

Integrating HIV Pre-Exposure Prophylaxis into Family Planning Care: A RE-AIM Framework Evaluation

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Abstract

We aimed to systematically evaluate the feasibility of integrating HIV prevention services, including pre-exposure prophylaxis (PrEP), into a family planning setting in a high-prevalence community. We used the RE-AIM Framework (Reach, Efficacy, Adoption, Implementation, Maintenance) to evaluate the integration of HIV prevention services into a family planning clinic over 6 months. Before the integration, PrEP was not offered. We implemented a staff training program on HIV PrEP. We determined the proportion of women presenting to the clinic who were screened, eligible for, and initiated PrEP through chart review. We assessed staff comfort with PrEP pre- and post-integration. We compared planned and actual implementation, interviewed staff to determine barriers and facilitators, and tracked systems adaptations. We assessed maintenance of PrEP after the study concluded. There were 640 clinical encounters for 515 patients; the rate of HIV counseling and PrEP screening was 50%. The rate was 10% in month 1 and peaked to 65% in month 3. Nearly all screened patients were eligible for PrEP (98.4%) and 15 patients (6%) initiated PrEP. Staff knowledge and comfort discussing PrEP improved after education. Facilitators included partnering with local experts, continuing education, clinical tools for providers, and patient education materials. Barriers included competing priorities during clinical encounters, limited woman-centered patient education materials, and insurance-related barriers. Embedding HIV prevention services in the family planning setting was feasible in this pilot. The proportion of women screened for PrEP rapidly increased. In this high HIV prevalence community, nearly all screened women were eligible and 6% initiated PrEP.

Keywords: HIV prevention, PrEP, women, family planning, implementation

Introduction

DESPITE REPRESENTING 20% of new HIV cases in the United States, women represent less than 5% of HIV pre-exposure prophylaxis (PrEP) users.^{1–3} High-risk sexual activities that increase risk for HIV also increase risk for unintended pregnancy, leading women to seek care in family planning clinics. Women's health and HIV prevention

experts have called for increasing access to HIV prevention and PrEP in settings that provide family planning services to women.^{4–6}

PrEP with daily tenofovir/emtricitabine (Truvada[®]) reduces transmission of HIV by up to 92%.^{7–9} CDC and American College of Obstetricians and Gynecologists (ACOG) recommend HIV PrEP for women who “engage in sexual activity in a high prevalence area” and inconsistently

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use condoms.^{10,11} Yet, local and state efforts to curb the HIV epidemic in the United States have not consistently translated into offering PrEP more widely to high-risk cisgender women.

Despite the synergies in offering PrEP alongside contraception, pregnancy termination, and sexually transmitted infection (STI) services, most family planning clinics do not offer PrEP. In 2016, Seidman et al. demonstrated low levels of PrEP knowledge and comfort prescribing PrEP among family planning providers. Most providers reported interest in learning about PrEP and the majority of family planning patients reported they would consider using PrEP, but providers reported multiple barriers to PrEP provision, including insufficient provider education, patient costs, and time constraints.¹² While suggestions have been made regarding implementation of PrEP for women,⁶ the feasibility, facilitators, and barriers to the integration of HIV PrEP services in family planning settings have not been systematically investigated.

We aimed to evaluate the feasibility and impact of integrating HIV PrEP services with family planning services using the RE-AIM Framework. RE-AIM is a widely accepted framework used to assess the feasibility, quality, and public health impact of a health intervention.¹³ The framework includes five dimensions: *Reach* intended population, *Efficacy* or *effectiveness*, *Adoption* by target staff, settings, or institutions, *Implementation* consistency and adaptations, and *Maintenance* of the intervention over time.^{13,14} We hypothesized that most family planning patients in a high-prevalence community would be eligible for PrEP and that some patients would be interested in initiating PrEP. We hypothesized that the integration would be feasible and that RE-AIM analysis would identify barriers and facilitators to PrEP integration.

