



Published in final edited form as:

AIDS Behav. 2015 July ; 19(7): 1305–1316. doi:10.1007/s10461-015-1002-5.

Risk factors for HIV acquisition in high risk women in a generalised epidemic setting

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Abstract

In South Africa young women bear a disproportionate burden of HIV infection however, risk factors for HIV acquisition are not fully understood in this setting. In a cohort of 245 women, we used proportional hazard regression analysis to examine the association of demographic, clinical and behavioural characteristics with HIV acquisition. The overall HIV incidence rate (IR) was 7.20 per 100 women years (wy), 95% Confidence Interval (CI) 4.50–9.80]. Women 18–24 years had the highest HIV incidence [IR 13.20 per 100 wy, 95% CI 6.59–23.62] and were almost three times more likely to acquire HIV compared to women 25 years and older [adjusted Hazard Ratio (aHR) 2.61, 95% CI 1.05–6.47]. Similarly, women in relationships with multiple sex partners had more than twice the risk of acquiring HIV when compared to women who had no partner or who had a husband or stable partner (aHR 2.47, 95% CI 0.98–6.26). HIV prevention programmes must address young women's vulnerability and sex partner reduction in this setting.

Keywords

Risk factors; HIV acquisition; women

Introduction

Into the fourth decade of the Human Immunodeficiency Virus (HIV) epidemic, South Africa has an estimated 6.1 million people living with HIV and contributes 17% of the global HIV burden, yet accounts for only 0.7% of the world's population (1, 2). With the majority of infections heterosexually acquired, young women bear a disproportionate burden of HIV. Sentinel surveillance among pregnant women shows that in 1990, HIV prevalence was 0.7% and by 1998 increased exponentially to 22.8%, peaking at 30.2% in 2010 (3). Data from national population based surveys show similar trends with high HIV prevalence in young women compared to young men (4, 5). HIV prevalence peaked at 36.0% among women aged 30–34 years and at 28.8% among men in the 35–39 year-old age group (5). HIV prevalence in women aged 15–19 years was 5.6%, eight times higher than males in the same age group, and increased to 17.4% among women aged 20–24 years and 28.4% in the 25–29 year age group (5). Within South Africa there is significant geographical variation in the distribution of HIV and the province of KwaZulu–Natal remains the worst affected. Whilst prevalence of HIV has been useful to understand the evolving epidemic, HIV incidence rate is a more sensitive measure to monitor the epidemic. Amongst rural and urban women the HIV incidence rates have been 6.5 and 6.4 per 100 women years (wy) respectively, and in young women <18 years 4.7 per 100 wy (6, 7). These data underscore the need to understand the risk factors contributing to these high HIV prevalence and incidence rates.

The key risk factors for HIV in the South African setting include a combination of structural, behavioural and biological factors. Poverty, labour migration, urbanisation, gender inequalities and gender-based violence contribute to an increased susceptibility to HIV (8, 9). At the individual level, being single, unemployed or not achieving a high school education increase vulnerability and predispose young women to HIV (5, 10, 11). High risk sexual behaviour such as engaging in multiple concurrent partnerships, transactional sex and age-disparate relationships, has contributed significantly to enhancing HIV risk and is exacerbated by the inability of women to negotiate condom use even in long-term partnerships (9, 12–15). Physiological changes of the genital tract as well as factors which affect the integrity of the genital tract epithelium such as sexually transmitted infections (STIs) and intra-vaginal insertion practices, either for cleansing or enhancing sex, may increase susceptibility to HIV by facilitating viral entry (9, 16–19). Recent data suggests that genital tract inflammation in women with symptomatic and asymptomatic STIs may upregulate HIV susceptible target cells at the mucosal level thereby aiding transmission (20). The association between hormonal contraception use, in particular progestogen-only injectable preparations, and HIV risk remains contentious as current biological and epidemiological evidence is limited (21, 22).

As the HIV epidemic evolves, it is important to understand the factors that contribute to vulnerability and risk so that the design of HIV prevention programmes are tailored taking these into account. In this prospective cohort study we explored factors associated with HIV acquisition in a generalised epidemic setting.

Methods

Study Setting and Study Population

The Centre for the AIDS Programme of Research in South Africa (CAPRISA) initiated the CAPRISA 002 study to advance the understanding of HIV-1 subtype C acquisition, pathogenesis and disease progression. Between August 2004 and May 2005 volunteers from the city of Durban, KwaZulu-Natal, South Africa were recruited for study participation. Eligibility criteria and screening and enrolment procedures have been described (23). Briefly women 18 years and older, self-identifying as sex workers or having had at least three partners in the 3 months prior to recruitment, and testing HIV negative were eligible for study participation.

Study procedures

Volunteers were provided with verbal and written information about the study objectives and procedures after which written consent for screening procedures and for long-term storage of specimens were obtained. Women agreeing to participate in the study received pre and post-test counselling for HIV testing and risk reduction counselling and had blood samples collected for HIV antibody testing. Women testing HIV positive were referred to support services for ongoing care and psychosocial support. All women testing HIV antibody negative or indeterminate were enrolled until the study endpoint of HIV infection or for a period of 24 months.

At baseline, trained nurses administered questionnaires to all participants to obtain information regarding socio-demographic history, risk behaviour, knowledge about HIV and more direct questions related to sex work. A complete physical examination was undertaken at baseline and at each monthly follow up visit. Urine pregnancy testing was done if clinically indicated or upon request by a participant (23). Pelvic examination for genital sample collection was undertaken at baseline and at six monthly intervals. Blood specimen collection was done at baseline (safety monitoring tests, serological testing for STIs and long-term storage), at each monthly visit (HIV testing) and at each six-monthly visit (safety monitoring tests and serological testing for STIs).

