tics without improvement. We treated her successfully with local applications of piperazine solution for only eight days, and more than 18 months elapsed since without any recurrence. In chronic cases of sterile pyuria, therefore, urine should be examined for parasites. If the ova or larvae of E vermicularis are found treatment should be by local irrigation of the urethra and bladder and vaginal application of piperazine salts.

The most probable source of repeated urinary tract infections in our patient was the vagina and not the intestinal tract. This explanation is based on three observations: (1) stools and parianal swabs from her and her husband were repeatedly negative for E vermicularis, (2) her condition responded to topical vaginal treatment, and (3) no antimicrobial or other drug was required to treat this condition.

Yours faithfully, Sarman Singh J C Samantaray

Section of Parasitology, Department of Medical Microbiology, All India Institute of Medical Sciences, New Delhi 110 029, India

References

- Belding DL, ed. Textbook of clinical parasitology. 2nd ed. New York: Appleton-Century-Crofts, 1952:422-34.
- 2 Sjovall A, Akerman M. Peritoneal granuloma in women due to the presence of oxyuris vermicularis. Acta Obstet Gynecol Scand 1968:47:361-72.
- 3 Adungo NI, Ondijo SO, Pamba HO. Observation of Enterobius vermicularis ova in urine: three case reports. East Afr Med J 1986; 63:676-8.

TO THE EDITOR, Genitourinary Medicine

Importance of Gardnerella vaginalis as an aetiological agent in bacterial vaginosis

Sir.

Ching et al have reported that using the PEM-GVA (plastic envelope method) Gardnerella vaginalis was isolated from 47 of 49 (96%) of women with clinical bacterial vaginosis (BV). The PEM broth medium showed an adherence of G vaginalis bacteria in 75% of these patients.

Using the PEM-GVA test, we undertook a study specifically directed at further investigating the in vitro adherence of *G vaginalis* in patients with symptomatic BV. We studied 103 consecutive women attending a local health clinic. We compared the results of pelvic examinations with results of the PEM-GVA and conventional techniques. 1-3

Table 1 shows that G vaginalis was isolated

Table 1 Isolation of Gardnerella vaginalis from 19 women with and 84 without bacterial vaginalis (BV) (figures are numbers (percentages) of women yielding G vaginalis)

Category	No	PEM-GVA	Conventional culture
BV	19	19 (100)	16 (84-2)
Non-BV	84	19 (100) 18 (21·4)	16 (84·2) 18 (21·4)
Totals	103	37	34

from 19 (100%) women with BV and 18 (21.4%) of the remaining 84 patients, who did not have BV. Appreciable adherence, as shown in table 2, occurred in 18 (95%) of the 19 women with BV and five (6%) of the 84 other patients (p < 0.0001). Table 3 shows that appreciable numbers of clue cells were found in 16 (84%) of the 19 women with BV, and four (5%) of the 84 other patients (p < 0.0001). When appreciable bacterial adherence and clue cells were absent, results correlated best with a BV negative predictive value of 98%.

A previously unreported observation was used in this study as a possible indicator of BV, namely the presence of gas bubbles in a patient's discharge. It was present in all the positive symptomatic confirmed cases.

In vivo adhesion of *G vaginalis* to epithelial cells may be important in the pathogenesis of BV. Whether any relation exists between the in vivo and in vitro adherence described previously is speculative. The results of this study, however, indicate that the PEM-GVA provides a rapid, sensitive, and specific method of growing *G vaginalis*.

Table 2 Adherence density of Gardnerella vaginalis in 103 women

Adherence density	BV	Non-BV
4+	14	1
4+ 3+	4	4
2+	1	5
1+	Ō	3
None	9	71
Total	19	84

Appreciable densities were 3+ and 4+.

Table 3 Clue cell density in 103 women

Clue cell density	BV	Non-BV
4+	10	0
4+ 3+ 2+	6	. 4
2+	2	4
1+	1	7
None	0	69
Total	19	84

Appreciable densities were 3+ and 4+.

This system also presents the microscopic visualisation of bacterial adherence and clue cells that correlate significantly with the clinical diagnosis of BV.

Yours faithfully, K A Borchardt* B S Adly* R F Smith† J Eapen‡ C B Bealt

*Center for Advanced Medical Technology, San Francisco State University, San Francisco.

†Contra Costa County Health Department, Martinez,

‡International Health Services, East Palo Alto, California, USA

References

- Ching LQ, Borchardt KA, Smith RF, Beal CB. A 24 hour plastic envelope method for isolating and identifying Gardnerella vaginalis (PEM-GVA). Genitourin Med 1988;64: 180-4.
- 2 Piot P, van Dyck E, Totten PA, Holmes KK. Identification of Gardnerella vaginalis. J Clin Microbiol 1982;15:19-24.
- 3 McFaddin JF. Biochemical tests for identification of medical bacteria. 2nd ed. Baltimore: Williams & Wilkins, 1980:4-12.
- 4 Scott TG, Smyth CJ, Keane CT. In vitro adhesiveness and biotype of Gardnerella vaginalis strains in relation to the occurrence of clue cells in vaginal discharges. Genitourin Med 1987:63:47-53.

TO THE EDITOR, Genitourinary Medicine

Acute urethritis due to Neisseria meningitidis group A acquired by oro-genital contact: case report

Sir

Following the recent report of urethritis due to Neisseria meningitidis, acquired from heterosexual oro-genital contact, we wish to report a similar case. A 16 year old schoolboy was referred by his GP in January 1989. He gave a history of pain in the left iliac fossa radiating to the groin for one day. He denied any urethral symptoms, and maintained that he had never had sexual intercourse. There was no significant past medical history. On examination he had a tender swelling adjacent to the left testicle and a profuse purulent urethral discharge. Gram negative intracellular diplococci were present on urethral smear.

A presumptive diagnosis of gonococcal urethritis and epididymitis was made and he was treated with 2 g intramuscular spectinomycin and a two-week course of doxycy-