Methods

Study design and setting

We conducted a 6-month prospective cohort study, including clinic staff and female patients at MedStar Washington Hospital Center's Family Planning and Preventative Care clinic (FPPC), from September 2017 to March 2018. FPPC is an outpatient family planning clinic within an urban tertiary care hospital in Washington, DC. FPPC is a training site for fellows, medical students, and residents in family planning, obstetrics/gynecology, family medicine, and pediatrics offering comprehensive reproductive health care for women, including contraception, abortion, STI testing and treatment, and gynecology. The majority of FPPC patients are younger than 30 years, African American, and publicly insured. Most self-refer for abortion and contraceptive services. Clinical services are not offered to male partners because the clinic is staffed by gynecologists. Before the study, PrEP was not offered to FPPC patients. IRB approval was obtained from MedStar Health Research Institute.

We developed a staff and provider training program before the integration of PrEP services. Providers (physicians) and staff (nurses, medical assistants, and receptionists) were independently trained by a local HIV PrEP expert, who was additionally available for questions and phone consultations. Providers were trained to counsel, prescribe, and follow-up patients who initiated PrEP. Family planning patients were to

be verbally screened for PrEP during the history-taking portion of the encounter and offered initiation if eligible based on CDC recommendations (Table 1). Patients interested in PrEP but unable to initiate through our clinic due to insurance barriers or patient preference were linked to local PrEP clinics and were included in the study as participants who did not initiate PrEP.

One week after training was completed, universal HIV prevention and PrEP integration were implemented. All patients were exposed to a 5-min video on PrEP (www.whatisprep.org)¹⁵ in the waiting room, following which providers assessed risk factors for HIV and discussed HIV prevention. Patients choosing to initiate PrEP completed same-day laboratories, were provided a 30-day prescription, and scheduled for a follow-up appointment in 4 weeks, at which time their HIV test was repeated and a 3-month supply of PrEP was prescribed. Quarterly follow-ups were subsequently scheduled. Women interested but not ready to initiate PrEP were offered another appointment to discuss HIV prevention.

We administered quantitative surveys and qualitative interviews to staff and providers before training ($N=8$), immediately post-training ($N=8$), and 1 ($N=7$), 3 ($N=6$), and 6 months ($N=6$) post-training. The written anonymous multiple-choice survey was administered by a research coordinator and assessed staff and provider self-rated knowledge, attitudes, efficacy, and experience with PrEP. All quantitative survey items used 5-point Likert scales. Semistructured interviews were administered individually by the research coordinator and assessed satisfaction with implementation, perceived barriers and facilitators to PrEP implementation, and suggestions for improvement.

Outcomes

Constructs, outcomes, measures, and data sources are summarized in Table 2.

Reach. We determined the number of clinical encounters and individuals seen for family planning visits during the study by reviewing the schedule in the electronic medical record (EMR). We reviewed the EMR for family planning patient encounters during the study period to extract data on

TABLE 1. SUMMARY OF CDC ELIGIBILITY FOR PRE-EXPOSURE PROPHYLAXIS USE FOR WOMEN WHO HAVE SEX WITH MEN^{2,3}

Substantial HIV risk	Sexual partner with HIV Recent bacterial STD High number of sex partners History of inconsistent or no condom use Commercial sex work Lives in high-prevalence area or network
Clinical eligibility	Documented negative HIV test before prescribing PrEP No signs/symptoms of acute HIV infection Normal renal function, no contraindicated medications Documented hepatitis B virus infection and vaccination status

PrEP, pre-exposure prophylaxis.