Laboratory Evaluations

HIV testing was done using two rapid antibody tests using the Determine HIV-1 test (Abbott Diagnostics, Johannesburg, South Africa) and the Capillus HIV-1/HIV-2 test (Trinity Biotech, USA) followed by HIV-1 RNA polymerase chain reaction (PCR) testing using the Cobas AmpliScreen Multiprep HIV-1 test version 1.5 and the Cobas AmpliPrep/Cobas Amplicor HIV-1 Monitor test version 1.5 (Roche Diagnostics, Branchburg, New Jersey, USA) if at least one rapid antibody test was negative (24). Confirmatory HIV ELISA testing was done on PCR positive samples (Enzygnost anti HIV1/2 plus Dade Behring, Deerfield, Illinois, USA) (24). HIV infection as endpoint was based on a positive HIV-1 antibody test with a previously documented negative HIV-1 antibody test; or the presence of HIV-1 RNA in the absence of HIV antibodies (23).

Safety monitoring tests comprised haematological and biochemical evaluations. Genital specimens were tested for *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, *Trichomonas vaginalis*, *Mycoplasma genitalium* and herpes simplex virus (HSV) type 2 using PCR and bacterial vaginosis (BV) was diagnosed on Gram-stained smears using Nugent's criteria (20). Syphilis [Becton Dickinson Macro-Vue RPR (rapid plasma reagin) card test, BD Diagnostic systems United States and Omega IMMUTREP TPHA (Treponema pallidum Hemagglutination Assay) test, Omega Diagnostics Group PLC], HSV infection (HerpeSelect-1 and HerpeSelect-2, Enzyme-linked immunosorbent assay, Focus Diagnostics, Cypress, CA) and hepatitis B (HBV) infection (Centaur XP, ADVIA Centaur XP Immunoassay System, SIEMENS AG, Germany) were diagnosed serologically.

Data Management

Demographic, behavioural, clinical and laboratory data were captured onto standardised case report forms (CRFs) which were coded with a participant identification number in order to maintain participant confidentiality. CRFs were faxed using the DataFax system (Clinical DataFax Systems Inc., Ontario, Canada), with all data verified by data encoders for quality checks and stored in a secure study specific database.

Measures

HIV-1 infection was the main outcome measure. Baseline information on socio-demographic factors (age, number of dependents, educational level), behavioural factors (age at sexual debut, relationship status, frequency and type of sex acts, male and or female condom use [defined as condom use] at last sex act, contraception type, douching, alcohol and substance use prior to sex), sex for compensation (time period involved in sex work, age started sex work, days per week engaged in sex work, number of sites worked at, number of sex clients, short sessions or overnight stays per week with clients and condom use), clinical factors, including pregnancy and laboratory measures (full blood counts, liver function tests, electrolytes, vitamin B12, folate, iron, glucose, calcium, phosphate, CD4+, CD3+ and CD8+ cell counts and STIs), and knowledge questions on HIV prevention and transmission were obtained.

Statistical Analysis

Baseline data were summarised using descriptive statistics, with continuous variables reported as means and standard deviations or medians and interquartile ranges, while categorical variables are reported as percentages and actual numbers. Unadjusted and adjusted proportional hazards regression analysis was performed to assess the impact of socio-demographic factors, risk behaviour, sex for compensation and clinical factors on HIV acquisition. Estimated time of HIV infection was defined as the midpoint between the last negative HIV ELISA and the first positive HIV ELISA or 14 days prior to a positive HIV-1 RNA test if the HIV ELISA is negative on the same day (23). Time to HIV infection was calculated from the date of enrolment until the estimated time of HIV infection. Participants who remained HIV negative were censored at their last visit. Factors with a p-value of less than 0.2 in the unadjusted analysis were included in the adjusted model. Given the biological plausibility of anal sex and the risk for HIV acquisition, this variable was included in a multivariable model as was contraception type although current evidence is

limited (25-28). Although some variables relating to sex work had a p-value of less than 0.2 in the unadjusted analysis, these were not included in the adjusted model, because a large proportion of the cohort did not identify themselves as sex workers. The second multivariable model was restricted to women reporting sex for compensation.

P-values less than 0.05 were considered statistically significant. Analysis was performed using SAS software version 9.3 (SAS Institute Inc., Cary).

Ethics Approval

Ethics approval for the CAPRISA 002 study was obtained from the Universities of Natal (E013/04), Cape Town (025/2004) and Witwatersrand (MM040202). The ethical approval for the secondary analysis of data was granted from the Biomedical Research Ethics Committee, University of KwaZulu-Natal (BE 092/11) and permission to review data stored in the CAPRISA 002 database was approved through the CAPRISA data sharing policy.

Results

A total of 775 women were screened of whom 509 (65.7%) were ineligible for the following reasons: tested HIV positive (n=462; 59.6%), reported less than three sexual partners in the previous three months (n=22; 2.8%), were pregnant (n=16; 2.1%), planned to relocate (n=4; 0.5%), younger than 18 years (n=3; 0.4%) or afraid of testing procedures (n=2; 0.3%) (23). Twenty-one (2.7%) women were eligible for study participation but did not return for enrolment and 245 (31.6%) were enrolled into the study.

