Lymphogranuloma venereum of the rectum in a homosexual man

Case report

ADRIAN MINDEL
From the Academic Department of Genitourinary Medicine, Middlesex Hospital Medical School, London

SUMMARY A male homosexual presented initially with bloody diarrhoea and a swelling in the left groin, which was unsuccessfully treated with erythromycin. Examination in hospital showed a rectal mass and an abscess in the left groin. Histological examination of the rectal mass and a positive lymphogranuloma complement fixation test result confirmed the diagnosis of lymphogranuloma venereum. This disease, although rare, should not be forgotten in the differential diagnosis of rectal problems in male homosexuals.

Introduction

Lymphogranuloma venereum (LGV) is a tropical sexually transmitted disease (STD). Only 35 cases were seen in the United Kingdom in 1980, most of these infections having been acquired abroad. Several recent reports from the United States have shown that rectal LGV is now being found among male homosexuals. We report a case of homosexually acquired LGV in the United Kingdom.

Case report

In December 1981 a 27 year old homosexual clerk noticed diarrhoea, which lasted a week and was followed in early January 1982 by constipation, tenesmus, and rectal bleeding. He attended his general practitioner and was treated symptomatically for haemorrhoids. Though this eased his symptoms they continued intermittently. In mid-January he developed a swelling in the left groin which was treated unsuccessfully with erythromycin by his general practitioner. He was admitted to the Hammersmith Hospital in February.

Examination showed a 6 x 3 cm fluctuant non-tender abscess in the left groin, which was incised under anaesthesia. Sigmoidoscopy performed at the same time showed an irregular mass situated at 2-5 cm from the anal margin and covered by thick mucosa which bled profusely. Histological examination showed a severely but patchily inflamed rectal mucosa with a heavy infiltration with inflammatory cells of all types especially plasma cells and several ill-defined histiocytic granulomata with multinucleated giant cells. The nature and location of the granulomata were considered atypical for Crohn's disease, and syphilis was considered a likely diagnosis. He was therefore referred to the department of genitourinary medicine at the Middlesex Hospital where he had previously been treated for rectal gonorrhoea and perianal warts in 1980.

He was seen at the Middlesex Hospital in March and stated that his only sexual contact had been with a Jamaican in November 1981. He had been both the active and the passive sexual partner and had also had oro-genital contact. Examination showed an unhealed wound of 3 cm in the left groin; the findings on proctoscopy were as before.

SPECIAL INVESTIGATIONS

A rectal smear showed numerous polymorphonuclear leucocytes and no Gram negative diplococci. Rectal culture results for Neisseria gonorrhoeae, herpes simplex virus, and Chlamydia trachomatis were negative. Cultures from the groin wound for C. trachomatis and bacteria also gave negative results. Treponemes were not seen on dark ground microscopy of material from the rectal mass; serological test results for syphilis were negative. Ova, cysts, and parasites were not seen on stool.
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microscopy. The erythrocyte sedimentation rate was 2 mm in the first hour and the white blood cell count 11 x 10^9/l (70% neutrophils). Estimation of urea, electrolytes, and liver function and a barium enema also showed no abnormalities. The lymphogranuloma complement fixation test (LGV-CFT) was positive at a titre of 512, which is compatible with an acute LGV infection.

TREATMENT AND FOLLOW UP
The patient was treated with oral oxytetracycline 500 mg four times daily for three weeks. The inguinal operation site healed within a week and the rectal mass completely resolved after three weeks. After three months the LGV-CFT was still positive at a titre of 32, and the serological test results for syphilis remained negative.

Discussion
Chlamydial infection of the rectum causes a wide range of symptoms. Many patients are asymptomatic, some have a diffuse proctitis, and a few with LGV have granulomatous masses in the rectum.4 Though rectal LGV has been recognised for many years, early reports of cases were predominantly in women and were considered to be a late complication of genital infection, either by lymphatic spread or “spill” during sexual intercourse.5 Cases of homosexually acquired LGV were reported from Seattle4 in three men with proctitis. The clinical features of these cases were similar to the present one, though they differed in one respect. Whereas the patients in Seattle were promiscuous homosexuals our patient had had sexual contact with only one partner in the previous three months.

The differential diagnosis of inflamed rectal mucosa with or without masses or ulcers in homosexuals includes Crohn’s disease, amoebiasis, syphilis, gonorrhoea, carcinoma, ulcerative colitis, and infection with herpes simplex virus, Yersinia spp, Campylobacter spp, and Clostridium difficile. Though LGV is extremely rare in Britain1 it should not be forgotten as part of the differential diagnosis.

References