Pathogenic microbial flora of genital ulcers in Sheffield with particular reference to herpes simplex virus and Haemophilus ducreyi

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SUMMARY The pathogenic microbial flora of genital ulcers in 161 (80 men and 81 women) unselected patients was studied prospectively. In only one case was Treponema pallidum responsible whereas herpes simplex virus was considered to be the cause of 130 (80·8%) genital ulcers. H ducreyi was isolated from 46 (28·6%) patients, most commonly as a secondary pathogen in herpetic lesions. Two or more pathogens were isolated from the ulcers in 67 (41·6%) patients, and in 21 (13%) patients no pathogens were isolated.

Our results indicate an urgent need for antiviral treatment to reduce the local reservoir of genital herpes, challenge traditional concepts about the prevalence of H ducreyi in Britain, and call for a reappraisal of its role in the causation of genital ulcers.

Introduction

Recent studies1 2 (unpublished data) of the microbial aetiology of genital ulceration show notable differences, particularly as regards the causes of non-syphilitic ulcers. This may partly reflect differences in technique but also genuine geographical differences in the incidences of genital pathogens. Whereas chancroid is considered to be endemic in tropical countries3 and genital herpes is infrequently reported, the converse is true in developed Western countries. In Britain, where chancroid has until recently been rarely diagnosed, and even then usually as an imported infection, the incidence of genital herpes has risen to epidemic levels in the past decade. The advent of reliable cultural techniques4 5 for the isolation of Haemophilus ducreyi in recent years has resulted in the reporting of localised outbreaks of chancroid in non-tropical countries6 8; recently we reported 22 cases in Sheffield9 which were confirmed by culture using a modified haemin-containing medium.

In this study, we examined the pathogenic flora of genital ulcers in men and women presenting to the department of genitourinary medicine, Royal Infirmary, Sheffield, over a 12-month period.

Patients and methods

One hundred and sixty-one unselected patients (80 men and 81 women), who presented for the first time with discrete genital ulcers between 1 December 1980 and 30 November 1981, were routinely assessed. Exudate from the bases of ulcers was examined by darkfield microscopy for the presence of Treponema pallidum and collected with cotton-tipped swabs for both tissue culture inoculation to detect herpes simplex virus and bacteriological culture for Haemophilus ducreyi, other pathogenic aerobic and anaerobic bacteria, and Candida species. In addition, all patients had blood samples taken at their first visit for syphilis serology by the rapid plasma reagin (RPR) and the Treponema pallidum haemagglutination (TPHA) tests and for the detection of herpes simplex antibodies by complement-fixation, ELISA, or neutralisation techniques. In patients returning for follow-up serological tests for herpes antibody were repeated three weeks after the initial visit and for syphilis after three weeks, six weeks, and three months.

HERPES SIMPLEX VIRUS ISOLATIONS Specimens of ulcer exudate obtained with cotton-tipped swabs were immediately inserted into viral transport medium and kept in the clinic refrigerator at 4°C before being transferred to the laboratory, normally within four hours. There they were stored
at −70°C before being inoculated into monolayers of one or more of the following cell lines: African grey monkey (Vero), human epithelium (Hep 2), rabbit kidney (RKB), human embryo lung (Flow 2002), and hamster kidney (BHK 21). Inoculated monolayers were incubated at 37°C for up to 14 days and observed daily for the cytopathic effect of herpes.

In patients with negative culture results for herpes a fourfold rise in antibody titre in the convalescent sera was considered to be indicative of recent herpes simplex infection. Primary and recurrent episodes of herpes infection were distinguished by analysis of the patient’s history, the severity and extent of the herpetic lesions, and the results of herpes antibody tests. Primary infections were diagnosed when there was no history of genital ulceration, there were numerous extensive skin lesions—usually bilateral and often accompanied by cervical or urethral lesions—and antibody titres were low or absent on initial serological testing. In contrast, patients with recurrent herpes almost always gave a history of episodes of genital ulceration, had few lesions usually confined to the skin, and had pre-existing antibodies at their first presentation usually at a high titre.

**HAEMOPHILUS DUCREYI ISOLATIONS**

Swabs containing ulcer exudate were inoculated directly at the time of examination on to two plates of modified haemin-containing medium, one of which contained 5% horse blood. The constituents of the base medium were as described with the addition of gelatin (Oxoid) in a concentration of 4 g/l. Inoculated plates were transported to the laboratory, normally within two hours, where they were incubated in an atmosphere containing 5% carbon dioxide at 33°C for up to seven days.

