HIV/AIDS

Stabilizing HIV prevalence masks high HIV incidence rates amongst rural and urban women in KwaZulu-Natal, South Africa

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Background In mature generalized human immunodeficiency virus (HIV) epidemics, as survival from accessing antiretroviral treatment (ART) increases, HIV prevalence data may be suboptimal and difficult to interpret without HIV incidence rates.

Objective To determine the HIV incidence rate among rural and urban women in KwaZulu-Natal, South Africa.

Methods We conducted a prospective cohort study from March 2004 to May 2007. Volunteers were recruited from a rural family-planning clinic and an urban clinic for sexually transmitted infections. Consenting, HIV-uninfected women aged 14–30 years were enrolled. Demographic, clinical, sexual and behavioural data were collected using standardized questionnaires with HIV risk reduction counselling and HIV testing. Pelvic examinations were completed at quarterly visits.

Results The HIV prevalence at screening was 35.7% [95% confidence interval (CI) 32.7–38.8] amongst rural women and 59.3% (95% CI 56.5–62.0) amongst urban women. A total of 594/2240 (26.5%) enrolled women contributed to 602 person-years (PYs) of follow-up. The median age was 22 years [inter-quartile range 18–23 years]. HIV incidence rate was 6.5/100 PY (95% CI 4.4–9.2) amongst rural women and 6.4/100 PY (95% CI 2.6–13.2) amongst urban women. HIV incidence rate of 17.2/100 PY (95% CI 2.1–62.2) was highest amongst urban women <20 years of age and 10.2/100 PY (95% CI 4.1–20.9) amongst rural women ≥25 years of age.

Conclusion HIV incidence rates are devastatingly high in young women in rural and urban KwaZulu-Natal, despite reports of stabilized HIV prevalence observed in current surveillance data. The diffuse nature of the HIV epidemic underscores the urgent need to enhance HIV prevention and treatment modalities.

Keywords HIV incidence, HIV prevalence, young women, South Africa
Introduction

South Africa has the highest burden of human immunodeficiency virus (HIV) infection in the world with an estimated 5.7 million people infected with HIV in 2008.1 The province of KwaZulu-Natal is at the epicentre of the pandemic, and women account for >60% of infections.2-4 The epidemic typology has been described as ‘hyperendemic, generalized and mature’5 to reflect >15.0% HIV prevalence in the general population.2 Annual, anonymous, antenatal surveys undertaken since 1990 have served as the mainstay of HIV surveillance for monitoring trends in the evolving epidemic, for priority setting and health-service planning. Data from these surveys in the past 3 years demonstrate a stabilization of national HIV prevalence estimates.6

With increasing access to antiretroviral treatment (ART),7 HIV prevalence is expected to increase resulting from increased survival.8 Notwithstanding the challenges to increasing ART access and coverage,7,9 knowledge of the epidemic is key to customizing responses.5,10,11 In the early stages of the epidemic where disease burden and mortality rates are low, HIV prevalence data provide a reliable marker for monitoring the epidemic.12 However, as the epidemic matures morbidity and mortality increases;5,13,14 incidence rate data8,13-17 are important to understand ongoing transmission dynamics and inform the prioritization of prevention efforts, whereas prevalence of established infections informs health service and care planning needs.5,10

As laboratory methods to differentiate established and new infections are being validated, cohort studies provide an option for ascertaining rates of new HIV infections. This prospective cohort study was undertaken in a rural and urban setting to better understand the stabilized HIV prevalence data in KwaZulu-Natal and to assess the feasibility of undertaking HIV prevention studies with incident HIV infections as endpoints.

Methods

Study design, setting and source population

The Centre for the AIDS Programme of Research in South Africa (CAPRISA) is an AIDS research institute in KwaZulu-Natal, South Africa conducting research that contributes to understanding HIV epidemiology, prevention and pathogenesis, as well as the links between tuberculosis and AIDS care. This prospective cohort study was undertaken at CAPRISA’s two clinical research sites.

The CAPRISA Vulindlela Clinical Research site (rural site) is located ~150 km north-west of Durban. This community has approximately 400 000 residents, with limited resources, infrastructure and employment opportunities accounting for high levels of poverty and unemployment.18 Health services are provided by seven public-sector primary health-care clinics and the closest referral hospitals are ~30 km away.

