BMJ Open Integration of sexually transmitted infection (STI) services into HIV care and treatment services for women living with HIV: a systematic review

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ABSTRACT

Objective To review and critically appraise the existing evidence on integration of sexually transmitted infection (STI) services into HIV care and treatment services for women living with HIV.

Design Systematic review.

Data sources Four electronic databases were searched through 16February 2017 using keywords for HIV, STIs and integration. Reference lists of included articles and other reviews were also screened.

Review methods We included studies that compared women living with HIV who received STI services integrated into HIV care and treatment services with those who received HIV care and treatment services without integrated STI services or standard of care.

Results Of 170 articles identified, 3 studies reported in 4 articles were included. Two studies evaluated comprehensive care for people living with HIV in the UK; in both cases, quality and uptake of STI services seemed to improve following integration. The third study conducted a comparative case study across different models of care in Swaziland: two clinics integrated with sexual and reproductive health services (including STI services), and two stand-alone HIV clinics (without STI services). Coverage for Pap smears among women living with HIV was higher at the fully integrated site, but there was no significant difference in the prevalence of sexual health screening or advice on sexual health. Reported client satisfaction was generally higher at the standalone HIV clinic, and a diverse range of factors related to implementation of different care models challenged the notion that integrated services are always superior or desired.

Conclusion While there is a limited evidence base for integrating STI services into HIV care and treatment services, existing studies indicate that integration is feasible and has the potential for positive outcomes. However, diverse population needs and health system factors must be considered when designing models of care to provide STI services to women living with HIV.

BACKGROUND

The United Nations 2030 Agenda for Sustainable Development defines a set of ambitious global health goals and targets, including the need to ensure healthy lives and promote

Strengths and limitations of this study

- Examining the evidence for integrating two related health services—sexually transmitted infection (STI) services and HIV services-for women living with HIV who have specific needs.
- Systematic search, screening and data abstraction process to evaluate the global literature on integration of STI services into HIV care and treatment services.
- While few studies were identified that met the inclusion criteria, the existing data indicate that integration can be feasible and can have positive outcomes, though differences were seen across settings and populations.

well-being for all at all ages (Goal 3).¹ To ensure the health sector implementation of this Agenda, WHO has developed global health sector strategies on HIV, viral hepatitis and sexually transmitted infections (STIs) which were adopted by the United Nations Member States in May 2016.²⁻⁴ Among the common priorities across the three strategies is the imperative to strengthen health systems to support integrated service delivery by identifying the best methods and approaches for delivering a continuum of high-impact, tailored health interventions and services to different populations and in different locations in order to achieve equity and ensure quality.

It is estimated that annually there are 357 million new cases of four curable STIs among people aged 15-49 years: Chlamydia trachomatis (131 million), Neisseria gonorrhoeae (78 million), syphilis (6 million) and Trichomonas vaginalis (142 million).⁴ That translates to nearly 1 million new infections per day. The prevalence of some viral STIs is similarly high, with an estimated 417 million people infected with herpes simplex type 2 and approximately 291 million women harbouring the human papillomavirus.⁴ STIs have clear negative health consequences, including fetal and neonatal death and disability, cervical cancer and infertility, as well as physical, social and psychological effects.⁴⁵ There is substantial evidence that STIs also increase HIV transmissibility as well as HIV acquisition risk by as much as 2–3 times in some populations.⁴⁶

For women living with HIV, a holistic and comprehensive approach to sexual and reproductive health and rights includes enhancing services to identify and treat STIs. Women living with HIV have high rates of STI coinfection; a recent systematic review of 37 studies found a mean STI prevalence among these women of 15.8% (SD 9.9).⁷ Screening and treatment for STIs have been recommended as a part of the package of care for people living with HIV by WHO^{8 9}; they are also part of the U.S. President's Emergency Plan for AIDS Relief (PEPFAR)'s minimum package of prevention services for people living with HIV.¹⁰ While many national HIV treatment guidelines (including those in the UK, the USA and South Africa) recommend that STI screening occurs within routine HIV care visits, published service statistics indicates that coverage rates for the recommended sexual history and STI screening are often very low.¹¹⁻¹⁴ For example, among a nationally representative sample of women in HIV treatment in the USA, the coverage of testing from 2008 to 2010 among sexually active women was only 49% for syphilis, 30% for chlamydia and 28% for gonorrhoea.15

