Sexually transmitted diseases in rape victims

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Abstract
From 1 January 1986 to 1 September 1989 124 women presented to the Ambrose King Centre (the department of genitourinary medicine of the London Hospital) alleging rape. Sexually transmitted diseases were found in 36 (29%) women (excluding candidosis and bacterial vaginosis). The commonest organisms detected were Neisseria gonorrhoeae and Trichomonas vaginalis, each being present in 15 patients. Eleven women had genital warts. Chlamydia trachomatis was isolated in six patients, two had herpes simplex virus infection and one patient had pediculosis pubis. Serological evidence of past hepatitis B infection was detected in five women and one patient had antibodies to human immunodeficiency virus. Eighteen of the 36 women (50%) had multiple infections. Six women had abnormal cervical cytology smears, three being suggestive of cervical intraepithelial neoplasia grades II–III. Although it is rarely possible to attribute infection to an assailant, these patients require further counselling, treatment and review. Rape victims are thus a population at risk of having sexually transmitted diseases and screening should be offered.

Introduction
The Sexual Offences Act (Amendment), 1976, in the UK defines rape as sexual intercourse with a woman without her consent, either by using force or fraudulent means. Vaginal penetration to any degree is sufficient. The risk of acquisition of sexually transmitted diseases (STDs) after sexual assault is unknown. A recent review highlighted the lack of data on the association between rape and STD, particularly conditions other than the traditional venereal diseases gonorrhoea and syphilis. Much of the available information has come from the microbiological assessment of women referred by police agencies in North America where the legal definitions of rape and child sexual abuse differ from those in the UK and where prophylactic treatment is more readily available. These studies have not always differentiated children from adults, girls aged less than 16 years representing up to 42% of victims in one study. More recently, infection with Chlamydia trachomatis has also been reported following rape.

The viral STDs, namely human papilloma virus (HPV), herpes simplex virus (HSV), cytomegalovirus (CMV), hepatitis B virus (HBV), human immunodeficiency virus (HIV) and molluscum contagiosum may be acquired through rape. However, only two studies and one case report have considered the risk of acquisition of these infections. In a study of 46 women one had a four week history of genital warts, having attended 12 months after rape. No voluntary sexual activity had occurred between the assault and her initial visit. HSV infection was not reported in these patients and none were tested for antibodies to HIV. In a more recent study, 204 women were reviewed within 72 hours of rape by an unknown assailant or assailants. No cases of genital warts were reported. However, 13 of 199 women had evidence of koilocytosis on cervical cytology at their initial visit. HSV was isolated in four of 170 women at presentation. One of 123 patients who had antibodies to HIV detected at her first attendance. No additional viral infections occurred in 52 patients during follow up ranging from two to 40 weeks, with a mean of eight weeks. The authors concluded that viral STD were not likely to be acquired as a consequence of sexual assault. Screening for markers of HBV infection was not carried out in either study. In addition, seroconversion to HIV has been reported in one woman within three months of a rape, where no other identifiable risk factors were present.

The difficulties in diagnosis and management of patients with viral STD have increased the workload of departments of genitourinary medicine in England. They require extensive counselling and follow up which may need to be intensified if the infections are diagnosed after rape. There is no simple confirmatory test for the clinical diagnosis of genital warts and the serological diagnoses of HBV and HIV infections may be associated with a lag period before a detectable response. The duration of infection is often impossible to establish particularly in relation to the timing of a rape. Genital warts may have a lengthy incubation period of up to nine months. HSV infection may be inapparent initially.
and remain latent until reactivation. Duration of infection may sometimes be estimated by document-
ing seroconversion in HSV, HBV and HIV infection.

The management of established infection with the
viral STDs is unsatisfactory. The associated long
term complications of these infections are an
additional burden on the rape victim. HPV is
associated with ano-genital neoplasia, \(^1\) HBV may
result in chronic liver disease and carcinoma and
HIV is complicated with life threatening opportunist-
ic infections and tumours.

The objective of this study was to assess the overall
frequency of STDs in rape victims who presented to
a department of genitourinary medicine.

**Methods**

Between 1 January 1986 and 1 September 1989 all
women attending the Ambrose King Centre, the
department of genitourinary medicine at the London
Hospital alleging rape were assessed according to a
standardised protocol. A detailed history of the
incident was taken, where appropriate, together with
a sexual and medical history. Signs of assault and
infection were documented at examination.

