis of bacterial vaginosis was done by Nugent's criteria, Amsel's criteria, Spiegel's criteria and by culture. The parameters that were necessary to decide the efficacy of the diagnostic tests, namely the positive predictive value, the negative predictive value and the sensitivity and specificity were calculated in comparison with Nugent's criteria, considering it as gold standard. The statistical analysis was done by using the Chi Square test or the Fisher's exact test as was appropriate.

**Diagnosis by Amsel's Criteria**
Amsel's composite criteria includes a homogeneous vaginal discharge, pH of the vagina > 4.5, the presence of clue cells in the gram stained vaginal discharge smears and a positive Whiff's test. According to Amsel, if 3 of the 4 criteria were positive, the patient was considered to have bacterial vaginosis [3].

Vaginal pH determination: Vaginal secretions or discharges were collected from the lateral vaginal walls with a cotton swab and they were then transferred onto strips of pH paper (Qualigens Fine Chemicals, India) and were compared with a standardized colourimetric reference chart to estimate the actual pH [4,7].

**Whiff's test**: A drop of vaginal discharge was mixed with a drop of 10% potassium hydroxide which was taken on a slide. A fishy smell indicated a positive test [8].

**Clue cells**: The vaginal discharge was smeared on clean glass slides, air dried, heat fixed and stained by Gram's Method by using an acetone alcohol (1:1) mixture as a decolouriser and dilute Carbol Fuchsin as the counter stain. The vaginal epithelial cells were completely covered by gram variable coccobacilli, so that their edges which normally had a sharply defined cell border, became indistinct or stippled and were considered as clue cells [9] [Table/Fig-1].

**Diagnosis by Culture**
The vaginal swabs were inoculated on appropriate culture media and they were incubated at 37ºC for 24 to 48 hours. For the diagnosis of bacterial vaginosis by culture [2].

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**Diagnosis by Culture**
The vaginal swabs were inoculated on appropriate culture media  and they were incubated at 37ºC for 24 to 48 hours. For the isolation of aerobes and facultative anaerobes, Columbia blood agar and Mac Conkey's agar were used [10]. For the isolation of *G. vaginalis*, Columbia human blood bilayer agar with Tween 80 and a *G. vaginalis* selective supplement was used [11]. These plates were incubated in a candle jar with a piece of wet, sterile cotton placed in it to provide a humid environment. For anaerobes, Columbia laked human blood agar with a neomycin supplement was used [12]. These plates were incubated in an Anaero Hi Gas Pack™ anaerobic jar (Hi Media Laboratories, Pvt. Ltd., Mumbai, India). Aerobes, facultative anaerobes and obligate anaerobes were identified by their colony morphologies, gram staining and standard biochemical reactions [10, 12]. All the media, reagents and discs were obtained from Hi Media Laboratories, Pvt. Ltd., Mumbai. Those women in whom the culture showed a predominant growth of *G. vaginalis* or an anaerobe or both were considered as positive for bacterial vaginosis by culture [2].

**Diagnosis by Nugent's Criteria**
Each bacterial morphotype was quantified under an oil immersion objective (1000 x) by using the following scheme: 1+, <1 per field; 2+, 1 to 5 per field; 3+, 6 to 30 per field and 4+, >30 per field. The large, gram-positive rods were considered as lactobacillus morphotypes; the small, gram-negative to gram-variable rods were considered as *G. vaginalis* and the *Bacteroides* spp. morphotypes and the curved gram-variable rods were considered as the Mobiluncus spp. morphotypes. The scoring was done as shown in [Table/Fig-2]. These scores were added up to yield a final score of 0 to 7 or more. The criterion for bacterial vaginosis was a score of 7 or higher; a score of 4 to 6 was considered as intermediate, and a score of 0 to 3 was considered as normal [3],[4],[5].

**Diagnosis by Spiegel's Criteria**
When the gram staining showed predominance (3 to 4+) of the lactobacillus morphotype with or without the Gardnerella morphotype, it was interpreted as normal. When the gram staining showed a mixed flora which consisted of gram-positive, gram negative, or gram-variable bacteria and when the lactobacillus morphotype was decreased or absent (0 to 2+), the gram stain was interpreted as consistent with bacterial vaginosis [4].

