EPIDEMIOLOGICAL REVIEW

Monitoring the effectiveness of HIV and STI prevention initiatives in England, Wales, and Northern Ireland: where are we now?

A E Brown, S E Tomkins, L E Logan, D S LaMontagne, H L Munro, V D Hope, A Righarts, J E Blackham, B D Rice, T R Chadborn, P A Tookey, J V Parry, V Delpech, O N Gill, K A Fenton

Primary and secondary prevention are essential components of the response to HIV and sexually transmitted infections (STIs). We present findings from nationally implemented HIV/STI prevention interventions. In 2003, of those attending STI clinics at least 64% of men who have sex with men (MSM) and 55% of heterosexuals accepted a confidential HIV test; 88% of all HIV infections in women giving birth in England were diagnosed before delivery; 85% of MSM eligible for hepatitis B vaccination received a first dose of vaccine at their first STI clinic attendance; 74% of STI clinic attendees for emergency appointments, and 20% of those for routine appointments were seen within 48 hours of initiating an appointment; the National Chlamydia Screening Programme in England found a positivity of 10% and 13% among young asymptomatic women and men, respectively. Prevention initiatives have seen recent successes in limiting further HIV/STI transmission. However, more work is required if current levels of transmission are to be reduced.

Prevention is an essential component of the response to HIV and sexually transmitted infection (STI) transmission. Despite the availability of effective antiretroviral therapy (ART), HIV infection is only treatable provided that an assiduous routine of medication is followed indefinitely (often with adverse side effects). HIV care and treatment are expensive and ART resistance is thought to be increasing in England. STI treatment costs are also substantial and if left untreated, can have serious long term sequelae and possibly facilitate the transmission of others, including HIV.

For HIV/STIs, primary prevention targets uninfected individuals, for instance, by reducing risk factors for disease acquisition. Examples include diagnosing HIV in pregnant women (to prevent vertical transmission), hepatitis B vaccination, and harm reduction measures (for example, needle exchanges). Secondary prevention targets infected individuals, aiming to reduce onward disease transmission or re-infection. Examples include the promotion of sexual health screening—for example, chlamydia screening among young people and promoting voluntary confidential HIV testing (VCT) in STI clinics. An HIV diagnosis provides access to ART, a timely STI diagnosis usually leads to treatment, and both allow an opportunity for partner notification and behaviour change counselling.

Major challenges remain in ensuring that prevention initiatives are effective. They need to be accessible (particularly for higher risk populations who may be socially vulnerable); timely; comprehensive (address all modes of HIV/STI transmission); implemented through functioning health systems; and subject to monitoring and evaluation.

The Health Protection Agency and its collaborators monitor the effectiveness of some primary and secondary prevention efforts in addition to providing national HIV/STI surveillance data. We present findings from nationally coordinated prevention monitoring programmes. This paper does not present an overview of all HIV/STI prevention activities that occur in the United Kingdom, but summarises information on prevention monitoring and disease outcomes, to demonstrate recent progress, and highlight areas that need further work.

DATA SOURCES

In the United Kingdom, the majority of HIV/STI prevention initiatives are implemented through STI clinics, primary care, and other community based services. The Health Protection Agency and its collaborators use nationally coordinated information systems to monitor prevention initiatives. Prevention monitoring systems (summarised in table 1) and their objectives are generally separate from the variety of infection surveillance systems used, but can overlap.

HIV testing

Monitoring the uptake of VCT and antenatal HIV screening relies on data from the Unlinked Anonymous Prevalence Monitoring Programme (UAPMP) surveys. The UAPMP survey of STI clinic attendees measures HIV prevalence (including undiagnosed cases) in England.

