Markers of sexually transmitted diseases in seminal fluid of male clients of female sex workers

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Objectives: To screen for certain STD markers in a group of male clients of female sex workers.

Method: Condoms with seminal fluid were collected at 10 “massage parlours” in Copenhagen. The seminal fluid samples were examined for HIV antibodies, markers of hepatitis B virus (HBV), Chlamydia trachomatis, and Mycoplasma genitalium.

Results: All samples (n = 332) were negative for HIV antibodies. Out of 327 samples examined for HBV markers 32 (9.8%) were positive for HBV core antibodies, one of which was also positive for HBV antigen. C trachomatis could be demonstrated in six out of 122 (4.9%) samples and M genitalium in one out of 122 samples.

Conclusions: The finding of a C trachomatis prevalence of 4.9% is considerable higher than expected in men with a presumed age of 35–55 years. The demonstration of a prevalence of HBV markers of 9.8% indicates that these clients have an increased risk of HBV infection, a finding that further consolidates the recommendation of HBV vaccination of sex workers. As shown in this study, STD transmission in commercial sex may also have the client as the source.

Keywords: sex workers; HIV; STD; seminal fluid; condoms

Introduction

Female sex workers are traditionally defined as women who sell sex for money or other goods. This definition, however, implies at least one other part, the person who provides money in exchange for the sexual service—that is, the client. In order to evaluate the relation between commercial sex and sexually transmitted diseases (STD), prevalences of STD should therefore be measured not only in female sex workers but also in clients.

The theoretical risk of HIV transmission from female sex workers to male clients and further on to the heterosexual community has, during the past decade, led to studies focusing upon STD prevalence and sexual behaviour among Danish sex workers1; however, unselected studies in male clients are difficult to perform.

In order to gather information about STD rates in an unselected group of male clients, condoms with seminal fluid were collected from “massage parlours” in Copenhagen and the contents were screened for certain STD markers.

Material and methods

Female sex workers from 10 massage parlours in Copenhagen known by the study group from a previous study agreed to participate.2 Used condoms were tied routinely with a knot and thrown in a plastic bag under the bed. Plastic bags containing the used condoms with ejaculates from clients who had visited the massage parlours within the previous 24 hours were collected in June 1991, January 1992, and January 1994 at the parlours open on these days. The bags were collected on 2 subsequent days in each period in order to minimise the risk of including samples from the same person twice. All sex workers, ranging from two to four women at each massage parlour, participated.

The seminal fluid samples were diluted with an equal volume of phosphate buffered saline (PBS) and centrifuged for 15 minutes at 2000 g. From the first sampling period in 1991–2, comprising 210 specimens, only the supernatant was kept, whereas both the pellet and the supernatant were kept frozen at –20°C from the second sampling period in 1994, comprising 122 specimens.

All samples were analysed for HIV and for markers of hepatitis B virus (HBV) using the supernatant, whereas only the 122 samples from which the pellets were stored could be used for direct detection of pathogens by the polymerase chain reaction (PCR). Pellets from the centrifuged seminal fluid were resuspended in the remaining supernatant and were processed for PCR by adding 100 µl of the material to 300 µl of a 20% w/v suspension of Chelex 100 (Bio-rad) in TE buffer (10 mM TRIS pH 8.0; 1 mM EDTA) and incubating the sample at 95°C for 10 minutes. A volume of 25 µl of the prepared specimen was used in PCR analysis.
HIV testing was preferred to hepatitis B core antibody (antiHBC) testing if the amount of material was insufficient.

HIV ANTIBODIES
The first 210 seminal samples were tested by an in house HIV-1 enzyme linked immunosorbent assay (HIV-1 ELISA), where the working dilution was 1:50 in PBS, and the next 122 samples were tested by the Abbott recombinant HIV-1/HIV-2 third generation enzyme immunoassay (Abbott HIV EIA) after dilution 1:2 in PBS. Positive samples were submitted to HIV-1 western blotting. 3

Paired serum and seminal samples collected from three HIV-1 positive people and from nine HIV negative people served as controls.