From March 2004 to April 2005, volunteers were recruited from the rural family planning clinic. The CAPRISA eThekwini Clinical Research site (urban site) is located in central Durban adjacent to a large public-sector primary-care clinic designated for the diagnosis and treatment of sexually transmitted infections (STIs) and tuberculosis. From July 2005 to December 2005, female volunteers were recruited from the STI clinic. Based on cross-sectional survey data, HIV incidence rates are highest in young women and establishing cohorts of young women aged 14–30 years, although this age cut-off reduces generalizability, targets women at highest HIV risk.

Study procedures

Volunteers aged 14–30 years received study information, provided written informed consent and were screened for eligibility to assess willingness to provide adequate locator data for study retention purposes, be sexually active (defined as having had vaginal intercourse at least once in the 3 months prior to screening) and test HIV negative. Women were excluded if planning to travel away from the study site for >3 consecutive months in the 12 months following enrolment; plans to migrate outside the study catchment area in the 12 months following enrolment; plans to become pregnant in the next 2.5 years post-enrolment; or enrolled in any other study of a vaginally applied product related to HIV prevention.

At enrolment and at each monthly visit, demographic, clinical, sexual and behavioural data were obtained from all women through nurse-administered interviews using structured questionnaires. Women were tested for HIV and pregnancy, received HIV and risk reduction counselling and were offered male and female condoms. At enrolment, quarterly and at study exit visits, pelvic examinations were completed and blood samples collected for archiving for prospective confirmation of HIV endpoint. Women were reimbursed minimally at each visit for time and travel costs as per the South African National Ethics Guidelines.19

Laboratory testing procedures

HIV testing was performed on-site using two rapid antibody tests (Determine HIV-1/2—Abbott Laboratories, IL, USA and HIV-1/2 SmartCheck assay—Globalemed.LLC, World Diagnostics, Inc.). Women testing concordantly positive, indeterminate or discordant at follow-up visits had blood sample collected at the same visit for HIV antibody testing (Enzygnost Anti-HIV-1/2 Plus, Dade Behring, Marburg, Germany) and HIV-1 RNA polymerase chain reaction (PCR) assay (COBAS AmpliScreen HIV-1 Test, version 1.5 (Qualitative) and AMPLICOR HIV-1 MONITOR Test, v1.5 (Quantitative); Roche Molecular Systems,
Inc., Branchburg, NJ, USA). Incident HIV infection was confirmed on positive HIV-1 RNA PCR (qualitative and quantitative) and/or HIV antibody positive, and negative on the last stored blood sample tested retrospectively. Urine pregnancy testing was performed on site (QuickVue One-Step hCG Urine Test; Quidel Corporation, San Diego, USA).

Ethical considerations
The protocol, informed consent and data collection forms were reviewed and approved by the University of KwaZulu-Natal Biomedical Research Ethics committee. The screening and enrolment visits were undertaken on different days to give eligible women sufficient time to consider their decision to participate in this study. At both sites, community engagement processes were established through Community Research Support Groups (CRSG) consisting of stakeholders including health-service providers and traditional leaders from community structures. The community liaison officers worked closely with CRSG members on a regular basis to disseminate information on the epidemiology of HIV epidemic in South Africa, HIV prevention strategies and the rationale for conducting HIV prevention research and HIV treatment.

At screening, women identified with HIV infection were referred to the CAPRISA AIDS Treatment Program operational at both sites. Pregnant women identified during screening or follow-up were referred to local public-sector facilities for follow-up antenatal care but continued with their monthly study visits. Women diagnosed with symptomatic STIs were managed syndromically on site as per the South African Department of Health guidelines. Contraceptive counselling and contraceptives were provided at both sites as part of the study visit. Women who became HIV infected were given the option of joining the CAPRISA Acute HIV-1 Infection Study or the CAPRISA AIDS Treatment Project for ongoing care and support.

Data collection and statistical analysis
Data were collected on standardized case report forms (CRFs) and faxed to a dedicated study database using DataFax (Clinical DataFax Systems Inc., Hamilton, Canada). For quality assurance, all CRF entries were manually and electronically verified and the quality of the data reviewed in real time. Data analysis was undertaken with Statistical Analysis Software (SAS) statistical package (version 9.1.3; SAS, NC, USA).