TABLE 2. RE-AIM CONSTRUCTS AND ANALYSIS PLAN

<i>Construct</i>	<i>Definition</i>	<i>Outcomes</i>	<i>Measures</i>	<i>Data source</i>
Reach	Individual-level measure of participation	Proportion and characteristics of women counseled on HIV prevention and screened for PrEP eligibility	Number of encounters	EMR
			Number of patients served	EMR
			Patient characteristics	EMR
			Proportion of patients counseled	EMR
Effectiveness	Achievement of program goals and objectives	Patient uptake of PrEP	Proportion of counseled patients eligible for PrEP	EMR
			Proportion of patients who initiated PREP	EMR
		PrEP continuation rate	One-month PrEP follow-up rate	EMR
			Staff and provider perception	Staff and provider knowledge of PrEP
Adoption	Assessment of the delivery setting	Description of the setting	Provider experience prescribing PrEP	Quantitative survey
			Provider comfort prescribing PrEP	Quantitative survey
			Proportion of staff and providers who participated in the intervention	Training log and quantitative survey
			Description of actual vs. planned implementation	Training and implementation logs
Implementation	Extent to which the intervention is delivered	Adherence to the planned intervention	Barriers and facilitators to adoption	Qualitative interviews
			Trends in patient counseling and screening	EMR
			Description of adaptations	Implementation log and qualitative interviews
Maintenance	Institutionalization of the program	Adherence to the planned integration beyond the study time period	Current provider assessment of program integration	Qualitative provider comments
			Staff and provider satisfaction with the integration	Qualitative interviews

EMR, electronic medical record; PrEP, pre-exposure prophylaxis.

age, race, ethnicity, and visit type (pregnancy termination, contraception, or other). We assessed the proportion of patients screened for PrEP by reviewing the clinical documentation.

Effectiveness. We evaluated the proportion of patients who initiated and continued PrEP by reviewing electronic prescriptions for PrEP and follow-up appointments for PrEP, respectively.

Adoption. We documented adherence to the planned training schedule. We determined the proportion of staff and providers who attended planned training through attendance logs. We assessed facilitators and barriers to the adoption of HIV prevention integration through surveys and interviews of staff and providers.

Implementation. The program goal was to offer universal HIV prevention counseling and PrEP eligibility screening. We logged adaptations and improvements made to the planned integration.

Maintenance. We surveyed and interviewed staff and providers to identify barriers to long-term integration of the

program and to assess provider satisfaction. We describe ongoing adaptations made to the program.

Analysis

We performed descriptive analysis of the RE-AIM outcomes described above. We assessed the association between patient characteristics, screening rates, and eligibility for PrEP using chi-square and Fisher’s exact tests. Statistical analysis was performed using Stata 14.2 (StataCorp LLC, College Park, TX, 2015). Qualitative data were recorded, transcribed, and deidentified. Qualitative data were examined for common themes, but thematic saturation was not expected due to the small sample size.

Results

Reach

There were 640 clinical encounters for 515 patients. Patients were predominately African American and >75% presented for abortion services (Table 3). Half ($n = 252$) were counseled on HIV prevention and screened for PrEP eligibility. Women who are African American race and those seeking abortion services were more likely to be screened for

TABLE 3. DEMOGRAPHICS, REASON FOR CARE, AND PRE-EXPOSURE PROPHYLAXIS SCREENING AMONG WOMEN PRESENTING FOR CARE AT A FAMILY PLANNING CLINIC IN WASHINGTON, DC

	Overall (N=515)	Screened (n=252)	Not screened (n=263)	p
Age in years	28.9 ± 7.5	27.6 ± 6.5	30.3 ± 8.1	<0.005
Race				<0.005
African American	344 (66.8)	200 (58.1)	144 (41.9)	
White	62 (12)	11 (17.7)	51 (82.3)	
Other/multiracial	90 (17.5)	32 (35.6)	58 (64.4)	
Unknown	19 (3.7)	9 (47.4)	10 (52.6)	
Ethnicity Hispanic/Latina	40 (7.7)	36 (32.5)	27 (67.5)	0.09
Reason for visit	N=640	n=271	n=369	<0.005
Abortion	486 (75.9)	249 (51.3)	236 (48.7)	
Miscarriage	33 (5.2)	13 (39.4)	20 (60.6)	
Contraception	82 (12.8)	8 (9.8)	74 (90.2)	
Other gynecologic care	40 (6.3)	1 (2.5)	39 (97.5)	

N=515 patients except where specified. Age presented as mean ± SD. All other data presented as n (%).

