136 Correspondence

Table Isolation of yeasts in three types of the urethra and seminal fluid, and have been specimen from 20 men

Specimen	No	No positive on:		
		Micro- scopy	Culture	- Total
Urethral secretion	20	1 3	1	1
Early morning urine Seminal fluid	20	10	14	14

many of these women have no recognised predisposing factors.

I therefore carried out a study to evaluate the possible role of sexual partners in the maintenance of such a high incidence of vaginal candidosis in our women. I examined 20 husbands or sexual partners of women with diagnosed candidal vaginitis, and collected three different specimens (urethral secretion, early morning urine, and seminal fluid) from each of the men. Moist sterile cotton tipped swabs were used to obtain urethral secretion, which was immediately squeezed out into a sterile phosphate buffer solution using a vortex mixer, and then centrifuged. A drop of the centrifuged deposit was examined by both wet film and Gram staining techniques for evidence of yeast cells. The remaining deposit was cultured on Sabouraud's medium at 37°C, and any growth after 24-48 hours was Gram stained and examined microscopically as before. Both the early morning urine (collected in sterile urine bottle) and seminal fluid (collected in sterile universal container) were also centrifuged and examined as for the urethral secretion. Diagnosis of infection by Candida spp was based on the presence of budding yeast cells or pseudohyphae in wet film and Gram stained smears of the centrifuged deposits of various specimens. This was confirmed by positive cultures of Candida spp on Sabouraud's agar plates.

The table shows the isolation of yeast in relation to the type of specimen. All the men were circumcised and none had any sign of balanitis. Although about half of the men admitted to vague symptoms of urethritis, only three showed any sign at all, and all three had yeasts in their urine and seminal fluid specimens. Where yeasts were isolated from the urethral swab and urine, they were also found in the seminal fluids. Many seminal fluids, however, were positive in the absence of yeast at the other sites.

The role of sexual transmission as a means of vaginal innoculation and colonisation has undergone considerable discussion within the past few years. Most studies, however, have centred on external penile colonisation, usually in the coronal sulcus,23 rather than in

mostly in uncircumcised men. The result obtained in this study shows that examination of seminal fluid for Candida spp is a better means of establishing its prevalence in the male urogenital tract. Yeasts that find their way into the urogenital tract are not always flushed out during micturition. Candidal invasion of the posterior urethra is recognised, and so also is prostatic invasion.4 Sexual transmission may yet be a more important factor in vulvovaginal candidosis. especially the recurrent forms, than has been previously recognised; seminal fluid acting as the source of innoculum during the powerful muscular contraction attending orgasm. I therefore recommend that, in evaluating the role of sexual transission in the epidemiology of candida vaginitis, investigation of the seminal fluid should take precedence over the traditional urethral and urine examinations.

Yours faithfully. B O Ogunbanjo

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TO THE EDITOR, Genitourinary Medicine

## Use of slide latex agglutination test for rapid diagnosis of vaginal candidosis

The recent letter from Sulaiman et al describing their results with the latex agglutination test is, in certain key respects, at variance with our own wide experience of this product. The candida slide test is intended as a rapid aid to the "patient-side" diagnosis of vulvovaginal candidosis. In this sense the only alternative is direct microscopy, as culture isolation requries at least 48 hours. The latex slide test is typically appreciably more sensitive than direct microscopy, as evidenced by Dr Sulaiman's study. In comparison with culture isolation, the latex test largely ignores symtomless (commensal) carriers and so, though sensitivity is slightly lower, the diagnostic positive predictive value is superior. In Dr Sulaiman's study 21% of culture positives came from symptomless women. Our own studies have suggested that the incidence of candida culture positive symptomless women can be considerably higher than this. though the only respect in which Dr Sulaiman's data significantly conflicts with our own experience is in the number of "false" latex positives (nine out of 23).

During the course of clinical trials of over 1000 unselected women attending two leading departments of genitourinary medicine in the United Kingdom, the lowest recorded specificity was 95.4% and the typical specificity value was 97-98%.<sup>2-4</sup> It is not readily apparent, from Dr Sulaiman's report, whether any of the nine specimens "falsely" positive by latex agglutination were culture positive.

The latex agglutination test should be evaluated against a carefully considered definition of vulvovaginal candidosis, and our own trials were based upon the definition described by Dr Sulaiman. In view of the relatively low prevalance in unselected women, the performances of the various diagnostic criteria are best assessed in terms of sensitivity and predictive values. In such circumstances the latex agglutination test invariably exhibits a higher diagnostic efficiency than any other single criterion.

Yours faithfully, D H Lewis

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TO THE EDITOR, Genitourinary Medicine

Sir,

I should be much obliged if you would grant me the courtesy of your columns to thank the