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Abbreviations: ARV, antiretroviral therapy; IDUs, injecting drug users; MSM, men who have sex with men; NCSNP, National Chlamydia Screening Programme; NSHPC, National Survey of HIV in Pregnancy and Childhood; SOPHID, Survey of Prevalent HIV Infections Diagnosed; STI, sexually transmitted infections; UAPMP, Unlinked Anonymous Prevalence Monitoring Programme; VCT, voluntary confidential HIV testing
<table>
<thead>
<tr>
<th>Prevention aim</th>
<th>Year aim introduced</th>
<th>Indicator</th>
<th>Data source</th>
<th>Area covered</th>
<th>1998</th>
<th>1999</th>
<th>2000</th>
<th>2001</th>
<th>2002</th>
<th>2003</th>
<th>2004</th>
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</thead>
<tbody>
<tr>
<td>All STI clinics to offer VCT to all attendees on their first screening for STIs, and subsequently according to risk*</td>
<td>2002</td>
<td>Percentage MSM attending STI clinics accepting VCT</td>
<td>UAPMP E, W &amp; NI</td>
<td>47%</td>
<td>48%</td>
<td>50%</td>
<td>57%</td>
<td>62%</td>
<td>64%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2002</td>
<td>Percentage heterosexuals attending STI clinics accepting VCT</td>
<td>UAPMP E, W &amp; NI</td>
<td>26%</td>
<td>27%</td>
<td>31%</td>
<td>39%</td>
<td>49%</td>
<td>55%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>To reduce the proportion of individuals newly diagnosed with HIV who have a “late diagnosis”</td>
<td>NA</td>
<td>Percentage of MSM newly HIV diagnosed with CD4 count below 200 cells x 10^9/l</td>
<td>CD4 surveillance England &amp; Wales</td>
<td>30%</td>
<td>29%</td>
<td>27%</td>
<td>24%</td>
<td>25%</td>
<td>21%</td>
<td>19%</td>
<td></td>
</tr>
<tr>
<td></td>
<td>NA</td>
<td>Percentage of heterosexuals newly HIV diagnosed with CD4 count below 200 cells x 10^9/l</td>
<td>CD4 surveillance England &amp; Wales</td>
<td>45%</td>
<td>46%</td>
<td>44%</td>
<td>42%</td>
<td>44%</td>
<td>40%</td>
<td>NA</td>
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<tr>
<td>The universal offer and recommendation of an HIV test as a routine part of antenatal care</td>
<td>1999</td>
<td>Percentage HIV infected women diagnosed before delivery</td>
<td>UAPMP and NSHPC England</td>
<td>40%</td>
<td>59%</td>
<td>71%</td>
<td>84%</td>
<td>82%</td>
<td>88%</td>
<td>NA</td>
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</tr>
<tr>
<td></td>
<td>1999</td>
<td>Percentage exposed infants becoming HIV infected</td>
<td>UAPMP and NSHPC England</td>
<td>17%</td>
<td>12%</td>
<td>9%</td>
<td>6%</td>
<td>7%</td>
<td>5%</td>
<td>NA</td>
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<tr>
<td>To monitor the proportion of HIV infected individuals on triple ARV or more</td>
<td>NA</td>
<td>Percentage of diagnosed HIV infected MSM on triple ARV therapy or more</td>
<td>SOPHID E, W &amp; NI</td>
<td>56%</td>
<td>66%</td>
<td>66%</td>
<td>64%</td>
<td>67%</td>
<td>65%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td></td>
<td>NA</td>
<td>Percentage of diagnosed HIV infected heterosexuals on triple ARV therapy or more</td>
<td>SOPHID E, W &amp; NI</td>
<td>50%</td>
<td>63%</td>
<td>65%</td>
<td>63%</td>
<td>64%</td>
<td>63%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>To offer all MSM hepatitis B vaccination at their first STI clinic attendance*</td>
<td>2003</td>
<td>Percentage MSM taking up first dose at their first STI clinic attendance</td>
<td>HepB3 survey England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>85%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2003</td>
<td>Percentage MSM taking up third dose at their first STI clinic attendance</td>
<td>HepB3 survey England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>39%</td>
<td>59%</td>
<td></td>
</tr>
<tr>
<td>To reduce waiting times at STI clinics</td>
<td>2004</td>
<td>Percentage patients with emergency appointments seen within 48 hours</td>
<td>STI waiting times survey England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>74%</td>
<td>1359</td>
<td>1843</td>
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<tr>
<td></td>
<td>2004</td>
<td>Percentage patients with routine appointments seen within 48 hours</td>
<td>STI waiting times survey England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>20%</td>
<td>3044</td>
<td>5520</td>
</tr>
<tr>
<td></td>
<td>2004</td>
<td>Percentage patients with walk-in appointments seen within 48 hours</td>
<td>STI waiting times survey England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>79%</td>
<td>4960</td>
<td>6307</td>
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<tr>
<td>To control genital chlamydial infection through early detection and treatment of asymptomatic infection among individuals aged under 25</td>
<td>2004</td>
<td>Chlamydia prevalence among women aged under 25</td>
<td>NCSP England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>10.1%</td>
<td>1538</td>
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<td>2004</td>
<td>Chlamydia prevalence among men aged under 25</td>
<td>NCSP England</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>13.3%</td>
<td>156/1172</td>
<td></td>
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<tr>
<td>To reduce the proportion of injecting drug users who share injecting equipment</td>
<td>NA</td>
<td>Percentage current injectors who report sharing needles/syringes</td>
<td>UAPMP E, W &amp; NI</td>
<td>32%</td>
<td>33%</td>
<td>31%</td>
<td>33%</td>
<td>34%</td>
<td>29%</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td></td>
<td>NA</td>
<td>Percentage current injectors who report sharing other injecting equipment</td>
<td>UAPMP E, W &amp; NI</td>
<td>54%</td>
<td>54%</td>
<td>52%</td>
<td>51%</td>
<td>52%</td>
<td>30%</td>
<td>NA</td>
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</tbody>
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*Health Protection Scotland; The Institute of Child Health (London); Collaborators on the Unlinked Anonymous Programme.
HIV infection) in attendees of 16/232 STI clinics in England, Wales, and Northern Ireland undergoing syphilis tests. Residual blood from syphilis testing is irreversibly unlinked from patient identifiers and anonymously HIV tested. Retained information includes sexual orientation and sexual health screen uptake (including VCT, further details are available on the STI website).