After 390 wy of follow up, 28 women acquired HIV, yielding an incidence rate (IR) of 7.20 per 100 wy [95% confidence interval (CI) 4.50–9.80]. Women aged 18–24 years had the highest HIV incidence (IR 13.20 per 100 wy, 95% CI 6.59–23.62) and among women 25 years and older it was 4.58 per 100 wy (95% CI 2.50–7.68). Three women were excluded from any risk factor analyses as they were in the window period of HIV infection at study entry.

The baseline characteristics overall and of those acquiring HIV are shown in Table I. The mean age of 242 women was 34.3 years [standard deviation (SD) 10.47, range 18–58], 185 (76.8%) women had one or more dependents and 156 women (64.5%) had an educational level which was above grade 8. The mean age at sexual debut was 17 years (SD 2.38, range 12–26). Although a large proportion (191; 78.9%) self-identified themselves as sex workers, having spent a median of three years [interquartile range (IQR) 1.3–8.3] in sex work, only 137 (56.6%) women reported being in a relationship with multiple partners at the time of the study.

Women who acquired HIV were approximately five years younger than those who remained HIV negative [t statistic (t)=2.20, p=0.029], reported being in relationships with multiple partners more frequently [Fisher's exact test hypergeometric probability (Fprob)=0.021, p=0.054] and had higher baseline serum vitamin B12 levels (t=-2.00, p=0.047).

Table II shows the socio-demographic, behavioural and biological variables associated with risk of HIV acquisition overall and by sex work. Younger women, aged 18–24 years were

almost three times more likely to acquire HIV compared to women 25 years and older [Hazard Ratio (HR) 2.85, 95% CI 1.29–6.28; Chi-square test statistic (χ^2)=6.73; $p=0.010$] and similarly women reporting many partners were at almost three times greater risk of HIV acquisition (HR 2.61, 95% CI 1.04–6.52; $\chi^2=4.18$; $p=0.041$). A higher educational level, above grade 8, was weakly associated with increased risk for HIV (HR 2.45, 95% CI 0.92–6.52; $\chi^2=3.21$; $p=0.073$), but no significant associations were found for HIV risk and engaging in anal sex (HR 1.49, 95% CI 0.68–3.29; $\chi^2=0.99$; $p=0.321$) or age at sexual debut (HR 0.88, 95% CI 0.73–1.06; $\chi^2=1.88$; $p=0.170$).

Among women reporting sex for compensation, HIV risk remained three-fold higher among 18–24 year-old women (HR 3.27, 95% CI 1.29–8.30; $\chi^2=6.24$; $p=0.013$) and ten-fold higher among those reporting many partners (HR 10.32, 95% CI 1.37–77.55; $\chi^2=5.15$; $p=0.023$). Weak associations were found between increased HIV risk and every additional overnight stay per week with clients (HR 1.33, 95% CI 0.94–1.88; $\chi^2=2.61$; $p=0.106$) and reduced risk for every additional year spent in sex work (HR 0.93, 95% CI 0.85–1.02; $\chi^2=2.34$; $p=0.126$).

In the final multivariable model, young age remained significant for HIV risk (HR 2.61, 95% CI 1.05–6.47; $\chi^2=4.27$; $p=0.039$).

Discussion

In this study young age and having multiple sex partners were associated with risk for HIV acquisition. Despite the introduction of HIV prevention and treatment programmes, the overall HIV incidence rate of 7.20 per 100 wy, and 13.20 per 100 wy among women aged 18–24 years remains unacceptably high (23). South Africa and many southern African countries continue to experience similar high HIV incidence rates. Longitudinal studies among women in South Africa have shown HIV incidence rates of 6.6, 8.5 and 6.5 per 100 wy particularly amongst young women in the province of KwaZulu-Natal (6, 10, 29). Among pregnant women from the same region, HIV incidence was 10.7 per 100 wy (30). In Malawi and Zimbabwe, among women recruited from postnatal or family planning clinics, the HIV incidence rates were 4.20, 4.86 and 4.78 per 100 wy in Lilongwe, Blantyre and Harare respectively (31). Overall, the highest incidence rate of 5.78 per 100 wy was among young women <25 years, whilst in Rwanda, among antenatal clinic attendees HIV incidence was 10.5 per 100 wy among women < 20 years (31, 32). These studies demonstrate young women's vulnerability and greater risk of acquiring HIV compared to older women. Our study shows that young age carries a three-fold greater risk of HIV acquisition.

The importance of measuring HIV incidence is key to understanding the dynamics of HIV disease to shape and modify effective responses. The persistence of high HIV incidence rates and the vulnerability of young women is incompletely elucidated in this setting. Recent studies have shown that residing in urban informal settlements, being unmarried or unemployed was associated with higher HIV incidence highlighting the underlying structural and social factors driving the epidemic (5, 10). Although common in this setting, high risk behaviours such as engaging in age-disparate relationships, was not found to predict HIV risk in a population-based study and is less likely to contribute to high HIV

incidence in young women (33). Sexually transmitted infections are known to contribute to HIV risk in women in this region and ongoing research into the immunology of the female genital tract may provide further insight into the biological susceptibility to HIV infection, particularly in young women (10, 20).

Although the vulnerability of young women is well recognised in this setting, risk for HIV is often underestimated since the perceived risk of HIV remains low (5). South Africa's epidemic is generalised with HIV prevalence at an unprecedented high level, in excess of 15% in the adult population, and new infection rates around 2% per year (5, 34). However, the majority of HIV-infected individuals are unaware of their HIV status, and this remains a barrier for both treatment access and prevention and helps sustain the epidemic. Poor knowledge of HIV transmission and limited access to and availability of health care services further promotes risk (5).