The baseline variables were summarized using descriptive statistics expressed as mean or median for continuous variables and as percentages for categorical variables. For statistical testing for differences, the Fisher’s exact test comparing categorical data and Wilcoxon rank sum test for continuous data were used. Mc Nemar’s agreement statistic was used for analysis of paired categorical data with two categories, Bowker’s test of symmetry for paired categorical data with more than two levels and Wilcoxon signed rank sum test for paired continuous data. Missing data were not included in the comparison between baseline and 12 months. All analyses were two-sided.

HIV incidence rate was estimated on the number of confirmed seroconversions, calculated per 100 person-years (PYs) at risk. The time at risk was defined as the time from enrolment to the last negative HIV test for women who remained HIV uninfected or the first confirmed positive HIV test result for those who became infected during the study. Pregnancy rate was estimated on the number of positive urine pregnancy tests, calculated per 100 PYs. Time at risk excluded the time from a positive pregnancy test to pregnancy outcome. Confidence intervals (CIs) for HIV and pregnancy incidence rate assumed a Poisson distribution.

Results
Screening, accrual and retention
A total of 2240 (981 rural and 1259 urban) volunteers were screened. HIV prevalence was 35.7% (95% CI 32.7–38.8) amongst rural women and 59.3% (95% CI 56.5–62.0) amongst urban women; the main reason for exclusion of volunteers. At both sites, HIV prevalence was highest amongst volunteers 25 years and older compared with younger volunteers (both sites P < 0.001) (Table 1). Other reasons for ineligible included volunteers being underage (13.8%), not returning for enrolment visit (5.8%), not sexually active (3.0%), requesting voluntary counselling and testing services only (1.2%), pregnancy (0.2%), planning to relocate from current address (0.1%), fear of study procedures especially pelvic examinations (0.2%), medical reasons (0.2%) and partner not agreeing to her participation in research (0.01%).

A total of 594 (26.6%) women were enrolled with a mean enrolment rate per month per site of 26 (rural) and 17 (urban) women with a screening to enrolment ratio of 1:2 and 1:10, respectively. The overall retention rate was 70.0% (95% CI 65.7–73.9) and 78.0% (95% CI 70.2–85.3) at the rural and urban sites, respectively. There were no differences in women who completed the study compared with those who did not return for follow-up with regard to demographic characteristics of age, completion of secondary schooling, relationship status, living with partner or whether they were from the rural or urban site (all P > 0.05). At exit, there were 477 and 117 women at the rural and urban site, respectively, with a cumulative of 602 PYs.

Demographic characteristics
All enrolled women were Black African with an overall median age of 22 years [inter-quartile range (IQR) 18–23 years]. Rural women compared with urban
women were younger; a higher proportion were <20 years of age and had not completed secondary schooling (all \( P < 0.001 \)). Majority of women reported to be in a stable relationship though they were currently not living with their sexual partner (Table 2).

### Sexual behaviour characteristics

Sexual behaviour data at baseline and at 12 months follow-up are presented in Table 3. Amongst rural women, there was a reduction in the mean number of sex acts reported in the 2 weeks preceding their Month 12 study visit (\( P = 0.03 \)), increase in reported condom use with vaginal sex (\( P < 0.001 \)), decline in the proportion reporting one sex partner in the past 3 months (\( P = 0.004 \)) with no changes in contraceptive-use patterns (\( P = 0.05 \)), although a higher proportion of women reported engaging in anal sex (0.8 vs 1.5%; \( P = 0.18 \)). In contrast, among urban women there was a decline in anal-sex practice (6.8 vs 1.3%, \( P = 0.03 \)), no decline in the mean number of sex acts in the past 2 weeks preceding their Month 12 study visit (\( P = 0.17 \)), or the percentage of women reporting one partner in the past 3 months (\( P = 0.37 \)), or changes in contraceptive-use patterns (\( P = 0.48 \)) and condom use during last vaginal sex act (\( P = 0.10 \)).

The practice of intra-vaginal substance use of water with salt, vinegar, soap or douching with antiseptics containing parahydroxybenzoyl benzoate or povidone iodine products or traditional herbs was used for personal hygiene. Despite counselling on potential HIV risk, we did not observe any changes in these practices in rural women (17.6 vs 12.0%, \( P = 0.07 \)). The self-report of STIs (genital ulceration and/or vaginal discharge) declined at 12 months amongst rural (25.4 vs 4.5%, \( P < 0.0001 \)) and urban women (28.2 vs 5.2%, \( P < 0.0001 \)); however, any abnormal pelvic examination finding (vaginal discharge, vaginitis, cervicitis, ulceration, abrasions, petechia, vesicles or genital warts) declined amongst rural women only (39.6 vs 15.4%, \( P < 0.0001 \)) and not amongst urban women (36.8 vs 30.9%, \( P = 0.35 \)).