The UAPMP surveys each clinic as discreet categorical units. The target population is sexually active individuals aged 15 years and over. The surveys include individuals receiving HIV related care in England, Wales, and Northern Ireland. Subsidiary information is collected from patient identifiers and anonymously HIV tested. Retained information includes sexual orientation and sexual health screen uptake (including VCT, further details are available on the STI website).

RESULTS: PREVENTION MONITORING UPDATE
VCT uptake

Overall, VCT uptake rose by 17% (95% CI 15% to 19%) from 47% (2956/6294) in 1998 to at least 55% (44 312/80 435) in 2003 for MSM and by 28% (95% CI 27% to 28%) from 27% (95% CI 26% to 28%) in 1998 to at least 64% (4920/7697) in 2003 for heterosexuals (fig 1A and B). Of those who did not accept VCT, at least 29% (817/2777) of MSM and 31% (11 312/36 142) of heterosexuals (fig 1C). The proportion of HIV infected women who were previously diagnosed. Patients for whom conflicting information was provided on HIV testing were excluded. Data source: Unlinked anonymous programme.

Descriptive epidemiology is the focus of the paper, but 95% confidence limits (95% CI) have been used to supplement main findings from the NCSP and the sentinel Unlinked Anonymous STI and IDU survey. All other prevention monitoring systems are comprehensive.

The UAPMP survey of injecting drug users (IDUs) collects self reported behavioural data (for example, injecting equipment sharing) in addition to measuring the prevalence of blood borne viruses among injecting drug users attending 63 specialist services in England, Wales, and Northern Ireland. Sharing rates are calculated for those who reported injecting in the previous 4 weeks.
England and Wales, 1994–2003. (Count under 200 cells

<table>
<thead>
<tr>
<th>Year</th>
<th>CD4 Surveillance Scheme</th>
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<tr>
<td>1994</td>
<td>30</td>
</tr>
<tr>
<td>1995</td>
<td>40</td>
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<tr>
<td>2002</td>
<td>110</td>
</tr>
<tr>
<td>2003</td>
<td>120</td>
</tr>
</tbody>
</table>

Figure 2 Late diagnosis* of HIV infection by exposure category, England and Wales, 1994–2003. (Percentage of patients with CD4 count under 200 cells ×10⁶/l within 90 days of diagnosis.) Data source: CD4 surveillance scheme.

36123) of heterosexuals are known to have been offered, but declined, VCT (fig 1C). Of those that declined VCT, 7% (56/817) of heterosexuals and 1% (83/11312) of heterosexuals were HIV infected.

The proportion of HIV infected individuals who could have been diagnosed during their attendance, but who left the clinic remaining unaware of their HIV infection, fell by 9% (95% CI 1% to 17%) from 60% (165/276) in 1998 to 51% (161/317) in 2003 among MSM and may have fallen by 7% (95% CI –1% to 15%) from 48% (104/217) in 1998 to 41% (160/394) in 2003 among heterosexuals.

Late diagnoses

In 2003, 33% (995/2982) of people with newly diagnosed HIV infection in England and Wales had CD4 counts below 200 cells ×10⁶/l. MSM are increasingly being tested at earlier stages of their infection* (fig 2). HIV infected heterosexuals were more likely to be diagnosed late.