A key approach to increasing coverage of STI services for people living with HIV is integration of these services within HIV care and treatment settings. As part of an effort to inform updated WHO guidelines on sexual and reproductive health and rights for women living with HIV, we sought to systematically review the evidence for the integration of STI services into HIV care and treatment services for women living with HIV. Our goal was to identify the models of integrating STI services into HIV care and treatment services for people living with HIV that have been evaluated and their associated outcomes.

METHODS Definitions

There is no agreed upon definition of service integration and no clear distinction between integrated and non-integrated services.¹⁶ WHO has defined integration as the 'management and delivery of health services so that clients receive a continuum of preventive and curative services, according to their needs over time and across different levels of the health system'.¹⁷ For the purposes of this review, we used this definition of integration and focus on the service delivery level, rather than at the policy or other levels.

We considered STI services to include sexual history screening, information and advice on avoiding re/infection, access to partner notification services and disclosure support, when appropriate, and testing, treatment and/ or syndromic management of any STIs other than HIV since we were looking at the integration of STI services with HIV services. The integration of cervical cancer screening by Pap tests (or smears) into HIV care and treatment settings absent of one or more of the aforementioned STI services was not considered sufficient to constitute integrated STI services.

Research question and inclusion criteria

The review addressed the following research question in population, intervention, comparison and outcomes (PICO) format: should STI services be integrated into HIV care and treatment services for women living with HIV?

Population

Women living with HIV.

Intervention

STI services, including STI testing, STI treatment and/ or syndromic management, integrated into HIV care and treatment services.

Comparison

HIV care and treatment services without integrated STI services or standard of care.

Outcomes

(1) Uptake of STI services, (2) STIs (reinfection), (3) client satisfaction and service quality (interpreted broadly to include receipt of sexual and reproductive health or HIV services, etc) and (4) provider knowledge and attitudes about STI services for women living with HIV.

To be included in the review, an article had to meet the following criteria:

- 1. Comparative study (any study design with pre/ post or multiarm outcome comparisons) examining women living with HIV who received STI services including STI testing, STI treatment and syndromic management—integrated into HIV care and treatment services, compared with women living with HIV who received HIV care and treatment services without integrated STI services or standard of care.
- 2. Evaluated one or more of the key outcomes outlined in the PICO question above.
- 3. Published in a peer-reviewed journal prior to 16 February 2017.

Because this review was designed to inform WHO guidelines on sexual and reproductive health and rights of women living with HIV,¹⁸ we excluded studies conducted only among men (primarily populations of men who have sex with men), but included studies conducted among both women and men. No restrictions were placed based on location of the intervention. No language restrictions were used on the search. We followed Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines for reporting systematic review results.¹⁹ A full review protocol was developed and is available from the corresponding author on request.

Search strategy

We searched the following electronic databases for articles published anytime through the search date of 16 February 2017: PubMed, CINAHL and EMBASE. Search terms were developed for STIs, HIV and integration and combined; the full search strategy for each database is available in online supplementary file 1. Secondary reference searching was conducted on all studies included in the review and three other related reviews.²⁰⁻²²

Screening abstracts

Titles, abstracts, citation information and descriptor terms of citations identified through the search strategy underwent an initial screening by a member of the study staff. Remaining abstracts then were screened in duplicate by two reviewers working independently with adjudication of differences by a senior member of the study team. Fulltext articles were obtained of all selected abstracts, and two independent reviewers assessed all full-text articles for eligibility to determine final study selection.

Data extraction and management

For each study, the following information was extracted: study citation, location, setting, target group, intervention description, study design, unit of analysis, selection of participants, sample size, age, gender and reported numerical outcomes and results.