### Incidence of HIV

As shown in Table 4, a total of 39 (6.6%) women became HIV infected during the study. The HIV incidence rate was 6.5/100 PY (95% CI 4.4–9.2) amongst rural women and 6.4/100 PY (95% CI 2.6–13.2) amongst urban women. The highest rate of 17.2/100 PY (95% CI 2.1–62.2) was among urban women <20 years of age, whereas the rate of 10.2/100 PY (95% CI 4.1–20.9) was highest amongst older rural women, \( \geq 25 \) years of age. HIV incidence rate was higher amongst older rural women compared with younger women [10.2/100 PY (95% CI 4.1–20.9) vs 3.6/100 PY (95% CI 1.7–6.9), \( P = 0.03 \)].

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**Table 1** HIV prevalence and screening outcome of rural (March 2004 to April 2005) and urban (July to December 2005) women aged 14–30 years in KwaZulu-Natal, South Africa

<table>
<thead>
<tr>
<th>HIV prevalence</th>
<th>Overall</th>
<th>Rural</th>
<th>Urban</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number screened</td>
<td>2240</td>
<td>981</td>
<td>1259</td>
</tr>
<tr>
<td>HIV prevalence</td>
<td>1096/2240</td>
<td>350/981</td>
<td>746/1259</td>
</tr>
<tr>
<td>% (95% CI)</td>
<td>48.9 (46.8–51.0)</td>
<td>35.7 (32.7–38.8)</td>
<td>59.3 (56.5–62.0)</td>
</tr>
</tbody>
</table>

**Age-specific HIV prevalence**

<table>
<thead>
<tr>
<th>&lt;20 years</th>
<th>101/378</th>
<th>47/245</th>
<th>54/133</th>
</tr>
</thead>
<tbody>
<tr>
<td>% (95% CI)</td>
<td>26.7 (22.4–31.5)</td>
<td>19.2 (14.6–24.8)</td>
<td>40.6 (32.3–49.5)</td>
</tr>
<tr>
<td>20–24 years</td>
<td>283/694</td>
<td>81/258</td>
<td>202/436</td>
</tr>
<tr>
<td>% (95% CI)</td>
<td>40.8 (37.1–44.6)</td>
<td>31.4 (25.9–37.5)</td>
<td>46.3 (41.6–51.1)</td>
</tr>
<tr>
<td>( \geq 25 ) years</td>
<td>575/870</td>
<td>85/180</td>
<td>490/690</td>
</tr>
<tr>
<td>% (95% CI)</td>
<td>66.1 (62.8–69.2)</td>
<td>47.2 (39.8–54.8)</td>
<td>71.0 (67.5–74.4)</td>
</tr>
</tbody>
</table>

**Screening outcome**

| HIV positive | 1096 | 350 | 746 |
| % (95% CI) | 48.9 | 35.7 | 59.3 |
| Age >30 years | 308 | 21 | 287 |
| % (95% CI) | 13.8 | 2.1 | 22.8 |
| Pregnant | 5 | 0 | 5 |
| % (95% CI) | 0.2 | 0.0 | 0.4 |
| Not sexually active | 68 | 65 | 3 |
| % (95% CI) | 3.0 | 6.6 | 0.2 |
| Voluntary counselling and testing | 27 | 0 | 27 |
| % (95% CI) | 1.2 | 0.0 | 2.1 |
| Planning to move away | 3 | 0 | 3 |
| % (95% CI) | 0.1 | 0.0 | 0.2 |
| Afraid of clinical procedures | 4 | 4 | 0 |
| % (95% CI) | 0.2 | 0.4 | 0.0 |
| Medical reasons | 4 | 0 | 4 |
| % (95% CI) | 0.2 | 0.0 | 0.3 |
| Partner refusing | 1 | 0 | 1 |
| % (95% CI) | 0.0 | 0.0 | 0.1 |
| Not returning for enrolment | 130 | 64 | 66 |
| % (95% CI) | 5.8 | 6.5 | 5.2 |

*Missing age for 298 women.*
Incidence of Pregnancy

A total of 96 women (16.2%) became pregnant during the study (Table 4), out of which 74 were rural and 22 urban. The pregnancy incidence rate was 16.6/100 PY (95% CI 13.1–20.9) among rural women and 22.4/100 PY (95% CI 14.1–34.0) among urban women. The highest pregnancy incidence rate of 30.6/100 PY (95% CI 6.3–89.3) was amongst young urban women <20 years of age.