Material from the posterior fornix of the vagina
was Gram-stained for candidal pseudohyphae/
splaces and "clue cells" as well as suspended in saline
and examined microscopically for *Trichomonas
vaginalis*. Candida species were also cultured from
the vaginal material in Sabouraud's medium and
*T. vaginalis* infection was confirmed by acridine
orange staining. Bacterial vaginosis was diagnosed by
the presence of "clue cells", positive amine test
and a vaginal pH of greater than 5.5. Gram-stained
smears from the urethra, endocervix and rectum
were examined for Gram-negative intracellular
diplococci. Material from the urethra, endocervix
and rectum and, if appropriate, the pharynx, were
directly plated on to brain heart infusion agar for
culture for *Neisseria gonorrhoeae*. Isolates were
identified on sugar solid media. Material from the
same sites were placed in 2-sucrose phosphate media,
and transported in liquid nitrogen to the laboratory
for culture for *C. trachomatis*. Cervical cytological
smears were performed in all the women. Genital
warts were clinically diagnosed. Cultures for herpes
simplex virus were taken as clinically indicated.
Specimens were transported in Viroluct transport
media (Medical Wire and Equipment Co Ltd) and
cultured in human embryonic lung cells. Evidence of
cytopathic effect was noted at weekly intervals for
three weeks.

A bimanual pelvic examination was performed.
Serological tests for syphilis and HBV were carried
out at the initial assessment. HBV serology was
performed using the Wellcozyme method. Serum for
antibodies to HIV was tested by ELISA methods and
confirmed using gel particle agglutination. An
additional serum sample was saved for possible
paired testing at a later date. Tests for antibodies to
HIV were taken only after counselling and informed
consent.

Follow-up was arranged after two weeks and three
months when Gram-stained smears and cultures for
*N. gonorrhoeae* and *C. trachomatis* were repeated.
Serological tests were also repeated at the last visit. A
third visit was not arranged if the initial assessment
was three months or more from the rape. Counselling
was initially provided by the clinic health advisers
who also informed the patients of other support
agencies and made referrals, where appropriate.

**Results**

**Epidemiology**

Of 141 women attending the Ambrose King Centre
from 1 January 1986 to 1 September 1989 following
sexual assault, 124 women (mean age 26 years,
range 16–45 years) alleged that rape had occurred.
This took place in Britain in 110 cases (89%),
Mediterranean countries in eight (6.5%), Africa in
four (3%), and North America in two (2%). One
hundred and five victims (85%) were Caucasian, 14
(11%) were Afro-Caribbean and four (3%) were
from other ethnic backgrounds. Fifty seven (46%)
presented within two weeks of the assault (range of
one day to six years). The number of clinic visits
made by the patients ranged from one to seven. After
the initial visit, 104 women (84%) attended for a first
follow-up at a mean of 31 days (range 3 days to 96
weeks) and 76 (61%) attended for a second review at
a mean of 11 weeks (range 1–100).

Most of the women, 69 (56%), were self-referred.
Other sources of referral are shown in table 1. The
rape had been reported to the police by 58 women

**Table 1** Referral patterns of rape victims

<table>
<thead>
<tr>
<th>Referral agency</th>
<th>Number of patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Self</td>
<td>69 (56)</td>
</tr>
<tr>
<td>General practitioner</td>
<td>29 (23)</td>
</tr>
<tr>
<td>Police</td>
<td>20 (16)</td>
</tr>
<tr>
<td>Voluntary agencies</td>
<td>6 (5)</td>
</tr>
<tr>
<td>Total</td>
<td>124 (100)</td>
</tr>
</tbody>
</table>

**Table 2** Epidemiological features of the rape

<table>
<thead>
<tr>
<th>Feature</th>
<th>Number of patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>No prior sexual experience</td>
<td>7 (6)</td>
</tr>
<tr>
<td>Known assailant</td>
<td>55 (44)</td>
</tr>
<tr>
<td>Oral penetration*</td>
<td>14 (11)</td>
</tr>
<tr>
<td>Voluntary intercourse after rape</td>
<td>50 (40)</td>
</tr>
</tbody>
</table>

*In addition to vaginal penetration.
Further epidemiological features of the assault are shown in Table 2. The rape was their first sexual experience for seven women. Fifty (43%) of 117 women with prior sexual experience had voluntary sexual intercourse with another partner after the assault. A past history of rape was given by four women (3%) and child sexual abuse by three women (2%). Twenty four women (19%) had been raped by more than one assailant and 13 (10%) had been subjected to anal penetration (in addition to vaginal penetration). Five women had evidence of vulval injury, all presenting within two weeks of the rape.