Our study demonstrates that women in relationships with multiple partners were more likely to acquire HIV. While multiple concurrent partnerships is recognised as a key driver of the epidemic in the Southern African region, the total number of lifetime partners has been found to be a significant predictor of HIV risk (12, 35). A meta-analysis of 68 studies from sub-Saharan Africa assessing risk factors for HIV acquisition reported that the number of lifetime partners increased the risk of HIV by almost four-fold [Odds Ratio (OR) 3.64, 95% CI 2.87–4.62] (35). In an urban mining community in Carletonville risk for HIV was almost five-fold higher in women reporting two or more lifetime partners (OR 4.88, 95% CI 3.01–7.89) (36). This risk was almost double for women reporting more than one partner three months prior to enrolment into HIV prevention intervention studies (adjusted OR 1.78, 95% CI 1.11–2.85) and similar for young women 15–24 years (OR 2.0, 95% CI 1.1–3.7) participating in the national household survey of South African youth, reporting more than one lifetime partner (4, 37).

In this setting unemployment levels remain high and resources are constrained (9, 38). Marriage is also uncommon in this setting, leading to instability of relationships (9). Transactional sex, which may range from serial monogamous relationships to occasional exchange of sex for money or goods, or to employment as a sex worker, provides a means of survival for women and their dependents, resulting in women engaging in multiple partnerships, either sequential or concurrent, in order to meet their basic needs (8, 9). Hence, emphasis and efforts on monogamy and multiple partner reduction cannot stand alone. The integration of structural approaches into current HIV policy, which address the socio-economic needs of women; and which create an environment of equal opportunities and rights, is crucial if behavioural and biomedical prevention efforts are to succeed. Nonetheless, rigorously including HIV counselling and testing (HCT) across all levels is important and key to knowledge of HIV status. HCT is especially important to help people learn their own and their partners' status so as to reduce HIV risk. Furthermore, repeat HCT must be promoted in hyperendemic regions.

The use of hormonal contraception was not found to be associated with HIV acquisition. Results from several studies exploring the use of hormonal contraception increasing the risk of HIV acquisition remain inconsistent (21, 22, 27, 39). Major limitations of these studies

have been the failure to test the hypotheses through robust study designs. Earlier studies showed some signals of an association between hormonal contraception and HIV acquisition (27). Similarly intra-vaginal insertion practices and pregnancy were not associated with HIV risk, though the outcome may be limited by the small sample size.

Whilst elevated serum vitamin B12 levels was associated with HIV acquisition, its causal role requires further exploration. Recent evidence from critically ill elderly medical patients suggests that elevated serum vitamin B12 levels are associated with increased mortality (40, 41). Although increased levels have been associated with systemic inflammatory markers, its role in mediating HIV risk remains unclear (40). While genital tract inflammation associated with HIV risk in this cohort has been reported, the role of vitamin B12 as a marker of systemic inflammation and HIV risk requires further evaluation (20).

The major strength of the study was the prospective cohort study design which allowed assessment of risk factors prior to HIV acquisition, however, there are several limitations as well. Firstly, participants had monthly study visits with risk reduction counselling, male and female condom provision, HIV testing with interviewer administered questionnaires; it is therefore possible that participants could have provided socially desirable responses given the sensitive nature of the sexual risk behaviour questions which could have biased the associations. Furthermore, given the wide age range of this cohort (18–58 years), it was difficult to adequately assess some risk behaviours which may be age dependent. Ideally, it would be important to follow up behaviour in those women who remained HIV negative to assess HIV risk over time. Secondly, the small sample size of the study is an important limitation, impacting on the precision of the study findings, therefore the particularly large confidence intervals around some of the point estimates. This may have also limited our ability to demonstrate stronger associations with some variables. Thirdly, the recruitment of high risk women limits the generalizability of the study findings as these may not be representative of women from elsewhere in the province. Finally, as the study excluded women younger than 18 years, it was not possible to measure HIV incidence and risk factors in this young age group in whom HIV incidence is high (7). Whilst there are ethical challenges in conducting research in adolescents, it is important that young people, especially girls are included into research studies in order to better understand the high rates of HIV acquisition (7, 42).

In conclusion, the findings of this study confirm that young women continue to bear the brunt of the HIV epidemic in this region. It is important that as transmission dynamics change over time, especially in hyperendemic settings, investments towards large-scale, fundamental changes in behaviour, social practices and community norms, address young women's vulnerability to HIV.

Acknowledgements

This work was supported by grants from the Comprehensive International Program of Research on AIDS (CIPRA) of the Division of AIDS (DAIDS), National Institute of Allergy and Infectious Disease (NIAID), National Institutes of Health (NIH), US Department of Health and Human Services (DHHS) [grant number 5U19 AI051794] and the National Research Foundation, South Africa [grant number UID 67385]. NN was partially sponsored by the University of KwaZulu-Natal for career development.

The authors would like to acknowledge the Centre for the AIDS Programme of Research in South Africa (CAPRISA) 002 Study Team and the participants of the CAPRISA 002 Study without whom this work would not have been possible.