Antenatal HIV testing

In England, 88% (697/790) of HIV infected women who gave birth in 2003 are calculated to have had their infection diagnosed before delivery. The majority of births to HIV infected women occurred in London where 89% (455/511) of women were diagnosed before delivery (fig 3). The proportion of children exposed to maternal HIV infection who acquire HIV is decreasing. Based on current estimated detection rates, 5% of children exposed to vertical transmission would have been infected in England in 2003, compared to 17% in 1998.

Uptake of antiretroviral therapy

Of MSM receiving HIV care, 65% (9991) were receiving at least three antiretroviral drugs in 2003 compared to 56% (5231) in 1998. Among heterosexuals, equivalent figures were 63% (9956) and 50% (2158). Thirty three per cent of MSM (5065) and 35% (5466) of heterosexuals were not receiving HIV therapy in 2003.

Uptake of hepatitis B vaccine among MSM

MSM were considered to be eligible for hepatitis B vaccination (dose 1) if they were not known to be either immune or fully/partially vaccinated. Overall, 85% (5598/6533) of eligible MSM were vaccinated with dose 1.

MSM eligible for the third vaccine dose (dose 3) included those who had had fewer than three doses, but excluded those who had had a booster dose, and those known to have immunity through blood testing following previous doses. The coverage rate for dose three was 39% (2588/6624) overall but showed regional variation (fig 4). Nearly one half (46%, 2588/5669) of MSM eligible for dose 1 completed the three dose course.

STI clinic waiting times

Nationally, 74% (1359/1843) of emergency appointments, 79% (4960/6307) of people attending walk-in clinics, and 20% (3044/15520) of people with routine appointments were seen within 48 hours. Lower proportions of 16–24 year olds and women of any age were seen within 48 hours. Full results have been published elsewhere.*1

Figure 3 Estimated proportion of HIV infected pregnant women diagnosed before delivery*, and of exposed children becoming HIV infected†, England, 1998–2003. (Includes previously diagnosed and those diagnosed through antenatal testing. †Assumes a vertical transmission rate of 26.5% in undiagnosed women and 2.2% in diagnosed women. *These data contain reports received by the end of September 2004 and are subject to reporting delay, particularly for 2003.) Data source: Unlinked Anonymous Programme and the National Study of HIV in Pregnancy and Childhood.

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among men and 32% among women. Chlamydia at STI clinics rose by 8%, and syphilis by 28%.

In 2003 there were 6606 new HIV diagnoses, more than double the number of HIV/STI diagnoses are increasing annually. In England, 16 413 young people were screened for chlamydia outside STI clinic settings during April 2003–March 2004. A 10% (1538/15 241, 95% CI 9.6% to 10.6%) and 13% (156/1172, 95% CI 11.4% to 15.4%) positivity among women and men aged under 25 was found respectively (fig 5). Women aged 16–19 were more likely to test positive for chlamydia than those aged 20–24; men aged 20–24 were twice as likely to be infected as younger men.

### Risk behaviours among IDUs

Thirty per cent (494/1677, 95% CI 28% to 32%) of injectors who had first injected in the previous 3 years reported ever having accessed a needle exchange service.

### DISCUSSION

Monitoring individual HIV/STI prevention initiatives in 2003 demonstrated successes. VCT uptake among STI clinic attendees was at the highest level ever recorded. Almost 90% of HIV infected women had their infection diagnosed before delivery in England leading to an increase in the proportion able to take advantage of interventions to prevent vertical transmission (for example, ARV and avoiding breast feeding), thereby decreasing the proportion of infants who become infected. The proportion of diagnosed HIV infected individuals on at least triple therapy has increased from around half in 1998 to almost two thirds in 2003. HIV infected MSM are being diagnosed at an earlier stage of their infection. Targeted screening for chlamydia has identified many asymptomatic infections that otherwise may have been missed.

However, high proportions of STI clinic attendees were not seen within 48 hours of seeking an appointment, delaying access to treatment and potentially heightening the risk of onward transmission. The elevated levels of injecting equipment sharing among IDUs is concerning.

### Are prevention initiatives reducing transmission?

Despite the individual success of many prevention initiatives, the number of HIV/STI diagnoses are increasing annually. In 2003 there were 6606 new HIV diagnoses, more than double the 2835 diagnoses in 1998. From 2002 to 2003, diagnoses of chlamydia at STI clinics rose by 8%, and syphilis by 28% among men and 32% among women.