**Sexually transmitted infections**

The frequency of sexually transmissible infections is shown in Table 3. Overall, 36 (29%) women had 57 separate STDs. Vaginal candidosis and bacterial vaginosis were diagnosed in 22 and seven women respectively but are excluded from these data. The commonest organisms detected were *N. gonorrhoeae* (no penicillinase producing organisms were isolated) and *T. vaginalis*, each being present in 15 patients. Genital warts were diagnosed in 11 women and *C. trachomatis* was found in six women. Of 123 women tested none had positive serological tests for syphilis. Clinical features of pelvic inflammatory disease were present in six women, of whom two had gonococcal cervicitis, one had trichomoniasis and one had bacterial vaginosis. No pharyngeal infections were associated with a history of oral penetration in 14 women. Anal penetration was associated with one case of rectal gonorrhoea. Of the seven women with no prior sexual experience, one had trichomoniasis and the other had candidal infection.

Of the two women with HSV infection one had a history of recurrent vulval disease and was seen four years after the rape. The timing of the primary attack was unknown. The other was seen eight days after being raped but had had symptoms for two weeks localised to the labia minora.

Antibodies to hepatitis B surface antigen (HBsAg) were detected in four women. Three were tested within two months of being raped, the fourth after six months. A fifth woman was both HBsAg positive and had antibodies to hepatitis B e antigen, being tested 18 days after the rape. Three of these women were from countries of high HBV prevalence and two were ex-injecting drug users. No cases of acute seroconversion were reported during follow up. Of the 87 women who had negative HBV serology, 37 (43%) were either initially tested or retested at three months or more.

The single patient (of 44 tested) with antibodies to HIV presented as a contact of her regular male partner who had gonorrhoea. She had been raped abroad two months previously. No oral or anal penetration had occurred. Samples from both her initial and two month follow up visits showed antibodies to HIV. Her partner remained HIV antibody negative and she denied other sexual contacts or risk factors. Overall, 24 (19%) patients expressed anxiety about HIV infection as a consequence of rape (Table 4). In eight women it was the reason for their initial attendance. Seven of this group declined HIV testing after counselling. HIV tests were performed at a mean of eight months (range one month to four years) in 43 women where the result was subsequently negative. After pre-test counselling, 42 women (34%) declined to proceed with the HIV test.

Eighteen of the 36 women (50%) with an STD had multiple infections (Table 5). Dual infections were present in 15 women and three had triple infections.

For 15 (12%) women an STD was found on follow up which had not been detected at their initial assessment. Four of these women had not had further intercourse from the time of the rape to the detection of the infection. The first woman had chlamydial infection which was detected at three months and was not found at her initial visit three weeks from the rape. The second woman presented with pediculosis pubis at five months, this infestation not being detected at her initial visit five days from the rape. The other two women had genital warts. Of the 11 women who had had intervening sexual intercourse four had gonorrhoea, three had trichomoniasis, one had both and three had genital warts.

**Table 3 Overall frequency of sexually transmissible infections in victims of rape**

<table>
<thead>
<tr>
<th>Sexually transmissible infections</th>
<th>Number of patients* (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gonorrhoea</td>
<td>15 (12)</td>
</tr>
<tr>
<td>Trichomoniasis</td>
<td>15 (12)</td>
</tr>
<tr>
<td>Genital warts</td>
<td>11 (8)</td>
</tr>
<tr>
<td>Chlamydial infection</td>
<td>6 (5)</td>
</tr>
<tr>
<td>Genital herpes</td>
<td>2 (2)</td>
</tr>
<tr>
<td>Pediculosis pubis</td>
<td>1 (1)</td>
</tr>
<tr>
<td>Hepatitis B markers</td>
<td>5/92</td>
</tr>
<tr>
<td>HIV antibody</td>
<td>1/44</td>
</tr>
</tbody>
</table>

*18 patients with multiple infections.

**Table 4 Women expressing HIV anxiety: HIV antibody tests**

<table>
<thead>
<tr>
<th>Year of study</th>
<th>Number of women</th>
<th>HIV anxiety (%)</th>
<th>HIV tests (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1986</td>
<td>21</td>
<td>3 (14)</td>
<td>8 (38)</td>
</tr>
<tr>
<td>1987</td>
<td>31</td>
<td>6 (19)</td>
<td>14 (45)</td>
</tr>
<tr>
<td>1988</td>
<td>44</td>
<td>12 (27)</td>
<td>18 (41)</td>
</tr>
<tr>
<td>1989*</td>
<td>28</td>
<td>3 (11)</td>
<td>4 (14)</td>
</tr>
<tr>
<td>Total</td>
<td>124</td>
<td>24 (19)</td>
<td>44 (36)</td>
</tr>
</tbody>
</table>

*Data up to 1 September 1989.
Table 5  Multiple sexually transmitted diseases in rape victims