Assessment of study rigour was conducted based on a nine-item tool, including items for (1) study design includes preintervention/postintervention data, (2)study design includes control or comparison group, (3) study design includes cohort, (4) comparison groups equivalent at baseline on sociodemographics, (5)comparison groups equivalent at baseline on outcome measures, (6) random assignment (group or individual) to the intervention, (7) participants randomly selected for assessment, (8) control for potential confounders and (9) follow-up rate $\geq 75\%$. This tool was based on the eight-item rigour assessment for systematic reviews of HIV behavioural interventions with a range of study designs (beyond randomised trials) developed by the Evidence Project.^{23 24}

Data analysis

Data were analysed according to coding categories and outcomes. Meta-analysis was not conducted due to the small number of included studies.

RESULTS

Online database searching yielded 406 hits, of which there were 235 unique citations after removing duplicates (figure 1). After initial screening of titles and abstracts, 104 citations remained for double-screening. These were double-screened and 40 were pulled for full-text review. Of these, one study (reported in two articles) was considered eligible for inclusion. In addition, two studies were included from the prior systematic review covering articles through 2007.²⁰ Therefore, a total of three studies reported in four articles met the criteria for inclusion in the review. Table 1 presents descriptions of the study settings, methods and outcomes. Table 2 presents an assessment of study rigour.

Of the three included articles, two evaluated the creation of new sexual health clinics within existing services for people living with HIV in the UK. The first study evaluated an integrated sexual healthcare clinic for HIV-positive women that included screening for STIs, along with other sexual health services including contraception, preconception counselling and cervical cytology.²⁵ The per cent of clients who were offered an STI screening increased from 70% before integration to 83% at 12-month follow-up. Other measures of service quality, including recording of sexual history and uptake of cervical cytology, also improved. In the second study, a sexual health clinic was similarly opened for HIV-positive adults that included STI screening, along with other services including hepatitis B, vaccination and cervical smear testing.²⁶ The per cent of clients receiving an annual STI screening increased from 26% before integration to 46% afterwards (p=0.003), while the per cent receiving an STI screening at their first appointment increased from 39% to 52% (p=0.065). Other measures of service quality, including documentation that an annual health discussion had taken place between provider and patient and uptake of cervical cytology, also generally improved.²⁶

The final study, reported in two articles,^{27 28} was a comparative case study examining the relationship between integration and the use and demand for sexual and reproductive health services (including STI services), and client satisfaction across different models of care in Swaziland: two clinics integrated with sexual and reproductive health services and two stand-alone HIV clinics (which did not provide STI services). In addition to employing a different study design than the previous two studies (a comparison across clinic models rather than a comparison before and after initiation of new services), this study had slightly higher overall rigour assessment as it randomly selected clients for assessment and adjusted for confounders (table 2). An exit survey (n=602) measured service use and satisfaction across models using logistic regression; a complementary qualitative assessment was also conducted. Women attending the fully integrated site were more likely to have had a Pap smear since testing positive than women attending the less integrated service models (35% vs 11%, 14% and 10%, p<0.001). However, clients attending the fully integrated site were not more likely to report having sexual health screening (45% vs 57%, 76% and 42% from most to least integrated, respectively) or advice on sexual health (16% vs 12%, 16% and 11% from most to least integrated). Client satisfaction was higher in the fully stand-alone clinic compared with both the partially integrated site (adjusted OR (aOR)

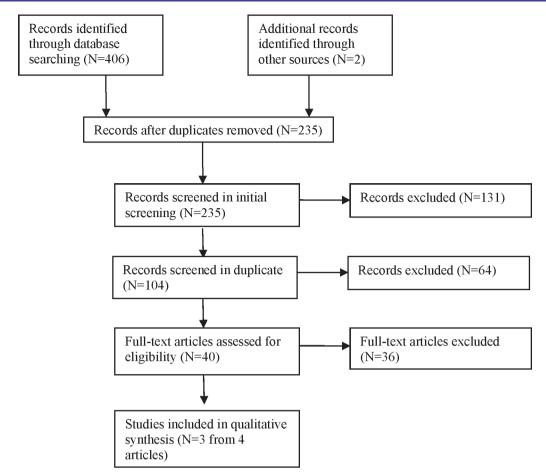


Figure 1 Disposition of citations during the search and screening process.