PrEP eligibility compared with all patients (58.1% and 51.3%, $p < 0.005$).

Effectiveness

Four physicians, two medical assistants, and two nurses completed the baseline survey ($N=8$). Seven of eight were very comfortable discussing sexual health and risk reduction with female patients. Seven of eight had heard of PrEP. None was previously trained on PrEP, prescribed PrEP, or referred a patient for PrEP. Self-reported knowledge of PrEP was poor or fair for all participants. Mean comfort screening for PrEP eligibility was 3.25 on a 5-point Likert scale (1=very uncomfortable, 5=very comfortable). The most common perceived barriers to offering PrEP were staffing and time constraints, lack of training, lack of institution clinical guidelines, and insurance coverage.

At month 3, knowledge of PrEP was rated as good or very good for all respondents ($N=6$). Mean comfort screening for PrEP eligibility was 4.3 on a 5-point scale. Mean physician comfort prescribing PrEP increased from 1.5 at baseline to 3 on a 5-point Likert scale ($n=3$). All surveyed physicians had discussed PrEP with patients and two of three had prescribed PrEP. The majority of respondents agreed that all providers should be trained to counsel and prescribe PrEP. By 6 months, five of six respondents reported good or very good knowledge of PrEP, mean comfort screening for PrEP eligibility decreased to 3.2, and mean physician ($n=3$) comfort prescribing PrEP decreased to 2.7.

Nearly all screened patients were eligible for PrEP (98.4%), all due to no or inconsistent condom use in a high-prevalence area. Fifteen patients (6%) initiated PrEP and three (1%) continued its use beyond 1 month.

Adoption

All staff and providers participated in the initial planned training, although not all completed research surveys. A theme that emerged during interviews was the request for additional training, especially during 3- and 6-month interviews. Planned pharmacy and billing training was not completed. Instead, providers were trained on billing- and pharmacy-related barriers to PrEP initiation and compliance.

Facilitators to adopting the integration of PrEP services included the development of tools that served as reference guides for providers (“job aids”). Staff mentioned the benefit of having patients watch the PrEP video in the waiting room, stating it oriented patients to PrEP and facilitated efficient discussions with providers.

Two systems barriers were identified by staff: insurance coverage and the need for ongoing training. Many patients seeking abortion services paid out-of-pocket and our clinic was not within their insurance network. Thus, the baseline laboratories required for PrEP initiation would not be covered by insurance if completed in our clinic. Patients with DC Medicaid were required to obtain their PrEP at specific pharmacies, so a list of participating pharmacies was provided to these patients. The difficulty of providing consistent training to rotating residents and medical students, as well as newly hired staff, was also a barrier. A suggestion was that a single provider should be designated as a “PrEP expert” or “champion” who would address complicated questions about PrEP and provide PrEP follow-up.

Other barriers included time constraints, lack of on-site psychosocial support for risk reduction and adherence counseling, sensitivity to the emotional needs of patients seeking abortion, and the importance of addressing the patient’s primary reason for the visit rather than HIV prevention. We could find no high-quality printed PrEP education materials for cisgender women at the initiation of the study. By the 6-month time point, ACOG had developed patient education materials on PrEP, which we utilized.

Implementation

Before implementation, women seeking care at FPPC were not routinely counseled on HIV prevention and none was offered PrEP. During the 6-month pilot, 50% of patients were screened for PrEP eligibility, rapidly increasing from 10% in the first month and peaking at 65% in the third and fourth months. Screening then decreased slightly (Fig. 1).