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Table I

Baseline socio-demographic, behavioural and biological characteristics of women, overall and by HIV status, Durban, South Africa, 2004/2007

Variable	Total (n = 242)	HIV Positive (n = 25)	HIV Negative (n = 217)	Test statistic = value ^a	p-value
<i>Demographic data</i>					
Age in years (Mean, \pm SD)	34.3 (10.47)	30.0 (10.37)	34.8 (10.39)	t=2.20	0.029
Age group (in years) % (n)					
18–24	26.54% (64)	44.0% (11)	24.4% (53)	Fprob=0.023	0.053
25	73.6% (178)	56.0% (14)	75.6% (164)		
Number of dependents % (n)					
0	23.2% (56)	16.0% (4)	24.1% (52)	Fprob=0.143	0.460
1	76.8% (185)	84% (21)	75.9% (164)		
Educational level % (n)					
Grade 8	35.5% (86)	20.0% (5)	37.3% (81)	Fprob=0.042	0.121
> Grade 8	64.5% (156)	80.0% (20)	62.7% (136)		
<i>Risk behaviour</i>					
Age in years at sexual debut (Mean, \pm SD)	17.0 (2.38)	16.5 (1.76)	17.1 (2.44)	t=1.18	0.239
Relationship status % (n)					
No partner or Stable/Married partner	43.2% (104)	24.0% (6)	45.4% (98)	Fprob=0.021	0.054
Many partners	56.6% (137)	76.0% (19)	54.4% (118)		
Mean sex acts per month (\pm SD)	10.3 (6.78)	10.3 (5.03)	10.3 (6.97)	t=-0.01	0.994
Type of sex act % (n)					
Ever had vaginal sex	100% (242)	100% (25)	100% (217)	–	–
Ever had anal sex	34.6% (83)	44.0% (11)	33.5% (72)	Fprob=0.100	0.374
Ever had oral sex	25.3% (61)	24.0% (6)	25.5% (55)	Fprob=0.192	1.000
Condom use at last sex act % (n)					
Yes	58.7% (142)	64.0% (16)	58.1% (126)	Fprob=0.147	0.670
No	41.3% (100)	36.0% (9)	41.9% (91)		
Any contraception % (n)	79.7% (192)	68.0% (17)	81.0% (175)	Fprob=0.064	0.185
Contraception type % (n)					
Condom only	41.5% (100)	36.0% (9)	42.1% (91)	Fprob=0.030	0.808
Hormonal	30.7% (74)	32.0% (8)	30.5 (66)		
None/Other ^b	27.8% (67)	32.0% (8)	27.2% (59)		
Douching after sex % (n)	9.5% (23)	4.0% (1)	10.2% (22)	Fprob=0.210	0.483
Sex after alcohol or substance use % (n)					
Yes	26.9% (65)	20.0% (5)	27.7% (60)	Fprob=0.144	0.484
No	73.1% (177)	80.0% (20)	72.4% (157)		
<i>Sex for compensation</i>					
Self-reported sex workers % (n)	78.9% (191)	72.0% (18)	79.7% (173)	Fprob=0.129	0.436
Years in sex work (Median, IQR)	3.0 (1.3 – 8.3)	2.5 (1.0 – 7.0)	3.0 (1.3 – 9.0)	Z=-1.36	0.179

Variable	Total (n = 242)	HIV Positive (n = 25)	HIV Negative (n = 217)	Test statistic = value ^a	p-value
Age at start of sex work (Mean, ±SD)	28.2 (9.02)	27.2 (10.16)	28.3 (8.92)	t=0.49	0.625
Days per week perform sex work (Mean, ±SD)	2.9 (1.40)	2.9 (1.55)	2.9 (1.39)	t=-0.21	0.835
Sites worked per year (Mean, ±SD)	2.6 (1.30)	2.8 (0.86)	2.6 (1.33)	t=-1.08	0.291
Clients per day (Mean, ±SD)	2.5 (1.19)	2.6 (1.24)	2.5 (1.19)	t=-0.36	0.722
Clients in the past week (Mean, ±SD)	3.1 (3.83)	2.4 (2.53)	3.1 (3.94)	t=1.01	0.323
Short sessions per week (Mean, ±SD)	4.2 (4.05)	3.6 (4.68)	4.3 (3.99)	t=0.63	0.527
Overnight stays per week (Mean, ±SD)	1.1 (1.20)	1.4 (1.34)	1.0 (1.19)	t=-1.46	0.147
Condom use in the last month % (n)					
Always	38.2% (73)	27.8% (5)	39.3% (68)	Fprob=0.033	0.650
Sometimes	42.9% (82)	50.0% (9)	42.2% (73)		
Never	18.9% (36)	22.2% (4)	18.5% (32)		
HIV knowledge					
How safe is anal sex compared to peno-vaginal sex? % (n)					
Same or more risk	62.8% (152)	56.0% (14)	63.6% (138)	Fprob=0.002	0.278
Less risk or don't know	37.2% (90)	44.0% (11)	36.4% (79)		
How safe is oral sex compared to peno-vaginal sex? % (n)					
Same or more risk	71.8% (173)	68.0% (17)	72.2% (156)	Fprob=0.163	0.644
Less risk or don't know	28.2% (68)	32% (8)	27.8 (60)		
If you have an STI are you more likely to get HIV? % (n)					
Yes	71.1% (172)	80.0% (20)	70.1% (152)	Fprob=0.085	0.665
No	1.7% (4)	0.0% (0)	1.8% (4)		
Only sometimes/unsure	27.3% (66)	20.0% (5)	28.1% (61)		
Do you think HIV can be treated? % (n)					
Yes	95.0% (230)	96.0% (24)	94.9% (206)	Fprob=0.159	0.516
No	2.1% (5)	4.0% (1)	1.8% (4)		
Unsure	2.9% (7)	0.0% (0)	3.2% (7)		
Clinical evaluation					
Ever pregnant prior to HIV % (n)					
Yes	10.3% (25)	12.0% (3)	10.1% (22)	Fprob=0.244	0.730
No	89.7% (217)	88.0% (22)	89.9% (195)		
Laboratory parameters					
Haemoglobin g/dL (Mean, ±SD)	12.7 (1.29)	12.8 (1.43)	12.7 (1.27)	t=-0.51	0.613
Eosinophils $1 \times 10^9/L$ (Mean, ±SD)	0.2 (0.24)	0.3 (0.25)	0.2 (0.24)	t=-0.81	0.420
Albumin g/L (Mean, ±SD)	44.7 (3.57)	44.8 (3.32)	44.7 (3.60)	t=-0.02	0.981
Vitamin B12 pg/mL (Mean, ±SD)	291.0 (93.14)	327.6 (112.19)	287.0 (90.25)	t=-2.00	0.047
Folate ng/mL (Mean, ±SD)	22.4 (8.45)	20.8 (8.54)	22.6 (8.45)	t=0.94	0.346
Serum iron $\mu\text{mol/L}$ (Mean, ±SD)	12.3 (6.41)	12.1 (5.36)	12.3 (6.53)	t=0.16	0.877
Random glucose mmol/L (Mean, ±SD)	5.2 (2.02)	4.7 (0.85)	5.3 (2.11)	t=2.72	0.008
Sodium mmol/L (Mean, ±SD)	137.1 (2.22)	138.2 (2.08)	137.0 (2.21)	t=-2.66	0.008
Potassium mmol/L (Mean, ±SD)	4.0 (0.32)	4.1 (0.41)	4.0 (0.32)	t=-0.35	0.730