Of the women who became pregnant, 50 (52.1%) reported not having used any form of contraception, 32 (33.3%) reported relying on partners using male condoms only, 9 (17.3%) and 5 (5.2%) reported to be on injectable or oral forms of contraception, respectively. Not surprisingly, women on hormonal contraception who became pregnant reported being non-adherent to contraceptive use.

Of the 96 pregnancies, 62 (64.6%) continued to full term, 20 (20.8%) were still pregnant at exit, 5 (5.2%) resulted in spontaneous abortion or had elective termination of pregnancy and in 9 (9.4%) the outcome was unknown. Should women in a trial of an investigational new drug become pregnant, for safety reasons they would be required to go off product for the duration of the pregnancy and the pregnancy time would translate to 58.7 PY.

Discussion

In this sample of rural and urban women in KwaZulu-Natal, HIV incidence rates of 6.5/100 PY and 6.4/100 PY, respectively, are devastatingly high.
Table 4: HIV incidence and pregnancy rates by age group among rural (March 2004 to May 2007) and urban (July 2005 to May 2007) women aged 14–30 years in KwaZulu-Natal, South Africa

<table>
<thead>
<tr>
<th>Enrolment and follow-up</th>
<th>Overall</th>
<th>Rural</th>
<th>Urban</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number enrolled</td>
<td>594</td>
<td>477</td>
<td>117</td>
</tr>
<tr>
<td>Person-years at risk (PYs)</td>
<td>602.0</td>
<td>492.8</td>
<td>109.2</td>
</tr>
<tr>
<td>Retention rate</td>
<td>71.4% (95% CI 67.5–75.0)</td>
<td>69.8% (95% CI 65.4–73.9)</td>
<td>77.8% (95% CI 69.0–84.7)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>HIV incidence rate</th>
<th>n</th>
<th>Cases PY</th>
<th>Per 100 PY (95% CI)</th>
<th>n</th>
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<td>32</td>
<td>492.8</td>
<td>117</td>
<td>7</td>
<td>109.2</td>
</tr>
<tr>
<td>&lt;20 years</td>
<td>244</td>
<td>11</td>
<td>260.7</td>
<td>227</td>
<td>9</td>
<td>249.1</td>
<td>17</td>
<td>2</td>
<td>116.2</td>
</tr>
<tr>
<td>20–24 years</td>
<td>241</td>
<td>20</td>
<td>249.5</td>
<td>166</td>
<td>16</td>
<td>174.8</td>
<td>75</td>
<td>4</td>
<td>74.7</td>
</tr>
<tr>
<td>≥25 years</td>
<td>109</td>
<td>8</td>
<td>91.8</td>
<td>84</td>
<td>7</td>
<td>68.9</td>
<td>25</td>
<td>1</td>
<td>22.9</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Pregnancy rate</th>
<th>n</th>
<th>Cases PY</th>
<th>Per 100 PY (95% CI)</th>
<th>n</th>
<th>Cases PY</th>
<th>Per 100 PY (95% CI)</th>
<th>n</th>
<th>Cases PY</th>
<th>Per 100 PY (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall</td>
<td>594</td>
<td>96</td>
<td>543.4</td>
<td>477</td>
<td>74</td>
<td>445.3</td>
<td>117</td>
<td>22</td>
<td>98.1</td>
</tr>
<tr>
<td>&lt;20 years</td>
<td>244</td>
<td>51</td>
<td>227.2</td>
<td>227</td>
<td>48</td>
<td>217.4</td>
<td>17</td>
<td>3</td>
<td>9.8</td>
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<tr>
<td>20–24 years</td>
<td>241</td>
<td>36</td>
<td>229.0</td>
<td>166</td>
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<td>17</td>
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<td>7</td>
<td>65.4</td>
<td>25</td>
<td>2</td>
<td>21.8</td>
</tr>
</tbody>
</table>