The program underwent several adaptations. Follow-up training was developed and offered through teleconference with all providers and staff participating. Training was developed for newly hired staff. Formal training was offered to medical students and residents, but they could seldom attend due to schedule conflicts. One physician in the clinic

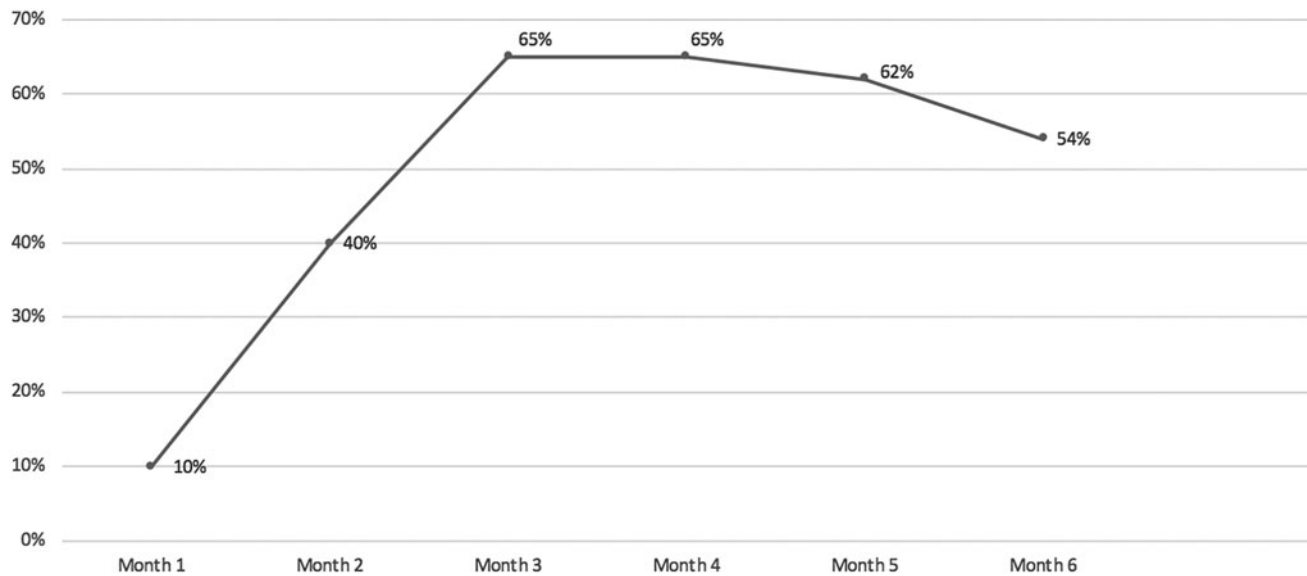


FIG. 1. Trends in counseling on HIV prevention and PrEP eligibility at a family planning clinic in Washington, DC during the 6-month PrEP Integration Program. PrEP, pre-exposure prophylaxis.

provided brief PrEP training to rotating medical students and residents, but this was implemented inconsistently. Job aids outlined PrEP initiation procedures (e.g., laboratory orders, prescription details, and contraindications) and billing codes while “quick-texts” standardized documentation. Similar tools were developed for PrEP follow-up appointments. The patient intake form was revised to include an item assessing the patient’s interest in discussing HIV prevention. Contraceptive option videos were looped with the PrEP video because some patients complained that the video repeated too frequently.

Maintenance

We continue to discuss HIV prevention with a majority of patients presenting for family planning services, facilitated by the updated intake form. New residents, students, and office staff are educated on the office protocols and reminded to counsel patients. Insurance coverage remains a barrier and we give women without in-network insurance coverage the ACOG brochure on PrEP and written referral information.

Discussion

This study systematically evaluated the integration of comprehensive HIV prevention services, including PrEP, in a family planning setting that primarily serves cisgender women. Staff and providers were eager to learn about PrEP and demonstrated improved knowledge and self-efficacy discussing and prescribing PrEP after training. Involvement of a local PrEP expert strengthened collaborations between sexual and reproductive health clinics, facilitated a referral process, and allowed family planning providers to develop a new skill set. PrEP counseling and screening grew rapidly in the first 3 months of implementation, with a peak screening rate of 65%.