Variable	Total (n = 242)	HIV Positive (n = 25)	HIV Negative (n = 217)	Test statistic = value ^a	p-value
Chloride mmol/L (Mean, ±SD)	103.0 (2.55)	103.9 (1.98)	102.9 (2.59)	t=-1.93	0.054
Calcium mmol/L (Mean, ±SD)	2.4 (0.12)	2.4 (0.09)	2.4 (0.12)	t=-0.67	0.501
Phosphate mmol/L (Mean, ±SD)	1.1 (0.18)	1.1 (0.17)	1.0 (0.18)	t=-1.43	0.155
HIV negative CD4 ⁺ count cells/μl (Median, IQR)	888 (742 – 1132)	969 (855 – 1068)	878 (738 – 1132)	Z=0.76	0.446
HIV negative CD3 ⁺ count cells/μl (Median, IQR)	1493 (1202 – 1839)	1522 (1324–1857)	1486 (1197 – 1822)	Z=0.58	0.564
HIV negative CD8 ⁺ count cells/μl (Median, IQR)	517 (402 – 688)	500 (420–760)	520 (402–680)	Z=-0.04	0.966
Bacterial vaginosis % (n)	52.7% (127/241)	68.0% (17)	50.9% (110/216)	Fprob=0.047	0.139
Sexually transmitted infection^c % (n)	31.3% (75/240)	44.0% (11)	29.8% (64/217)	Fprob=0.063	0.172

^aTest statistic are: t=t-tests or Z=Wilcoxon Rank Sums (Normal Approximation), for continuous variables; Fprob=Fisher's exact test hypergeometric probability, for categorical data

^bFemale sterilisation or rhythm/calendar method

^cAny STI present if participant tested positive for syphilis antibodies or *Trichomonas vaginalis*, *Neisseria gonorrhoeae*, *Mycoplasma genitalium*, *Chlamydia trachomatis* or herpes simplex virus (HSV) type II (PCR)

Socio-demographic, behavioural and biological risk factors for HIV acquisition in women, overall and by sex work, Durban, South Africa, 2004/2007