_H. ducreyi_ isolates were initially recognised by their dark-brown colonies, which had a pronounced tendency to remain intact and could easily be distinguished from other organisms on the colourless medium. On the blood-containing medium the colonies produced α-haemolysis. The identity of the organisms was confirmed by Gram-staining, negative catalase, H₂S, indole, and carbohydrate fermentation test results, and positive test results for alkaline phosphatase and nitrate reduction. Our isolates conformed to the taxonomic requirements of Killian and in all respects were identical to a reference strain (NTC 10945 type strain) and only differed from strains isolated from patients with classical chancroid in South Africa (kindly supplied by Dr P Piot, Institute Voor Tropische Geneeskunde, Antwerp, Belgium) and Kenya (kindly supplied by Dr A Ronald, Department of Medical Microbiology, Winnipeg, Manitoba, Canada) in being β-lactamase-negative when tested with chromogenic cephalosporin. All 16 isolates tested by intradermal injection of rabbits gave the pathogenic responses described by Hammond et al.

**AEROBIC AND ANAEROBIC BACTERIAL ANALYSIS**

Swabs of ulcer exudate were normally transferred to the laboratory before being inoculated on to blood agar media (Oxoid blood agar base plus 10% lysed horse blood) and MacConkey’s medium. When a delay greater than two hours between specimen collection and media inoculation was expected, swabs were placed in Amies transport medium. Inoculated plates were incubated for up to seven days both aerobically and anaerobically. Pathogenic isolates were identified using standard methods.

**Results**

Separate analysis of the isolates from ulcers in men and women gave similar results (table). At least one pathogenic agent was found in 140 (87%) patients, and in 67 (41.6%) patients there were two or more pathogenic organisms in their ulcer flora.

| Pathogenic microbrial flora isolated from the genital ulcers in 161 patients |
|-----------------------------|-----------------|------------------|
| **Men** (n = 80) | **Women** (n = 81) | **Total** (n = 161) |
| Herpes simplex virus | 52 | 63 | 115 (71.4) |
| Treponema pallidum | 1 | 1 | 2 (0.6) |
| Haemophilus ducreyi | 24 | 22 | 46 (28.6) |
| Other aerobic bacteria | 20 | 20 | 40 (24.8) |
| Staphylococcus pyogenes | 6 | 6 | 12 (7.5) |
| β-haemolytic streptococci | | | |
| Group A | 2 | 1 | 3 (1.9) |
| Group B | 8 | 5 | 13 (8.1) |
| Group C | 1 | 1 | 2 (1.2) |
| Group G | 4 | 3 | 7 (4.3) |
| Enterococci | 1 | 5 | 6 (3.7) |
| Coliforms | 3 | 5 | 8 (5.0) |
| Neisseria gonorrhoeae | 1 | 1 | 2 (1.2) |
| Anaerobic bacteria | 5 | 4 | 9 (5.6) |
| Bacteroides species | 3 | 3 | 6 (3.7) |
| Anaerobic cocci | 3 | 2 | 5 (3.1) |
| Candida species | 2 | 14 | 16 (9.9) |
| No pathogens isolated | 13 | 8 | 21 (13.0) |

**HERPES SIMPLEX VIRUS**

Herpes simplex virus was the most commonly isolated pathogen; in 52 patients it was the sole pathogen. In addition to the 115 patients in whom direct virus isolation from ulcers was successful, 15 other patients had a history of recurring genital ulceration with at least one episode of genital herpes confirmed by culture before their presentation or serological evidence of current herpes simplex infection or both. In these culture-negative
patients it is likely that the virus was the initiating cause of their genital ulceration. Thus, a total of 130 (80-8%) patients had genital herpes, of whom 78 had primary episodes and 52 recurrent episodes.

**TREPOEMA PALLIDUM**

*T pallidum* was the cause of genital ulceration in only one patient; he had acquired his infection in Mexico. No other patients had clinical, microscopical, or serological evidence of syphilis either at initial presentation or at subsequent follow-up.

**HAEMOPHILUS DUCREYI**

The commonest bacterial pathogen was *H ducreyi*, but in almost all of 46 patients infected with this organism there was evidence of a separate initiating cause for their genital ulcers; in 38 patients the organism was found in the lesions of genital herpes. Although *H ducreyi* was the sole pathogen isolated from ulcers in 10 patients, seven of these had a known history of recurrent genital herpes and the other three had secondary infection of ulcers caused by trauma, Behçet's syndrome, and excoriated penile psoriasis. The frequency of *H ducreyi* isolation was not appreciably different in patients with either herpetic or non-herpetic genital ulceration. The organism was isolated in 24 (30-8%) of 78 patients with primary herpes, 14 (26-9%) of 52 patients with recurrent herpes, and eight (25-8%) of 31 patients with non-herpetic ulceration.

*H ducreyi* was found in the ulcers of nine (47%) of 19 patients who had had sexual contacts in countries outside Britain in the four weeks before presentation compared with 37 (26%) of 142 who had had local contacts only; most *H ducreyi* infections had apparently been acquired locally. Moreover, in only two patients had there been contact with a known case of chancroid. Although ulcers containing *H ducreyi* were invariably tender with a purulent or necrotic base clinical features varied. Multiple ulcers (74%) and lymphadenitis (70%) were common in patients with *H ducreyi* infection but there was no case of bubo formation in this series.