We identified multiple facilitators to offering PrEP in our family planning clinic. Ongoing training was perceived as critical for staff and providers. This is consistent with na-

tional research demonstrating that lack of education and training is a major barrier to PrEP implementation.¹² Clinical tools such as job aids, order sets, and documentation aids improved efficiency for providers. Education materials in the waiting room prepared patients for discussions about HIV prevention. Data from family planning clinics in Georgia suggest that women want waiting-room materials on PrEP, such as brochures and posters, and that 39% wanted doctors to initiate discussions on PrEP.¹⁶ Woman-centered PrEP materials continue to be developed and a list of patient and provider resources can be found at hiveonline.org. It was also suggested that one clinician be identified as the “PrEP Champion,” to take the lead in addressing systems issues.

Lack of insurance coverage due to being out of network was consistently identified as an insurmountable barrier to PrEP uptake for interested patients at our location. We referred these patients to a nearby clinic. This is likely to be an issue in similar settings where patients pay out of pocket for sexual and reproductive health services. Insurance coverage of PrEP prescription was not a barrier; however, patients with Medicaid required referral to specific pharmacies. Insurance- and pharmacy-related issues illustrate a few of the structural barriers and biases in PrEP access. We hypothesize that decreased reported PrEP knowledge and confidence among providers at 6 months were due to turnover in trainees and staff. This reflects a missed opportunity for resident training on HIV prevention and PrEP and highlights the importance of training new providers, especially in sites with high turnover. Finally, providers questioned if discussing PrEP during visits sought for contraception and abortion was patient centered.

Limitations of our study include the single site, small sample size, and the 6-month time line. It is unknown whether universally screening and offering PrEP would significantly impact the workflow in high-volume clinics. Six months is a short time frame in which to assess implementation, and thus, factors related to maintenance are not fully explored. We reviewed EMR documentation of counseling and PrEP eligibility to assess the screening rate, which

TABLE 4. PRACTICES THAT SUPPORT INTEGRATING COMPREHENSIVE HIV PREVENTION SERVICES INTO FAMILY PLANNING CLINICS

Cultivation of relationships with local PrEP experts and organizations providing HIV prevention services
Development of a referral process for patients who need to initiate PrEP elsewhere due to insurance barriers
Development of a PrEP training program for staff and providers, implemented at regular intervals
Incorporation of PrEP training into on-boarding for staff, providers, and trainees
Assessment of insurance coverage and insurance networks of the patient population to anticipate insurance-related barriers
Utilization of woman-centered PrEP patient education materials
Designation of a “PrEP Champion” to lead systems initiatives and manage more complicated PrEP clinical scenarios

PrEP, pre-exposure prophylaxis.

would not account for counseling that occurred but was not documented. We hypothesize that the increased rate of screening among African American women and women seeking abortion was related to an increased risk perception by providers, but we did not assess reasons for nonscreening or the impact of bias among providers. Inequitable access to PrEP due to provider bias based on patient sociodemographic characteristics has been documented among both physicians and medical students.^{17,18} We also did not follow-up patients who were referred elsewhere for PrEP initiation.

Much remains unknown about the cascade of PrEP uptake for cisgender women. Prior research indicates that the majority of women attending family planning clinics who are at risk for HIV are interested in learning about PrEP.^{19–22} Our PrEP initiation rate was 6% and only three participants continued PrEP beyond 1 month, suggesting a disconnect between PrEP interest and utilization of PrEP. We hypothesize that many high-risk women are interested in PrEP but perceive that the barriers to PrEP use outweigh the benefits. Future research should investigate the cascade of events leading to PrEP initiation, continuation, and adherence among women and how family planning providers can better support PrEP uptake among interested women.

In this pilot, embedding comprehensive HIV prevention and PrEP services into family planning clinics was feasible and well received by staff and providers. Based on our experience, we developed a list of practices that support offering comprehensive HIV prevention in women’s health clinics (Table 4). Family planning clinics that provide abortion and contraception in high HIV prevalence areas are an ideal setting for PrEP provision, given that they predominantly serve women who are eligible for and would benefit from PrEP.

Author Disclosure Statement

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