<table>
<thead>
<tr>
<th>Sexually transmitted disease</th>
<th>Genital warts</th>
<th>HBV</th>
<th>HIV</th>
<th>TV</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chlamydial infection</td>
<td>2*</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Gonorrhoea</td>
<td>2*</td>
<td>0</td>
<td>0</td>
<td>8</td>
</tr>
<tr>
<td>Herpes simplex</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>TV</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>8</td>
</tr>
<tr>
<td>Total</td>
<td>8</td>
<td>1</td>
<td>1</td>
<td>8</td>
</tr>
</tbody>
</table>

*One woman had concurrent TV.
†This patient also had antibodies to HBsAg.

contraception was prescribed for 13 women. Three pregnancies were associated temporally with the rapes and in one case the woman had no current sexual partner. Two pregnancies were terminated, the third patient was not followed up after her third visit.

Discussion

STDs (excluding vaginal candidosis and bacterial vaginosis) were present in 36 (29%) of 124 rape victims attending the Ambrose King Centre during the study period. The commonest organisms detected were N. gonorrhoeae and T. vaginalis, each being found in 15 women. Although C. trachomatis is one of the commonest STDs in the United Kingdom it was isolated in only six women. There are many variables in the isolation of C. trachomatis. Culture from a single swab from each site may be associated with a false negative result.12

No women were found to have reactive serological tests for syphilis. Previous studies have shown similar results.3-7  Syphilis is now an uncommon infection in this country, probably due to continued screening and contact tracing. These findings should reinforce current screening policies rather than lead to complacency.13

The most common viral infection was with HPV in the form of genital warts, being present in 11 (9%) patients. In five women the warts were not detected at their initial visit but only on follow up. This suggests that the infection either did not pre-date the rape or that at the initial visit the patient had subclinical disease. Eight of this group had one or more coexistent STD. It is possible that cases of wart virus infection were missed because of the long incubation period and that additional cases may have been detected on extended review. The use of the colposcope may have increased the yield of HPV detection but has the disadvantage of being perceived as an additional trauma by the rape victim. Although DNA typing was not undertaken in this study it may be desirable in providing medico-legal evidence.

Two women had evidence of symptomatic HSV infection. In both cases these were unlikely to have been associated with the rape. However, inapparent HSV infection would not have been detected in this study since viral cultures were only performed on clinical grounds. HSV is often complicated by psychological morbidity.14  This problem could be exaggerated if the infection were acquired following rape.

No acute cases of HBV infection were detected. One low risk carrier of HBV was identified 18 days after the rape. Probably all of the women with HBV markers acquired their infection prior to their assault. This is the first study to present data on HBV in victims of rape. However, if the assailant is

Table 6  Contraceptive methods used by women in the study at the time of rape

<table>
<thead>
<tr>
<th>Method of contraception</th>
<th>Number of patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(n = 124)</td>
</tr>
<tr>
<td>None</td>
<td>70 (56)</td>
</tr>
<tr>
<td>Combined oral contraceptive</td>
<td>25 (20)</td>
</tr>
<tr>
<td>Condom</td>
<td>16 (13)</td>
</tr>
<tr>
<td>Sterilisation</td>
<td>6 (5)</td>
</tr>
<tr>
<td>Intra-uterine contraceptive device</td>
<td>5 (4)</td>
</tr>
<tr>
<td>Diaphragm</td>
<td>1 (1)</td>
</tr>
<tr>
<td>Depo-provera</td>
<td>1 (1)</td>
</tr>
</tbody>
</table>
suspected of being an infective risk, post-exposure HBV prophylaxis should be considered.\textsuperscript{15}

The increasing trend in the number of HIV tests performed from 1986 to 1988 shown in table 5 may have reflected publicity surrounding HIV at that time. Eight women attended solely because they perceived themselves to be at risk of HIV infection following rape, one decided not to be tested after counselling. Of the 24 (19\%) women expressing anxiety about AIDS, seven declined HIV testing after counselling.