0.45, 95% CI 0.25 to 0.81) and the partially stand-alone site (aOR 0.53, 95% CI 0.31 to 0.90), but there was no difference in satisfaction between the fully integrated and fully stand-alone sites.²⁶ There was no difference in satisfaction by gender (aOR comparing men to women: 0.88, 95% CI 0.53 to 1.43).²⁷ A very small number of clients were referred for STI screening (1–2 per clinic).²⁸

DISCUSSION

Despite significant STI burden among women living with HIV and increased attention to integration of services, there is only modest evidence on the impact of integrating STI services into HIV care and treatment centres. We identified just three studies from two countries that provided comparative data on our outcomes of interest. In two studies from the UK, quality and uptake of STI services seemed to improve following introduction of new sexual health clinics into existing HIV services. However, in Swaziland, people living with HIV generally reported higher satisfaction at a stand-alone HIV clinic than at integrated service sites, and clients attending the integrated sites were no more likely than those at non-integrated sites to have had a sexual health screening. In this latter study, a diverse range of factors related to implementation of different care models challenged the notion that integrated services are always superior or desired; instead,

the authors concluded that different models may work for different types of clients and the diversity of client needs and preferences, along with health system factors, should be considered when developing integrated services.²⁷

Our review identified only a small number of articles that met the inclusion criteria, limiting the evidence base from which we can draw conclusions. Further, the study designs were cross-sectional or serial cross-sectional with modest sample sizes, limiting the inferences that can be drawn. The number and type of outcomes were also limited. The evidence gap does not necessarily reflect the lack of integrated STI and HIV services, but rather the paucity of comparative evaluations of such integration. Notably, because many national HIV guidelines indicate that STI screening should occur within routine HIV clinical care visits, evaluations may have focused on routine programme monitoring and quality assurance rather than comparing models of service delivery.²⁹ For example, in Scotland, an audit of facilities serving people living with HIV found that less than half of sites met the guidelines for offering STI screening and recording sexual history among at least 80% of patients within 4weeks of diagnosis, with wide variability in performance across sites.³ In Thailand, quality improvement initiative among 12 HIV care sites documented a significant increase in coverage of STI services, including an increase from 0%

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	Outcomes	 Pap smear since testing HIV positive (wormen) A: 35%, B: 11%, C: 14%, D: 10% C. Sexual health screening since testing HIV positive (men and women) A: 45%, B: 57%, C: 76%, D: 42% Counselling on condom use (men and women) A: 76%, B: 78%, C: 95%, D: 87% A: 70%, B: 78%, C: 95%, D: 87% A: 10%, B: 39%, C: 56%, D: 77% A: 11%, B: 39%, C: 56%, D: 77% A: 11%, B: 39%, C: 56%, D: 77% A: 11%, B: 39%, C: 56%, D: 77% A: 16%, B: 12%, C: 16%, D: 11% Mean number of providers seen during women) A: 16%, B: 12%, C: 16%, D: 11% A: 16%, B: 12%, C: 55%, D: 41% A: 16%, B: 44%, C: 59%, D: 41% Clients who would have liked to receive STI services (men and women) A: 44%, B: 48%, C: 59%, D: 41% Client satisfaction score24 was lower in models B (aOR 0.45, 95% CI 0.25 to 0.81) and C (aOR 0.53, 95% CI 0.21 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from model 1 (aOR 0.84, 95% CI 0.31 to 0.90) and not statistically significantly different from
	Sample size	Sample size: Total: 602 men and women; 475 women A: 71 men and women B: 163 men and women D: 190 men and women D: 190 men and
	Study design	<i>Study design:</i> Cross- sectional Unit of analysis: Individual Selection of participants: Systematic random sampling
sported outcomes	Intervention	Among the four models of HIV care evaluated, the most fully integrated clinic provided reproductive health services including FP, STI screening and care, and Pap smears, theoretically provided by one provider in one room (model A). A partially integrated clinic provided HIV and reproductive health services, FP, STI and Pap smears by different providers in different rooms within the same building (model B). A third partially stand-alone model provided HIV treatment on the campus of a larger district hospital where reproductive health services were provided in different buildings (model C). A fourth fully stand-alone HIV clinic did not offer reproductive health services (model D).
Descriptions of included studies and reported outcomes	Location, setting and target group	Location: Manzini, Swaziland Target group: HIV-positive women and men attending an HIV clinic
Table 1 Description	Study	Church <i>et al</i> ²⁸ and Church <i>et al</i> ²⁷