<table>
<thead>
<tr>
<th>Methods of diagnosis</th>
<th>Results</th>
<th>Nugent's score &gt; 7</th>
<th>Nugent's score (0-6)</th>
<th>Total</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amsel's criteria</td>
<td>Bacterial vaginosis</td>
<td>78</td>
<td>19</td>
<td>97</td>
<td>&lt; 0.01</td>
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<tr>
<td></td>
<td>Normal</td>
<td>22</td>
<td>408</td>
<td>430</td>
<td></td>
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<tr>
<td>Spiegel's criteria</td>
<td>Bacterial vaginosis</td>
<td>100</td>
<td>29</td>
<td>129</td>
<td>&lt; 0.01</td>
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<tr>
<td></td>
<td>Normal</td>
<td>0</td>
<td>398</td>
<td>398</td>
<td></td>
</tr>
<tr>
<td>Culture</td>
<td>Bacterial Vaginosis</td>
<td>51</td>
<td>37</td>
<td>88</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td></td>
<td>Normal</td>
<td>49</td>
<td>290</td>
<td>339</td>
<td></td>
</tr>
</tbody>
</table>

[Table/Fig-2]: Comparison of diagnosis of bacterial vaginosis by culture, Amsel's and Spiegel's criteria with the gold standard Nugent's criteria

**Results**
The results of the diagnosis of bacterial vaginosis which was done by Amsel's criteria, culture, Nugent's criteria and Spiegel's criteria are shown in [Table/Fig-3]. We diagnosed 100 (19%) cases of bacterial vaginosis by Nugent's method, 129 (24%) cases by Spiegel's method, 97 (18%) cases by Amsel's criteria and 88 (16.7%) cases by culture. In comparison with Nugent's criteria, the positive predictive value, the negative predictive value, and the sensitivity and specificity of Amsel's criteria were found to be 80.4%, 94.8%, 78% and 95.6% and those of Spiegel's criteria were found to be 77.5%, 100%, 100% and 93.2%. The culture was 51% sensitive and 88.7% specific, the positive predictive value was 85.5% and the negative predictive value was 58%. The statistical analysis showed that all the 4 methods could be used as a means of diagnosis of bacterial vaginosis (p < 0.01)
DISCUSSION

Here, we conducted a study on 100 cases of bacterial vaginosis which were diagnosed by the gold standard method, Nugent’s criteria [5,6]. It classifies gram stained vaginal smears into normal, intermediate and bacterial vaginosis, based on the gram stain scoring system. The standardized score had an improved inter center reliability as compared to the Spiegel’s criteria which divided the gram stained vaginal smears into only 2 categories, normal or bacterial vaginosis [4,5,6]. In a previous study where women with intermediate flora were followed up to 3 months, some of them developed bacterial vaginosis, some continued to have intermediate vaginal flora and some reverted to the normal flora patterns [13]. So, it is evident that women with intermediate flora must be considered separately. Hence, the Spiegel’s criteria which divides women into only 2 categories, bacterial vaginosis and normal, is not as popular as Nugent’s method. There are many studies which have tried to formulate better the gram stain scoring systems, but they are not as popular as the Nugent’s method of diagnosis of bacterial vaginosis [14,15].

Previous studies have shown that the diagnosis of bacterial vaginosis by Amsel’s criteria was less sensitive than the gram stain interpretation [3,16]. This low sensitivity may be because many cases of bacterial vaginosis are asymptomatic. In the present study, Amsel’s method was found to be 78% sensitive and 95.6% specific as compared to Nugent’s method. The diagnosis by Amsel’s criteria requires a minimum of 3 to 5 vaginal swabs from each patient [3,16]. It was observed that routinely, only a single swab was sent to the laboratory to rule out bacterial vaginosis in the hospitals where the study was carried out. This might be the reason why the diagnosis of bacterial vaginosis by Amsel’s criteria was unpopular in these places. But Amsel’s method is very popular as a means of diagnosis of bacterial vaginosis as has been reported in every research paper on bacterial vaginosis [3,4,6].

Culture is the gold standard method for the diagnosis of most of the bacterial diseases; however, culture cannot become the gold standard for the diagnosis of bacterial vaginosis, as the organisms which are involved in bacterial vaginosis cannot be isolated in the laboratory easily and as normal women also have this flora in their vagina in small numbers.

The rate of bacterial vaginosis, when it was diagnosed by Nugent's scoring system, was 19%. Indian studies which were conducted on the general population have shown a similar prevalence [2,17].

CONCLUSION

Amsel’s and Spiegel’s criteria were comparable with Nugent’s criteria for the diagnosis of bacterial vaginosis. Culture was the least sensitive method for the diagnosis of bacterial vaginosis.

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