By the in house HIV-1 ELISA the positive serum and seminal samples had absorbance values > 2-0 and > 1-0, respectively. The negative serum and seminal samples had mean absorbance values of 0-12 (range 0-03–0-17) and 0-02 (range 0-01–0-07), respectively. The cut off value for seminal fluid was defined at 0-15. The Abbott HIV EIA positive serum and seminal samples had absorbance values > 2-0 whereas the paired negative serum and seminal samples had mean absorbance values of 0-04 (range 0-03–0-11) and 0-06 (range 0-02–0-14), respectively. The cut off value for seminal fluid by this assay was defined at 0-15. Seminal samples with absorbance values above 0-15 were submitted to HIV-1 western blotting.

HEPATITIS B MARKERS
A total of 327 of the 332 samples (98%) were tested for antiHBC using a competitive enzyme immunoassay (Corzyme, Abbott Laboratories, USA). The Corzyme is an enzyme immunoassay for the qualitative determination of total antibody to hepatitis B (antiHBC) in human serum or plasma. The Corzyme is not validated for detection of antiHBC in seminal fluid samples.

In order to use seminal fluid samples as a predictor of antiHBC in serum we tested seminal fluid and serum samples from four men positive for antiHBC in serum. Using the cut off value defined by positive and negative controls supplied by the manufacturer all four serum samples were positive, but all four seminal samples were negative. By titration of the serum samples in order to estimate the titre, two samples had a very high titre of antiHBC, whereas two turned negative after a dilution of 1:5. The seminal samples corresponding to the two high titre serum samples showed an antiHBC optical density (OD) value between the cut off value and the negative control value, whereas the antiHBC OD value of the seminal samples corresponding to the two low titre serum samples were close to the negative control OD value. We defined a new antiHBC cut off for seminal fluid samples as the cut off value (for serum samples) × 1-95.

All seminal fluid samples classified as positive for antiHBC according to the new cut off value were tested for hepatitis B surface antigen (HBsAg) by an enzyme immunoassay (Auszyme, Abbott Laboratories, USA).

CHLAMYDIA TRACHOMATIS
C trachomatis was detected by amplification of the cryptic plasmid using primers CP24/CP27.4 Each reaction tube received an internal processing control containing the primer binding sites in order to assure the validity of negative results. All positive results were confirmed by amplification of the C trachomatis 16S rRNA gene using primers 1A/1B.5

This PCR method was previously validated against C trachomatis culture by investigating 100 seminal samples from men with infertility where it was found to detect 3/3 culture positive plus one additional PCR positive culture negative. The latter specimen was positive also in another laboratory using another PCR technique (unpublished results).

Results
Altogether 332 condoms were collected, 210 condoms in 1991/1992 and 122 condoms in 1994.

HIV ANTIBODIES
All seminal samples from the condoms were tested for HIV antibodies and all were negative.

HBV MARKERS
A total of 337 seminal samples were examined for antiHBC, of which 32 (9-8%) were positive. Among the 32 antiHBC positive samples, one was positive and 31 were negative for HBsAg.

Discussion
In Denmark, with a population of five million inhabitants, an estimated number of at least 5000 sexual transactions involving about 1500 female sex workers take place every day, the majority in massage parlours. According to the female sex workers, male clients are mainly 35–55 years old, most of them are married or cohabiting, representing a broad spectrum of the society. In representative samples of Danish men, 13%–14% have had sexual contact with sex workers once or more in their
life, and 48% of these men have had a history of one or more STD, figures that are in line with those from other European countries.

Behavioural information about female sex workers and clients that are based upon studies among STD clinic attenders may be biased, as STD clinic attenders are selected, may have a higher refusal rate when participating in behaviour studies, and may not be discovered as female sex workers or clients if they do not want to identify themselves as such. When comparing clients recruited at an STD clinic with those recruited outside, a history of STD was much more frequently reported among the STD clinic sample in a study from Glasgow and a study from Amsterdam, and clients with relatively high risk behaviour were strongly represented among the STD clinic sample in one of these studies.