Continued

Table 1 Continued					
Study	Location, setting and target group	Intervention	Study design	Sample size	Outcomes
Coyne et al ²⁵	<i>Location:</i> Slough, UK <i>Target group:</i> HIV-positive women attending an HIV clinic	The Garden Clinic, for HIV- positive women, started a specific clinic (FP Plus) to provide HIV-positive women clients with screening for STIs, contraception, pre- conception counselling and cervical cytology. The Garden Clinic already worked on a model of integrated sexual healthcare, and FP Plus was staffed by doctors and senior nurses trained in both STI management and FP.	<i>Study design:</i> Serial cross-sectional <i>Unit of analysis:</i> Individual <i>Selection of participants:</i> Non-random	S <i>ample size:</i> Total: 60 women Before: 30 women After: 30 women	 Using condom only as contraception (women) Before: 30% After: 7% Cervical cytology in 12 months (women) Before: 63% After: 83% After: 83% After: 83% After: 63% After: 63% After: 63% After: 63% After: 77% After: 77% After: 97% STI screen offered in 6 months (women) Before: 70% After: 83%
Hamlyn <i>et al</i> ²⁶	<i>Location:</i> London, UK <i>Target group:</i> HIV-positive adults attending an HIV clinic	At the Caldecot Centre, an audit showed the need for a specific sexual health clinic for HIV-positive patients. Nurse- led STI clinics were set up on both a booked appointment and an emergency walk-in basis.	<i>Study design:</i> Serial cross-sectional <i>Unit of analysis:</i> Individual <i>Selection of participants:</i> Non-random	Sample size: Total: 200 men and women Before: 100 men and women; 48 women women; 48 women	 Positive STI result at first appointment (men and women) 2. Positive STI result at annual STI screen (men and women) 1.5 (33%), C: 12 (31%) 2. Positive STI result at annual STI screen (men and women) 1.5 (33%), C: 10 (38%) 3. STI screen at first appointment (men and women) 1.5 (33%, I vs C NS (p=0.065) 4. Annual STI screen (men and women) 1.46%, C: 39%, I vs C Sig (p=0.003) 5. Screening for hepatitis B (men and women) 1.46%, C: 26%, I vs C Sig (p=0.003) 5. Screening for hepatitis B (men and women) 1.46%, C: 24%, I vs C NS (p=0.129) 7. Documented annual sexual health discussion (men and women) 1.70%, C: 37%, I vs C Sig (p<0.0001)
aOR adjusted OB: C. c	omparison aroup. FP family pl	aOB adjusted OB: G. comparison group: EP family planning: 1 intervention group: NS not significant: Sig significant: STI sexually transmitted infection	significant: Sig significant	STI sexually transmitted	infection

aOR, adjusted OR; C, comparison group; FP, family planning; I, intervention group; NS, not significant; Sig, significant; STI, sexually transmitted infection.

