Growth of *Candida* species in liquid culture medium for *Trichomonas vaginalis*

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**SUMMARY** The growth of *Candida* spp from vaginal specimens in Bushby's liquid medium for *T vaginalis* was compared with that on Sabouraud's agar medium, and isolation was significantly greater in Bushby's medium (*p*<0.001). Isolations missed (4.43%) in Bushby's medium probably represented vaginal carriage of small numbers of *Candida* spp.

**Introduction**

Feinberg and Whittington observed that *Candida* spp grew well in culture medium for *Trichomonas vaginalis*. Both these organisms cause vaginal disease, and it would be cost effective if they could be isolated from a single liquid medium. To compare the isolation rate of *Candida* spp on conventional Sabouraud's agar with that obtained in modified Feinberg-Whittington medium a prospective study was carried out in the department of genitorinary medicine at this hospital.

**Patients and methods**

**DESIGN STUDY**

All women patients (1029) who attended during a six week period in March and April 1982 were included. A loopful of material was taken from the lateral vaginal wall, Gram stained, and examined for *Candida* spp. Three specimens were taken for culture from each patient using a nichrome wire loop. A sample from the cervical os was inoculated into Bushby's modified Feinberg-Whittington medium. Two samples were taken from the lateral vaginal wall and inoculated either into Bushby's medium or onto Sabouraud's agar (Oxoid) and incubated at 37°C for two days. In the first three week period Bushby's medium was inoculated first, but the order of inoculation was reversed in the second period.

**IDENTIFICATION OF CANDIDA SPP**

Identification of *Candida* spp by their typical colonial appearance on Sabouraud's agar was confirmed by microscopical examination for budding yeast cells. In Bushby's medium microscopical examination revealed budding yeast cells and pseudomycelium. Selected isolates of *Candida* spp were tested by Analytical Profile Index (API) to identify the species. These isolates were also tested for germ tube production. A random selection of 207 isolates of *Candida* spp was surveyed for germ tube production to determine what proportion were *Candida albicans*.

**GROWTH KINETICS OF C ALBICANS IN BUSBY'S MEDIUM**

An isolate of *C albicans* was emulsified in 0.85% saline (about 3·0 × 10⁶ colony forming units (cfu)/ml) and 1 ml of a range of dilutions was inoculated into 9 ml aliquots of Bushby's medium. The growth curves were plotted by means of a nephelometer to determine the rate of increase in optical density. Viable counts were performed on the dilutions by standard microbiological techniques.

The growth kinetics in Bushby's medium of seven strains that had grown only on Sabouraud's agar on primary isolation was compared with that of three others that grew in both media. About 5 × 10⁵ cfu were inoculated into Bushby's medium and the optical density measured by nephelometry.

**Results**

**ISOLATION OF CANDIDA SPECIES**

The table shows that *Candida* spp were isolated from 226 (22.0%) of the 1029 patients in either or both media. In the first three week period isolation of *Candida* spp in Bushby's medium was significantly (*p*<0.001) greater than on Sabouraud's agar, whereas there was no significant difference in yields from the two media in the second period when Sabouraud's agar was inoculated first. Over the...
TABLE  Growth of Candida spp from 1029 women

<table>
<thead>
<tr>
<th>Medium</th>
<th>No of specimens yielding candidal growth</th>
<th>In weeks 1-3*</th>
<th>In weeks 4-6†</th>
<th>In total period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sabouraud’s only</td>
<td>4</td>
<td>6</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Bushby’s only</td>
<td>23</td>
<td>13</td>
<td>36</td>
<td></td>
</tr>
<tr>
<td>Both</td>
<td>101</td>
<td>79</td>
<td>180</td>
<td></td>
</tr>
<tr>
<td>Neither</td>
<td>429</td>
<td>374</td>
<td>803</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>557</td>
<td>472</td>
<td>1029</td>
<td></td>
</tr>
</tbody>
</table>

*Bushby’s medium inoculated before Sabouraud’s agar;
† Sabouraud’s agar inoculated before Bushby’s medium.

The isolation was significantly (p<0.001) greater in Bushby’s medium. Samples from 10 women, however, failed to grow Candida spp in Bushby’s medium although growing on Sabouraud’s agar, which is equivalent to 4-4% false negative results (95% confidence limits of p are 1-7% and 7-1%).

Of the 207 isolates of Candida spp taken at random, 195 (94.2%) were identified as Candida albicans by a positive germ tube test.

INOCULUM SIZES

Fig 1 shows the growth curves obtained by inoculation of varying numbers of C albicans into Bushby’s medium. As few as 300 cfu/ml gave rise to appreciable growth within 24 hours, and even an inoculum as small as 1-5 cfu/ml would establish growth by the fifth day.

GROWTH KINETICS OF VARIOUS ISOLATES OF C ALBICANS

Isolates that grew only on Sabouraud’s agar on primary isolation were all identified as C albicans. Fig 2 shows the growth kinetics of seven such strains compared with that of three others that grew on both media. The latter showed a faster growth rate in the first 18 hours and a slightly greater growth at 28 hours.

Discussion

The significantly greater isolation of Candida spp from Bushby’s medium than from Sabouraud’s agar would imply that a liquid medium like the former would suffice for both Candida spp and T vaginalis. In a small percentage of patients, however, primary isolation was only from Sabouraud’s agar. The reasons for this are not entirely clear, but these patients were probably harbouring very small numbers of Candida spp, which represented vaginal carriage as opposed to infection. Failure to grow in Bushby’s medium may therefore be attributed to chance, particularly as examination of the case notes of the 10 women concerned showed that they had no clear cut signs or symptoms of candidosis. Moreover, a Gram stained smear of material (taken from the lateral vaginal wall before the specimens for culture) gave negative results for Candida spp in these cases. The order in which the media were inoculated played a major role in their respective isolation rates, as can be judged by comparison of the data from the first and second time periods (see table). Not surprisingly, this indicated that the major part of the material went to the first medium inoculated.

The percentage of the 207 isolates of Candida spp identified as C albicans (94.2%) was similar to that reported by Hurley et al. and Oriel et al.
Growth curve experiments showed that an inoculum of less than 100 cfu/ml of *C albicans* would give rise to appreciable growth within 48 hours. All 10 isolates that failed to grow in Bushby's medium were identified as *C albicans*, but they may have been a biotype that did not grow well in this medium. A comparison of the growth characteristics of seven of these strains with three that had been isolated from both media (see Fig 2) showed that the growth yield after 28 hours was not noticeably reduced. Of note, however, was that the seven strains which grew only on Sabouraud's agar had a longer lag phase, but that once they entered the log phase the mean generation time was reduced.

No attempt was made in this study to determine the minimum inoculum of *Candida* spp required to establish growth on Sabouraud's agar, so that a direct comparison in this respect with Bushby's medium is not possible.

When both *Candida* spp and *T vaginalis* are present in the same tube of Bushby's medium, *T vaginalis* may be overgrown with the *Candida* spp by the second day of incubation (Y J Erdman, unpublished observation). Extra care should therefore be taken in microscopic examination of tubes that contain *Candida* spp.

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References