Only one woman had antibodies to HIV and she did not present with HIV anxiety. It was not possible to directly attribute this to the rape since antibodies were present at the initial visit two months after the rape had occurred. The HIV status of her assailant was unknown. Supporting data were presented in Jenny's series from Seattle\textsuperscript{a} in which no sero-conversions were recorded in 123 patients, of whom 52 (42\%) returned for a second follow up visit at a mean of eight weeks. A recent case report of HIV seroconversion after rape\textsuperscript{a} has highlighted this potential risk for rape victims. This has prompted discussion about post-rape prophylaxis using zidovudine.\textsuperscript{9} As in the case of needle stick injury,\textsuperscript{16} no data are available on the efficacy of this treatment. The topical use of substances with anti-HIV activity,\textsuperscript{17} such as nonoxynol-9, following rape has been suggested but these products have not been evaluated, do not consider the possibility of multiple sites being involved and may even be harmful.\textsuperscript{18} Discussion about HIV with a woman in such a vulnerable state following rape may not be appropriate in all cases.\textsuperscript{19} Our practice is to save a serum sample at the initial visit to be tested when an informed decision can be made and repeated, as necessary. However, the risk of HIV transmission is small, at least in the low prevalence areas so far studied, but this may well change with time. The risk of HIV transmission may also be affected by the use of condoms by assailants, the occurrence of anal penetration, anogenital trauma and rapes by multiple assailants.

Three of six abnormal smears showed changes suggestive of significant CIN abnormalities. In view of the natural history of CIN and the relatively short time period to detection in this study it is unlikely that these abnormalities were related to the rape. However, they represent important incidental findings in need of assessment which may otherwise have remained undetected.\textsuperscript{20}

The presence of an STD cannot necessarily be attributed to an assault. It is possible that the infection was present before the rape. This could be excluded in one patient with \textit{T. vaginalis} infection who had been a virgin. An important confounding factor in linking an STD to an assailant is the occurrence of voluntary sexual intercourse between the rape and medical review, introducing the possibility of STD acquisition after the rape. Fifty (43\%) of 117 women with prior sexual experience had voluntary sexual intercourse with another partner after the rape. Of those women found to have an STD, five of their sexual partners were traced. All had corresponding infections. However, this does not elucidate the source of the STD. Despite the assailant being known to 55 (44\%) women we did not have the opportunity to screen any of them. If an alleged assailant does present for STD screening it would be prudent for medico-legal purposes to store specimens, in particular gonococcal isolates, which can be both serotyped and auxotyped,\textsuperscript{21} if necessary, at a reference laboratory. These specimens could be compared to stored isolates from the victim to either help corroborate or refute the link between assailant and victim. In a similar way, immunotyping of chlamydial isolates could be performed if facilities permit.

Results of medical investigations may be required as evidence in pending legal cases. Culture of \textit{N. gonorrhoeae} and \textit{C. trachomatis} can be used as firm evidence for the presence of infection. Chlamydial antigen detection methods using immunofluorescence or enzyme immune assay (EIA) are becoming more widely available. They are useful in clinical management for screening populations at risk of chlamydial infection. They should not be used in medico-legal evidence\textsuperscript{22} because of their lower specificity. However, EIA modified with blocking techniques has improved predictive values (Schachter \textit{et al}, International Society for Sexually Transmitted Disease Research, Abstract no. 19, Copenhagen, 1989), but the problem of false positive results still exists.

Intercurrent antibiotics used by 10 women may have masked infection in the six women in whom no STD was detected. The antibiotics used were all potentially active against some STD.

When raped 36 (29\%) women had adequate protection against pregnancy (either using the combined oral contraceptive, an intra-uterine contraceptive device or had been sterilised). Of those with inadequate contraception three became pregnant and two had terminations. Post-coital hormonal contraception had been given to 13 women; it had not been prescribed for the three women who became pregnant because they all presented more than five weeks from the assault. In patients with chlamydial infection pelvic inflammatory disease may result from termination of pregnancy\textsuperscript{23} if these infections are not sought and treated.

The range of STD detected mitigates against the use of prophylactic antibiotic regimes following rape. Thorough investigation and treatment of identified infection is a more satisfactory alternative.\textsuperscript{22} However, epidemiological treatment may be
appropriate if a patient cannot return for follow up and compound regimes have been suggested.\textsuperscript{2}

Screening for STDs is only part of the overall management of rape victims. Psychological and emotional support should be made available. The patients in our study were offered immediate access to health advisers within the clinic for initial counselling. Further tiers of psychological support were provided by other statutory and voluntary agencies as appropriate.

In conclusion, we have shown that rape victims are a population at risk of having STDs. However, in only a minority of cases is it possible to attribute the STD to an assailant. Screening for STDs should now be offered in the management of rape victims. It may also be necessary to make provision for storing specimens for further analysis where medico-legal cases are likely.

We thank all the staff of the Ambrose King Centre for their help in the sympathetic management of the patients reported in this study. We also thank Dr J Treharne of the Institute of Ophthalmology, London, for performing chlamydial isolation.

Address correspondence to Dr S Estreich.

16 Centers for Disease Control. Public health service statement on management of occupational exposure to HIV, including consideration regarding zidovudine postexposure use. MMWR 1990;39(no.RR-1).