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Table 2 Study rigour	rigour								
Study	Study design includes preintervention/ postintervention data	Study design includes control or comparison group	Study design includes cohort	Comparisc groups Study Comparison equivalent design groups equivalent at baseline includes at baseline on on outcom cohort sociodemographics measures	Comparison groups equivalent at baseline on outcome measures	Random assignment (group or individual) to the intervention	Participants randomly Control for selected for potential Follow- assessment confounders up rate	Control for potential confounders	Follow- up rate
Church <i>et al</i> ²⁸	No	Yes	No	N/A	N/A	No	Yes	Yes	N/A
Coyne <i>et al ²⁵</i>	Yes	No	No	N/A	N/A	No	No	No	N/A
Hamlyn <i>et al ²⁶</i> Yes	Yes	No	No	No (except for age)	N/A	No	No	No	N/A
N/A, not applicab	N/A, not applicable for the given study design.								

to 94% in median coverage for syphilis serology, and an increase from 0% to 67% in cervical cancer screening.²⁹ As part of the quality improvement initiative, a checklist of services that included STI screening was developed and a new standard operating procedure was initiated offering routine gynaecological examination and Pap smears to all HIV-infected women. In settings where STI services are available in HIV clinics, quality improvement and performance monitoring frameworks may be valuable approaches to improve outcomes, while also documenting service delivery strategies and coverage.

Further documentation of diverse service models using comparative designs would help to inform programme implementation. However, there is evidence from community-based clinic settings that STI screening can be increased at relatively low cost through strategies such as the strategic placement of specimen collection materials or automatic specimen collection as part of a routine visit, the use of electronic health records and the use of patient reminders (via text, telephone and postcards).³¹ Such strategies would likely work in HIV care and treatment settings as well.

Several service integration initiatives hold promise for better provision of STI services within HIV care and treatment settings. Many PEPFAR-supported sites have rolled out elements of positive health, dignity and prevention which includes STI services for people living with HIV, so in future years more evidence of STI/HIV integration may be forthcoming, particularly from high-HIV burden settings.³² The integration of cervical cancer screening in HIV treatment sites in resource-limited settings in recent years has demonstrated feasibility and acceptability.^{33 34} The creation of private spaces for pelvic examinations provides an opportunity for STI screening among HIV-infected women. In the USA, women living with HIV had 1.9 times the odds of having a Pap smear in the last year if gynaecology services were co-located with their primary HIV care³⁴ and a recent study found that women with HIV were significantly more likely to have been screened for gonorrhoea and chlamydia if they had also had a Pap smear in the past year.³³

As country initiatives in the elimination of mother-tochild transmission and Option B+ (immediate initiation of pregnant women on antiretroviral therapy) are expanded through the Start Free Stay Free AIDS Free framework of the Global Action Plan,35 more women living with HIV are being diagnosed in antenatal and postnatal care settings where HIV care and treatment is also being provided. With the WHO strategy for dual elimination of HIV and syphilis,³⁶ antenatal and postnatal care settings may become key sites for the integration of STI services with HIV care and treatment. Further, antenatal care settings and other maternal, neonatal and child health services may require integration with STI services regardless of whether HIV care and treatment services are also provided, as the assumption that sexual and reproductive health services are already a comprehensive, integrated set of services is often not accurate.

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In line with a people-centred approach, grounded in principles of human rights, gender equality and health equity, the WHO global strategies on HIV, STIs and viral hepatitis aim to facilitate people's access to information, improve access to treatment and comprehensive long-term care when needed and challenge pervasive stigmatisation and discrimination. The provision of STI services within HIV treatment settings is one way to reach many national STI screening guidelines for women living with HIV. Despite the limited evidence base, the studies identified in this review did document the feasibility of integrating STI services into HIV care and treatment settings and the potential for positive outcomes of such integration. Programmes should be encouraged by this evidence to critically consider whether integration might prove beneficial in their particular settings to improve access to comprehensive, holistic services that address the sexual and reproductive health needs of women living with HIV.

Correction notice This paper has been amended since it was published Online First. Owing to a scripting error, some of the publisher names in the references were replaced with 'BMJ Publishing Group'. This only affected the full text version, not the PDF. We have since corrected theseerrors and the correct publishers have been inserted into the references.

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Contributors CEK developed the data collection tools with input from SAH. CEK and SAH led the search, screening and data abstraction processes. CEK drafted the initial manuscript. All authors developed the study protocol, reviewed and approved the final manuscript draft and interpretations.

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