**OTHER AEROBIC BACTERIA**

Other important aerobic bacteria were isolated from genital ulcers in 40 patients, but in only eight were they the sole pathogens isolated.

Only *Staphylococcus pyogenes* and β-haemolytic streptococci appeared to be primary pathogens. In the one patient in whom *Neisseria gonorrhoeae* was isolated from a vulval ulcer, herpes simplex virus and *H ducreyi* were isolated from the same genital lesion. It is, therefore, unlikely that she had true cutaneous gonorrhoea.

Anaerobic bacteria were never found in the absence of other microbiological causes of genital ulceration but may have had a secondary pathogenic role in some patients.

*Candida* species appeared to be secondary invaders of herpetic lesions in 11 patients but in five they may have initiated genital ulceration by causing pruritus which lead to subsequent excoriation.

**Discussion**

In common with other recent studies we have found that genital ulcers often have a polymicrobial flora and frequently contain two or more pathogens. We found no obvious sex bias in the distribution of the major pathogens.

Herpes simplex virus was the most common isolate and probably the initiating cause of genital ulceration in 81% of our patients. This is a much higher frequency than in other studies and contrasts with 31-40% reported in North American studies,1 14 and only 4% in Kenya.2 Although the rapid increase in locally diagnosed cases of genital herpes during the past 10 years results partially from increased clinical awareness and improved methods of virus isolation, in the main the rise simply reflects the natural history of a condition where recurrences are common and effective treatment lacking. In Sheffield we have also found a high prevalence of HSV type 1 isolates5 (possibly related to changes in sexual practice), contributing to the ever-increasing reservoirs of genital herpes in the community.

Infectious syphilis is rare within our relatively stable population because contact tracing and effective treatment of the few sporadic cases seen in recent years has rapidly removed the local reservoir of infection. It is interesting to compare the single case of imported treponemal ulceration in this study with the reported frequencies of 17% in Denver,1 12% in Seattle (unpublished data), and 11% in Kenya.2

*H ducreyi* was isolated from the genital ulcers of 28% of our patients, which compares with 1-2% in recent American studies1 and 60% in Kenya.2 In most infected patients the organism appeared to be a secondary invader rather than a primary pathogen. The sex distribution of infections was about equal and the great majority were locally acquired. As only two cases were linked, clearly our results were not simply due to a local outbreak of chancroid, although a local reservoir of individuals with asymptomatic genital carriage of *H ducreyi* may well exist. This view conflicts with the observations of others,2 6 16 but evidence to support our contention was found in this study. In three men, all of whom presented initially with ulcers infected both with
herpes simplex virus and H. ducreyi, we were able to re-isolate H. ducreyi from the preputial swabs after their ulcers had completely healed. All had been treated with saline washes alone. In a separate study, we have isolated H. ducreyi from endocervical swabs taken from 11 women and the preputial swabs from three men, most of whom had no history of genital ulceration and none of whom had current genital ulcers.

Organisms resembling H. ducreyi have been seen on smears from chancreoid ulcers in previous studies, and recently Ursi et al. described a “ducreyi-like bacillus”, which they isolated from genital ulcers in 25 of 155 patients in Swaziland. This organism differed from H. ducreyi in that it preferred a microaerophilic atmosphere, did not require X factor for growth, produced hydrogen sulphide and xylose fermentation, but gave a negative result to the alkaline phosphatase test. In another recent study Caine et al. (unpublished data) suggested that Haemophilus influenzae and Haemophilus parainfluenzae may cause some genital ulcers. We did not routinely examine stained smears from ulcers, because of the low specificity and sensitivity of this investigation, and are satisfied that our isolates were indeed H. ducreyi, which have a high requirement for haemin and share identical cultural and biochemical characteristics with test strains of the organism. Our local strains may possibly have had a lower pathogenicity in humans than those found in the tropics, although we have so far failed to show any diminished pathogenicity in rabbits. We are aware that strains isolated from typical cases of chancreoid in Africa are frequently β-lactamase producers whereas our local strains are not; we are, therefore, investigating them for other possible differences in plasmid content and antimicrobial sensitivity. Although our methods for isolating other aerobic and anaerobic bacteria were less sensitive than in other studies, which detected commensals as well as pathogenic organisms, we nevertheless found bacteria other than H. ducreyi, which were fairly common secondary invaders of ulcers caused by herpes simplex virus and trauma but were relatively rare as primary pathogens. In some cases antibiotic treatment was necessary to eradicate these organisms and so ensure healing of the ulcer.

This study shows that the most frequent initiating cause of genital ulceration in Sheffield is herpes simplex virus and that herpetic lesions are frequently subject to secondary infection with H. ducreyi and other bacteria. The immediate and long-term morbidity associated with genital herpes makes accurate diagnosis essential, and the increasing local reservoir of genital herpes in this community necessitates the urgent development of effective antiviral treatment.

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References