Table II

Variable	Overall				By Sex Work							
	#HIV Cases/PY at risk	HIV Incidence (95% CI)	Univariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)	#HIV Cases/PY at risk	HIV Incidence (95% CI)	Univariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)
<i>Demographic characteristics</i>												
Age group (in years)												
25	14/305.90	4.58 (2.50–7.68)	1.00	–	1.00	–	10/248.35	4.03 (1.93–7.40)	1.00	–	1.00	–
18–24	11/83.32	13.20 (6.59–23.62)	2.85 (1.29–6.28)	0.010 ($\chi^2=6.73$)	2.61 (1.05–6.47)	0.039 ($\chi^2=4.27$)	8/59.36	13.48 (5.82–26.56)	3.27 (1.29–8.30)	0.013 ($\chi^2=6.24$)	2.44 (0.79–7.56)	0.123 ($\chi^2=2.38$)
Number of dependents												
None	4/79.27	5.05 (1.37–12.92)	1.00	–	–	–	3/62.58	4.79 (0.99–14.01)	1.00	–	–	–
1 or more	21/309.13	6.79 (4.21–10.38)	1.36 (0.47–3.95)	0.576 ($\chi^2=0.31$)	1.92 (0.65–5.68)	0.241 ($\chi^2=1.37$)	15/244.31	6.14 (3.44–10.13)	1.29 (0.37–4.46)	0.686 ($\chi^2=0.16$)	1.45 (0.40–5.20)	0.570 ($\chi^2=0.32$)
Educational level												
Grade 8	5/148.20	3.37 (1.10–7.87)	1.00	–	1.00	–	4/115.79	3.45 (0.94–8.85)	1.00	–	1.00	–
Grade >8	20/241.02	8.30 (5.07–12.82)	2.45 (0.92–6.52)	0.073	1.92 (0.65–5.68)	0.241 ($\chi^2=1.37$)	14/191.92	7.29 (3.99–12.24)	2.10 (0.69–6.39)	0.190 ($\chi^2=1.72$)	1.45 (0.40–5.20)	0.570 ($\chi^2=0.32$)
<i>Behavioural characteristics</i>												
Age at sexual debut (per 1 year increase)												
			0.88 (0.73–1.06)	0.170 ($\chi^2=1.88$)	0.86 (0.71–1.05)	0.149 ($\chi^2=2.08$)			0.96 (0.78–1.19)	0.707 ($\chi^2=0.14$)		
Relationship status												
No partner or stable/married partner	6/175.52	3.40 (0.09–18.92)	1.00	–	1.00	–	1/117.30	0.85 (0.02–4.75)	1.00	–	1.00	–
Many Partners	19/211.75	8.97 (5.40–14.01)	2.61 (1.04–6.52)	0.041 ($\chi^2=4.18$)	2.47 (0.98–6.26)	0.056 ($\chi^2=3.65$)	17/190.41	8.93 (5.20–14.29)	10.32 (1.37–77.55)	0.023 ($\chi^2=5.15$)	8.55 (1.11–6.02)	0.040 ($\chi^2=4.23$)
Average no. of sex acts per month (per 1 act increase)												
			1.00 (0.94–1.06)	0.954 ($\chi^2<0.01$)	–	–			0.99 (0.93–1.06)	0.801 ($\chi^2=0.06$)		
Ever had anal sex												
No	14/253.73	5.52 (3.02–9.26)	1.00	–	1.00	–	9/192.44	4.68 (2.14–8.88)	1.00	–	1.00	–
Yes	11/133.33	8.25 (4.12–14.76)	1.49 (0.68–3.29)	0.321 ($\chi^2=0.99$)	1.65 (0.73–3.74)	0.230 ($\chi^2=1.44$)	9/115.11	7.82 (3.57–14.84)	1.66 (0.66–4.19)	0.281 ($\chi^2=1.16$)	1.68 (0.63–4.47)	0.301 ($\chi^2=1.07$)
Ever had oral sex												
No	19/293.53	6.47 (3.90–10.11)	1.00	–	–	–	13/225.89	5.76 (3.06–9.84)	1.00	–	–	–
Yes	6/94.75	6.33 (2.32–13.78)	0.97 (0.39–2.44)	0.955 ($\chi^2<0.01$)	–	–	5/80.88	6.18 (2.01–14.43)	1.07 (0.38–2.99)	0.902 ($\chi^2=0.02$)	–	–
Condom use at last sex act												
Yes	16/225.00	7.11 (4.06–11.55)	1.00	–	–	–	11/173.32	7.11 (6.35–11.36)	1.00	–	–	–
No	9/164.22	5.48 (2.51–10.40)	0.77 (0.34–1.75)	0.534 ($\chi^2=0.39$)	–	–	7/134.39	5.21 (2.09–10.73)	0.82 (0.32–2.12)	0.684 ($\chi^2=0.17$)	–	–