*Excludes time pregnant.
2.4% amongst men and women aged 15–48 years. Young women in the age group 20–29 years had the highest rate of 5.6% compared with 0.9% amongst men in the same age group.11 Several studies21,22,27 and this study confirm the persistently high steady-state HIV incidence rates, demonstrating the underlying transmission dynamics despite the scaled-up prevention and treatment efforts, which have failed to address the HIV prevention needs of young women. This study highlights the importance of augmenting HIV prevalence data with incidence rate data to get a more accurate indication of trends of HIV infection.16 Although these studies are expensive and may not be feasible to undertake at the same frequency as cross-sectional surveys, it is important in mature HIV epidemics to supplement HIV prevalence data with HIV incidence rate data to minimize com- placency emanating from a stabilized prevalence and to provide a more informative and sensitive marker of the state of the epidemic.27

The sexual-behavioural data highlight the com- plexity of HIV risk and prevention for women across their life course. Although marriage is rare, many women are sexually active and the duration of their relationships vary substantially. In these settings where economic options for women are limited and their survival is dependent on having a man in their life, child-bearing is an important facet of their identity. In this context, the traditional ‘ABC’ (abstinence, be faithful, condom use) prevention approach is misleading and has limited applicability.11,28 The higher abnormal pelvic examination findings in urban women is not surprising given that they were recruited from the STD clinic and are likely to be at higher risk of acquiring and reporting STIs. The strengthening of sexual reproductive health services, including comprehensive pregnancy prevention counselling, provision of services for reducing HIV transmission to infants and reducing maternal mortality rates has been gaining increasing attention;29–34 and in the context of HIV programme planning,33,34 is as important for primary and secondary HIV prevention.

Although this study has some important findings, the benefit of longitudinal data and rigorous endpoint ascertainment, it also has several limitations. A key limitation is the retention rate in the study as HIV risk behaviours of those lost to follow-up are likely to be different from those completing the study. As these are research-naïve cohorts, the potential challenges in undertaking longitudinal studies underscore the importance of placing more effort on retention for future studies.35,36 HIV risk behaviours were assessed using self-report and even though the staff received substantial training to minimize socially desirable responses, there is no guarantee for valid self-reported data.37 The anal sex reporting rates, for example, could be indicative of women’s response to information received from study staff in the HIV risk-reduction counselling session on the additional increased risk of HIV acquisition through anal sex compared with vaginal sex and could partially account for the lower reported rates of anal sex in these study populations compared with other reported studies from this region.21,22 Given its potential confounding effect in testing efficacy of vaginal products, accurate measurement of anal sex practice is important.35

Pregnancies were determined using chemical testing and could be an overestimate of real pregnancy rates.38 The potential bias with selecting women not intending to fall pregnant could impact HIV incidence rates substantially; however, only 0.2% of the pregnant volunteers were excluded at screening and none excluded as a result of planning to become pregnant, and this should have minimal impact on HIV incidence in this study. It is also clear that self-reported intention not to fall pregnant does not correlate with actual intentions and the importance of the selection criteria used. Although external validity is important, the strategy used is not generalizable even though it is important for identifying women at high risk of acquiring HIV infection. While the pregnancy rates were high but lower than when contraceptives are not provided by study sites,39–41 women in the rural site recruited from the family-planning clinic were more likely to already be on a reliable form of contraceptive in contrast to urban women recruited from the STD clinic. A key lesson for future clinical trials of investigational new drugs is the importance of counselling and on-site provision of contraceptive services to support fertility control and maintain the scientific integrity of the trial through minimizing product hold. Lastly, the urban HIV incidence rate is less stable than the rural estimate given the smaller size of the cohort and the shorter duration of follow-up.

Notwithstanding these study limitations and the HIV prevalence appearing to be stabilizing,7,6,27 these results highlight the uniquely diffuse, generalized, hyperendemic nature of the epidemic in South Africa and the unprecedented high HIV prevalence and incidence rates across rural and urban KwaZulu-Natal. Although these populations are not representative of all HIV epidemics, the high prevalence and incidence rate data underscore the importance and appropriateness of including these cohorts for testing new HIV prevention modalities. The public health imperative for the conduct of pivotal prevention trials in this select population in this region is particularly important to alter HIV epidemic trajectories in this region and globally.

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Conflicts of interest: None declared.

KEY MESSAGES
- The HIV incidence rates are disturbingly high among young women in rural and urban KwaZulu-Natal, South Africa, despite reports of stabilized HIV prevalence observed in current surveillance data.
- The diffuse nature of the epidemic highlights the need to enhance HIV prevention and treatment modalities for young women.

References
19 Department of Health. *Guidelines for Good Practice in the Conduct of Clinical Trials with Human Participants in*


