Isolation of intestinal spirochaetes from homosexuals

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SUMMARY Spirochaetes were isolated from rectal swabs of two homosexuals and the faeces of a third, using simple isolation techniques not previously applied to specimens of this type. The ease of culture of these organisms will enable their distribution and pathogenicity to be studied, particularly in relation to their significance in homosexuals.

The sexual transmission of enteric pathogens is recognised, and a high incidence of gastrointestinal parasites has been reported in communities of homosexuals. A non-treponemal spirochaete has been associated with a purulent rectal discharge in a homosexual.

We report the isolation of spirochaetes using simple techniques from rectal swabs and faeces of three homosexuals. The culture medium, blood agar containing spectinomycin, was initially described for the isolation from pig faeces of Treponema hyodysenteriae, the aetiological agent of swine dysentery.

Isolation methods

Rectal swabs were taken from passive homosexual males attending the Venereal Disease Clinic at the General Infirmary at Leeds. The swabs, transported in Stuart's medium, were inoculated on to 5% horse blood agar containing 400 mg/l spectinomycin. Anaerobic incubation in jars (90% H₂, 10% CO₂) was continued at 37°C for 5 to 14 days. Growth of spirochaetes was detected as spreading, film-like, grey colonies producing complete haemolysis along the lines of spread of the inoculum.

Gram-staining, using dilute carbol fuchsin as a counterstain showed large, irregularly-coiled, Gram-negative spirochaetes (Fig. 1). Electron microscopy, with negative contrast, showed organisms of 0.3-0.5 μm diameter by 10-20 μm with pointed ends, containing four or five axial fibrils (Fig. 2).

Results

Spirochaetes were isolated from three of 25 rectal swabs, after five days incubation. We failed to isolate spirochaetes from 74 diarrhoeal faeces sent to the diagnostic laboratory but did culture the organism from one specimen of faeces from a homosexual. The four isolates of spirochaetes were from three patients, and their cases are presented below.

CASE 1

A 34-year-old homosexual, with a past history of anal herpes simplex virus and syphilis, attended the Venereal Disease Clinic for a "check-up" after a holiday in New York. Proctoscopy was normal and there were no gastrointestinal symptoms. Faecal examination for ova, cysts and parasites was negative, but spirochaetes were cultured from the specimen. A rectal swab taken at the same time was negative. Neisseria gonorrhoeae was not isolated and...
Fig. 2. Negatively stained electronmicrographs. In A, four axial fibrils are seen. Original magnification $A \times 74,000$, $B \times 18,500$.

Syphilis serology indicated past infection. No chemotherapy was given and no spirochaetes were cultured from a rectal swab taken four months later.

Case 2
A 38-year-old homosexual attended with urethral gonorrhoea. Cultures for *N gonorrhoeae* from rectum
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and pharynx, and syphilis serology were negative. Proctoscopy was normal and there were no gastrointestinal symptoms. Spirochaetes were grown from a rectal swab. The patient received 3.5 g ampicillin and 1 g probenecid, followed by a 14-day course of oxytetracycline 250 mg, four times daily, for postgonococcal urethritis. A rectal swab taken 14 days after the first was still positive for spirochaetes.

CASE 3

A 37-year-old homosexual with a past history of treated syphilis and gonorrhoea attended for a "check-up" after a holiday in Athens. Proctoscopy was normal and there were no gastrointestinal symptoms. Spirochaetes were cultured from a rectal swab. Cultures for N gonorrhoeae were negative and syphilis serology indicated past infection. No chemotherapy was given and no further specimens obtained.

Discussion

It is difficult from the three cases described to assess the significance of colonisation by spirochaetes. Further studies are required on these easily cultured organisms. Previous descriptions of isolation techniques for intestinal spirochaetes have been complex, as strict anaerobiosis and special growth factors were thought to be necessary, but with the introduction of the technique described, it will be possible to assess the significance of these organisms in venereal disease and in bowel disorders.

Electron microscopy has been used to demonstrate spirochaetes in the intestines of healthy individuals and in patients with a variety of gastrointestinal disorders. An association between intestinal spirochaetosis and diarrhoea has been reported: however, we have so far failed to isolate spirochaetes from cases of diarrhoea.

The organism we isolated morphologically resembles Treponema vincentii, but the structure of spirochaetes is variable and may be altered by culture. Cultural characteristics resemble those of Treponema hydysenteriae but the isolate from case I was non-identical with pathogenic and non-pathogenic porcine spirochaetes, by growth-inhibition testing (Dr R Lemcke, personal communication, 1981).

Further studies are in progress to characterise these organisms more fully, to determine their pathogenicity and estimate their incidence in homosexuals and in heterosexuals.

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References


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