Variable	Overall				By Sex Work				p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)
	#HIV Cases/PY at risk	HIV Incidence (95% CI)	Univariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)	#HIV Cases/PY at risk	HIV Incidence (95% CI)					
Contraception type													
Condom only	9/161.76	5.56 (2.54–10.56)	1.00	–	1.00	–	7/127.41	5.49 (2.21–11.32)	1.00	–	1.00	–	–
Hormonal	8/118.78	6.75 (2.91–13.31)	1.21 (0.47–3.14)	0.692 ($\chi^2=0.16$)	1.03 (0.38–2.76)	0.958 ($\chi^2<0.01$)	5/91.16	5.48 (1.78–12.80)	1.00 (0.32–3.15)	0.999 ($\chi^2<0.01$)	0.86 (0.27–2.75)	0.799 ($\chi^2=0.07$)	–
None or other ^a	8/108.16	7.40 (3.19–14.57)	1.33 (0.51–3.45)	0.558 ($\chi^2=0.34$)	1.51 (0.57–4.03)	0.411 ($\chi^2=0.68$)	6/88.32	6.79 (2.49–14.79)	1.24 (0.42–3.69)	0.699 ($\chi^2=0.15$)	1.52 (0.49–4.77)	0.469 ($\chi^2=0.52$)	–
Douching after sex													
No	24/356.29	6.73 (4.32–10.02)	1.00	–	–	–	18/280.89	6.41 (3.80–10.13)	1.00	–	–	–	–
Yes	1/32.11	3.11 (0.08–17.35)	0.46 (0.06–3.40)	0.448 ($\chi^2=0.58$)	–	–	0/25.99	0.00	–	–	–	–	–
Sex after alcohol or substance use													
No	20/285.24	7.01 (4.28–10.83)	1.00	–	–	–	13/215.56	6.03 (3.21–10.31)	1.00	–	–	–	–
Yes	5/103.98	4.81 (1.56–11.22)	0.68 (0.26–1.82)	0.447 ($\chi^2=0.58$)	–	–	5/92.15	5.43 (1.76–12.66)	0.90 (0.32–2.51)	0.834 ($\chi^2=0.04$)	–	–	–
Sex for compensation													
Sex worker													
No	7/81.51	8.59 (3.45–17.69)	1.00	–	–	–	–	–	–	–	–	–	–
Yes	18/307.71	5.85 (3.47–9.25)	0.68 (0.28–1.63)	0.390 ($\chi^2=0.74$)	–	–	–	–	–	–	–	–	–
Time in sex work (per 1 year increase)													
Age at start of sex work (per 1 year increase)			0.93 (0.85–1.02)	0.126 ($\chi^2=2.34$)	0.93 (0.85–1.02)	–	–	–	0.93 (0.85–1.02)	0.126 ($\chi^2=2.34$)	0.94 (0.85–1.05)	0.289 ($\chi^2=1.12$)	–
Days per week perform sex work (Per 1 day increase)			0.98 (0.93–1.04)	0.564 ($\chi^2=0.33$)	0.98 (0.93–1.04)	–	–	–	0.98 (0.93–1.04)	0.564 ($\chi^2=0.33$)	–	–	–
Number of sites per year (per 1 site increase)			1.08 (0.79–1.48)	0.648 ($\chi^2=0.21$)	1.08 (0.79–1.48)	–	–	–	1.08 (0.79–1.48)	0.648 ($\chi^2=0.21$)	–	–	–
Clients per day (per 1 client increase)			1.12 (0.80–1.57)	0.503 ($\chi^2=0.45$)	1.12 (0.80–1.57)	–	–	–	1.12 (0.80–1.57)	0.503 ($\chi^2=0.45$)	–	–	–
Clients past week (per 1 client increase)			1.10 (0.76–1.59)	0.628 ($\chi^2=0.23$)	1.10 (0.76–1.59)	–	–	–	1.10 (0.76–1.59)	0.628 ($\chi^2=0.23$)	–	–	–
Short sessions per week (per 1 session increase)			0.93 (0.75–1.16)	0.536 ($\chi^2=0.38$)	0.93 (0.75–1.16)	–	–	–	0.93 (0.75–1.16)	0.536 ($\chi^2=0.38$)	–	–	–
Overnight stays per week (per 1 stay increase)			0.97 (0.83–1.12)	0.651 ($\chi^2=0.20$)	0.97 (0.83–1.12)	–	–	–	0.97 (0.83–1.12)	0.651 ($\chi^2=0.20$)	–	–	–
Condom use with client in the last month			1.33 (0.94–1.88)	0.106 ($\chi^2=2.61$)	1.33 (0.94–1.88)	–	–	–	1.33 (0.94–1.88)	0.106 ($\chi^2=2.61$)	1.22 (0.84–1.77)	0.291 ($\chi^2=1.11$)	–
Clinical evaluation													
Always	5/119.27	4.19 (1.36–9.78)	1.00	–	–	–	5/119.27	4.19 (1.36–9.78)	1.00	–	–	–	–
Sometimes	9/132.17	6.81 (3.11–12.93)	1.61 (0.54–4.82)	0.391 ($\chi^2=0.74$)	–	–	9/132.17	6.81 (3.11–12.93)	1.61 (0.54–4.82)	0.391 ($\chi^2=0.74$)	–	–	–
Never	4/56.27	7.11 (1.94–18.20)	1.68 (0.45–6.26)	0.440 ($\chi^2=0.60$)	–	–	4/56.27	7.11 (1.94–18.20)	1.68 (0.45–6.26)	0.440 ($\chi^2=0.60$)	–	–	–

Variable	Overall				By Sex Work				p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)	Univariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)
	#HIV Cases/PY at risk	HIV Incidence (95% CI)	Univariate Hazard Ratio (95% CI)	p-value (χ^2)	Multivariate Hazard Ratio (95% CI)	p-value (χ^2)	#HIV Cases/PY at risk	HIV Incidence (95% CI)						
Ever pregnant prior to HIV														
No	22/342.24	6.43 (4.03–9.73)	1.00	–	–	–	22/342.24	5.89 (3.37–9.57)	1.00	–	–	–	–	–
Yes	3/46.99	6.38 (1.32–18.66)	0.99 (0.30–3.32)	0.990 ($\chi^2 < 0.01$)	–	–	2/36.15	5.53 (0.67–19.98)	0.93 (0.22–4.06)	0.927 ($\chi^2 < 0.01$)	–	–	–	–
<i>Laboratory evaluations</i>														
Sexually transmitted infection^b or bacterial vaginosis														
No	5/148.75	3.36 (1.09 – 7.84)	1.00	–	–	–	4/116.34	3.44 (0.94 – 8.80)	1.00	–	–	–	–	–
Yes	20/238.49	8.39 (5.12 – 12.95)	2.47 (0.93–6.58)	0.071 ($\chi^2 = 3.26$)	2.49 (0.91–6.85)	0.077 ($\chi^2 = 3.13$)	14/189.38	7.39 (4.04–12.40)	2.13 (0.70–6.46)	0.184 ($\chi^2 = 1.77$)	1.89 (0.59–6.08)	0.287 ($\chi^2 = 1.14$)	1.89 (0.59–6.08)	0.287 ($\chi^2 = 1.14$)

^aFemale sterilisation or rhythm/calendar' method

^bAny STI present if participant tested positive for syphilis antibodies or *Trichomonas vaginalis*, *Neisseria gonorrhoeae*, *Mycoplasma genitalium*, *Chlamydia trachomatis* other herpes simplex virus (HSV) type II (PCR)