The number of diagnoses is determined by interactions between transmission dynamics, migration of individuals from high prevalence areas, and the relative effectiveness of targeted testing. Increases in diagnoses represent both successes in increased testing in high risk populations as well as continued disease transmission within that population. The relative contribution of these factors is difficult to disentangle.

However, HIV incidence may be increasing among MSM attending STI clinics, whilst HIV prevalence among recently initiated injectors (those who began injecting in the previous 3 years) was at the highest level recorded in 2003, perhaps a result of elevated levels of equipment sharing.

The reasons for continuing transmission despite apparently successful prevention initiatives require consideration:

### Do prevention monitoring systems show the whole picture?

In the United Kingdom, methods for monitoring prevention initiatives are pragmatic, simple, inexpensive and reach accessible populations—not necessarily those at greatest risk. Consequently, data are generally derived from those using health services, and will exclude high risk populations who have poor access to services.

Monitoring systems also collect limited data. While rates of STIs are higher among black and ethnic minority populations, few UK prevention monitoring systems collect ethnicity data, making it difficult to monitor the specific impact of prevention initiatives on such vulnerable populations.

Each prevention monitoring system has its own limitations.

### HIV testing

All patients included in the Unlinked Anonymous STI survey are undergoing syphilis tests, therefore, VCT in this population may not represent all STI clinic attendees. The survey cannot monitor the frequency of repeat HIV testing (which may positively bias the results), or outcomes among first time attendees.

Diagnosis detection rates among HIV infected pregnant women are calculated by aligning diagnosis reports to the NSHPC with UAPMP prevalence data. Since data are anonymised, records are not individually matched. Limited mismatching may occur with respect to time and geography. Detection rates are minimum estimates and may rise as late reports are received by the NSHPC.

CD4 surveillance suggests MSM with HIV are being diagnosed earlier in their infection. However, the high proportion of heterosexuals categorised as having a “late diagnoses” may not accurately reflect recent efforts in VCT promotion; a high proportion of heterosexuals are infected abroad and may not have been resident in the United Kingdom long enough to have had an earlier diagnosis.

### ARV

HIV infected individuals accessing health care show an increase in ARV uptake. However, it is difficult to calculate what proportion of HIV infected individuals should be on therapy. Guidelines state that individuals who have a CD4 count below 200 cells $\times 10^6/l$ should begin therapy. Such guidelines are not directed from a public health perspective to prevent HIV transmission, but on the basis of individual need/readiness; patients may delay, interrupt, or stop therapy for many reasons.

The proportion of diagnosed HIV infected individuals with low CD4 counts on ARV is not routinely calculated because these specific fields are incomplete for a minority of records. In the future, this proportion may be calculated through cross linking to other HIV reporting databases.

### Hepatitis B vaccination

The HepB3 study demonstrates that high proportions of MSM are vaccinated with dose 1 on their first STI clinic attendance, but lower proportions complete the three dose course. Patient identifiers are not collected, so it is impossible...
to monitor movement of patients between clinics. This, combined with reporting delay, may lead to an underestimate of the true performance.

**STI clinic waiting times**

Although the national waiting times survey show a high proportion of attendees cannot get a timely appointment, it is not possible to calculate the median waiting time since data is collected in categories.

**Chlamydia screening programme**

While the NCSP has improved access for chlamydia screening since its implementation in England, it is not currently possible to calculate national coverage. When the programme is fully rolled out throughout England, coverage will be calculated by dividing the number of people aged 16–24 screened by the total eligible population (sexually active population aged 16–24).

**Risk behaviour monitoring**

The Unlinked Anonymous IDU survey only includes those in contact with services for drug users and therefore the data may not be generalisable to all injectors, specifically, those not in contact with services, who may have different levels of risk behaviour.

**Are prevention initiatives effective?**

The effectiveness of prevention initiatives to reduce transmission requires assessment. For instance, although the promotion of VCT has reduced the proportion of HIV-infected individuals leaving the clinic remaining undiagnosed, it may not target those who have been recently infected, who may be more infectious. Research is required to elucidate the role of “recently HIV infected” people in contributing onward HIV transmission. For IDUs, there is evidence of a recent shift in needle exchange provision towards pharmacy-based services. Studies suggest that IDUs using pharmacy-based services may be more likely to share equipment owing to absence of harm reduction counselling in these settings.