Among STD clinic attenders in Denmark, 17% of the heterosexual men reported to have had sex with a female sex worker, the proportion increased with reported number of sex partners. From the same study, which is an ongoing surveillance study, chlamydial infection was present in 6–1% of those with a history of ever being a client of a female sex worker but, again, this figure gives no information about the number of clients who are infected when the sexual transaction takes place.

Information about STD prevalence among Danish sex workers is from recent studies based upon interview. In one of these studies, 111 out of 253 female sex workers reported STD examination within the past year resulting in the finding of C trachomatis in 8%, M genitalium in 4%, and hepatitis B markers in 1%. The frequency of STD (other than HIV) among these sex workers was correlated with the number of non-paying sexual partners and with the inconsistency in condom use. In contrast, HIV infection is closely related to drug use. HIV infection has not been found in non-drug using sex workers in Denmark, and in only a few cases from other European countries.

Studies focusing upon HIV prevalence among male clients of female sex workers have recruited participants from STD clinics and by advertising. A few cases have been positive but as the clients were not randomly selected they can only give very limited information about the risk of HIV transmission from the male client to the female sex worker. The participants in our study, who were all HIV negative, were selected as to the type of sex workers they visited, but were otherwise unselected, as the sample of sex workers routinely refused clients who would not use a condom.

The risk of HIV transmission by non-drug related sex workers seems, therefore, to be very low in many European countries. The advantage, however, is that it has given rise to a very high level of condom use. Consequently, nowadays, female sex workers rarely transmit HIV or STD.

Men aged 20–29 years account for 65% of all laboratory reported male chlamydial cases in Denmark, those aged 30 years or more account for 19%, and those aged 40 years or more account for only 5%. Studies of the prevalence of chlamydial infections in unselected groups of Danish men, have been performed among those aged 17–26 years already in, or liable for, military service, where 5.7% and 7.9% respectively had urethral chlamydial infection.

The prevalence of chlamydial infections in an unselected group of men aged between 35 and 55 years, which is the presumed age of the clients in this study, is not known, but if this age group accounts for 10–15% of all chlamydial infections a prevalence of 0.6–1.2% should be expected, based upon the reported prevalence among younger men. The finding that C trachomatis could be demonstrated in 4.9% of the seminal samples in this study, therefore, suggests that these men had a four to eight times higher rate of infection than the general population based on the above calculations. Previous studies were based on urine sampling and enzyme immunoassays, whereas this study used the more sensitive PCR technique, a fact that may overestimate the difference between the expected and the observed prevalence. Furthermore, the fact that the rather informal sampling method could have caused some sample to sample carryover cannot be totally excluded. However, since we did not find more than one M genitalium PCR positive specimen this threat to the validity of the results may be minor.

M genitalium has been found in men with non-gonococcal, chlamydia negative urethritis, and a causal connection has been suggested. The prevalence of this microorganism in asymptomatic men has, however, not been elucidated, so it is difficult to relate the finding of this microorganism in one of the samples to other studies.

The presence of antiHBc together with a negative test for HBsAg is a marker of a past but now non-transmissible infection. The finding of an antiHBc prevalence of 9–8% may be considerably underestimated as antiHBc could only be detected in the seminal samples from two controls with high antiHBc serum titres, whereas it was undetectable in two controls with low serum titres. Denmark is a low endemic area for HBV. In spite of this, the measured prevalence was at least twice as high as that found in Danish blood donors, in the same range as is found among heterosexual STD clinic attenders born in Denmark, but lower if the STD clinic attenders were from intermediate and high endemic areas. Therefore, it seems reasonable to conclude that the clients from whom the seminal samples in this study originated had an increased risk of being HBV infected, and so female sex workers should consider HBV vaccination.

HBV is the only STD for which a protective vaccine has been developed, whereas the only protection against other STDs is the consistent use of condoms.

It is well documented that intact latex condoms provide a continuous mechanical barrier
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