**Are prevention initiatives implemented on the correct scale?**

Since the prevalence of an infection drives onward transmission, recent increases in the prevalent pool of diagnosed and undiagnosed infections may limit the ability of prevention programmes to reduce transmission levels.

For example, the success in reducing the proportion of infants exposed to maternal HIV infection who become infected, has not substantially reduced the absolute number, because prevalence among pregnant women has increased. Continuing investment in prevention activities is essential, and activities need to adapt to meet the challenge of the evolving epidemics.

**What next?**

While individual prevention initiatives have had an impact, and rates of ongoing transmission would have been higher in their absence, investment needs to be strengthened and sustained in order to reduce HIV/STI transmission. Current prevention initiatives partially accommodate the diversity of populations at high risk of infection, but require flexibility if they are to match the evolving epidemic. Evaluation of prevention initiatives, both individually and in combination, is needed to measure how much they reduce transmission. Novel monitoring tools require development to assess prevention initiatives aimed at populations who have poor access to health services, but who play an important part in HIV/STI transmission.

Local and national surveillance systems are essential in ensuring that the effectiveness of prevention initiatives are continually reviewed and updated to meet the diversity of needs of the populations at risk of HIV/STIs.

**ACKNOWLEDGEMENTS**

We thank David Goldberg, Daniel Thomas, Brian Smyth, Sarah Dougan, Elizabeth Rudd, Christine McGarrigle, and all others who contributed to the writing and editing of the annual report Focus on Prevention.

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Confidential reports of HIV-infected pregnant women are collated at the Institute of Child Health by the National Study of HIV in Pregnancy and Childhood through surveillance schemes run in collaboration with the Royal College of Obstetricians and Gynaecologists and the British Paediatric Surveillance Unit of the Royal College of Paediatrics and Child Health. Research at the Institute of Child Health benefits from R&D funding received from the NHS Executive.

Finally, we thank Dr Helen Ward and Susie Huntington for their useful comments on drafts of this paper.

**CONTRIBUTORS**

AB, LL, SL, HM, VH, AR, BR, and TC analysed the data from the Unlinked Anonymous STI clinic survey, Unlinked Anonymous pregnant women survey, National Chlamydia Screening Programme, HepB3 survey, Unlinked Anonymous IDU survey, STI waiting times survey, SOPHID and CD4 surveillance scheme respectively with support from VD, NG, and KF; PT coordinates the programme to reduce transmission levels.

We gratefully acknowledge the continuing collaboration of the National Chlamydia Screening Programme, HepB3 survey, Unlinked Anonymous IDU survey, STI waiting times survey, SOPHID and CD4 surveillance scheme respectively with support from VD, NG, and KF; PT coordinates the National Study of HIV in Pregnancy and Childhood and collaborated with the analysis of the Unlinked Anonymous Pregnant Women Surveys; JD oversees the data for SOPHID, CD4 surveillance, and the NCSP; NG is the programme manager and is responsible for data from the Unlinked Anonymous Programme; JP is responsible for laboratory aspects for the Unlinked Anonymous Surveys and assisted with the interpretation of data; all authors were involved in interpretation of the results and drafting the paper with ST and JB substantially contributing; AB undertook the main writing of the paper.

**Key messages**

- Prevention activities have been seen recent successes in limiting further HIV/STI transmission in England, Wales, and Northern Ireland.
- High proportions of STI clinic attendees are having voluntary confidential HIV tests. HIV-infected pregnant women are diagnosed before delivery; MSM are receiving hepatitis B vaccinations at their first clinic attendance.
- Prevention monitoring activities may be missing at-risk populations who are likely to have an important role in onward HIV/STI transmission.
- Existing prevention initiatives may only be having a limited impact on the current rate of HIV/STI transmission and require development to match evolving epidemics.

**WHAT NEXT?**

- Prevention activities have been seen recent successes in limiting further HIV/STI transmission in England, Wales, and Northern Ireland.
- High proportions of STI clinic attendees are having voluntary confidential HIV tests. HIV-infected pregnant women are diagnosed before delivery; MSM are receiving hepatitis B vaccinations at their first clinic attendance.
- Prevention monitoring activities may be missing at-risk populations who are likely to have an important role in onward HIV/STI transmission.
- Existing prevention initiatives may only be having a limited impact on the current rate of HIV/STI transmission and require development to match evolving epidemics.
REFERENCES


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