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Racial/Ethnic and HIV risk category disparities in PrEP discontinuation among patients in publicly-funded primary care clinics.

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Abstract

Objective: Dissemination of pre-Exposure Prophylaxis (PrEP) is a priority for reducing new HIV infections, especially among vulnerable populations. However, there are limited data available on PrEP discontinuation following initiation, an important component of the PrEP cascade.

Design: Patients receiving PrEP within the San Francisco Department of Public Health Primary Care Clinics (SFPCC) are included in a PrEP registry if they received a PrEP prescription, were not receiving post-exposure prophylaxis, and not known to be HIV positive.

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Methods: We calculated PrEP discontinuation for patients initiating PrEP at any time from January 2012 to July 2017 and evaluated their association with demographic and risk variables using Cox regression analysis.

Results: Overall, 348 patients received PrEP over the evaluation period. The majority (84%) were men, and the cohort was racially/ethnically diverse. The median duration of PrEP use was 8.3 months. In adjusted analysis, PrEP discontinuation was lower among older patients (aHR 0.89; 95% CI: 0.80–0.99; $p=0.03$); but higher among Black patients (compared with White patients) (aHR 1.87; 95% CI: 1.27–2.74; $p=0.001$), patients who inject drugs (aHR 4.80; 95% CI 2.66–8.67; $p<0.001$), and transgender women who have sex with men (compared with men who have sex with men) (aHR 1.94; 95% CI: 1.36–2.77; $p<0.001$).

Conclusion: Age, racial/ethnic, and risk category disparities in PrEP discontinuation were identified among patients in a public-health funded primary care setting. Further efforts are needed to understand and address PrEP discontinuation among priority populations to maximize the preventive impact of PrEP, and reverse HIV-related disparities at a population level.

Keywords

PrEP discontinuation; racial/ethnic disparities

Introduction

Pre-exposure prophylaxis has been shown to have high efficacy in HIV prevention in several clinical trials and clinical cohorts.^[1–7] In the US, although PrEP uptake is increasing each year, uptake has been lower among Black and Latino patients than White patients in demonstration projects, clinical cohorts, and national reviews of pharmacy records.^[7–11] Similar disparities in adherence rates and PrEP discontinuation were observed in the US PrEP Demo project.^[12]

Continued PrEP use over time is an important step in the PrEP continuum, but there are limited data about PrEP discontinuation rates, particularly within primary care settings with diverse patient populations. In a Kaiser study of 972 patients who initiated PrEP, approximately 30% discontinued PrEP during the 3 year study follow-up period.^[13] In this clinical cohort, the risk of discontinuation was highest among women and those with a history of alcohol use. Similar discontinuation rates were noted in another clinical cohort of PrEP users within a San Francisco sexual health clinic.^[14] In an LGBT focused health center in Chicago, only 43% were still in care after 12 months, with uninsured patients less likely retained in care.^[15] However, these cohorts were in clinical settings focused on PrEP, and not necessarily reflective of patients in general primary care practice. Furthermore, both cohorts were nearly all men and approximately 70% White, not reflecting the racial/ethnic demographics of new HIV diagnoses in the US.^[16] As PrEP dissemination expands, more data are needed on implementation within primary care settings with diverse patient populations.^[17, 18]

In this study, we assessed PrEP discontinuation rates among individuals who initiated PrEP within the San Francisco Department of Public Health Primary Care Clinics (SFPCC), an

integrated safety-net primary care network that provides care for uninsured and publically-insured patients within San Francisco. We evaluated PrEP use and factors associated with PrEP discontinuation among patients who initiated PrEP from 2012–2017.

Methods

Patients receiving PrEP within the SFPCC were initiated on PrEP by their primary care providers within one of 14 primary care clinics. PrEP initiation, prescription refills (filled through the patient's preferred commercial or community pharmacy), adherence counseling and follow-up visits, and laboratory testing was determined by the prescribing clinician. Clinical PrEP guidelines were developed in 2015 to support PrEP implementation including recommendation for quarterly laboratory testing, and clinic follow-up, and management of adverse events. Standard patient retention practices varied between clinics, and included follow-up phone outreach for patients with missed visits, but not proactive outreach for those with overdue visits or laboratory testing. All of the patients receiving PrEP are included in a registry to monitor PrEP use and related outcomes. The study was reviewed and approved by the University of California, San Francisco Institutional Review Board.

Patients were included in this study if they received PrEP at one of 14 SFPCC clinics and met the following criteria based on medical record review: 1) verified to have been provided a PrEP prescription by a clinician; 2) were not receiving post-exposure prophylaxis; and 3) were not known to be HIV infected through surveillance reporting or SFPDPH laboratory testing. Patients initiating PrEP from January 1st, 2012 through December 31st, 2016, with follow-up through July 2017 were included in this analysis. We obtained dates of PrEP initiation and discontinuation, age and race/ethnicity, insurance and housing status, self-reported substance use, mental health disorders, visit data, and PrEP prescription data via electronic health record abstraction. Race/ethnicity was reported as a single categorical variable based upon the availability of the data within the electronic health record. PrEP indication was collected through chart review and classified into the following exclusive categories: sero-different relationship, men who have sex with men (MSM), people who inject drugs (PWID), transgender women who sex with men (TGWSM), or high-risk heterosexual.

Mental health diagnoses recorded for patients included anxiety, depression, post-traumatic stress disorder (PTSD), schizophrenia, and bipolar disorder; the latter two were considered serious mental illness (SMI). Data were captured on stimulant use (any amphetamine or cocaine/crack use mentioned by a medical provider) and heavy alcohol use (mention of heavy or disordered alcohol use by a medical provider) if it was reported in the medical chart. Patients with documentation from a provider of homelessness or unstable housing were classified as having housing instability.

PrEP panel management and patient navigation support were initiated at four of the 14 SFPCC clinics evaluated in this study starting in November of 2015. The primary care-based PrEP navigators within SFPCC were placed within four clinics: two youth clinics, an HIV care clinic which expanded to provide PrEP, and an LGBT clinic with embedded youth and transgender clinic services. The PrEP navigators provided ongoing support and active

outreach for patients who initiated PrEP and referrals to case management and other services as needed at two of the clinics, while pharmacists provided panel management (use of clinic registries to monitor patients who were due for visits, laboratory testing, or refills) without active outreach at the others. The navigators and/or the pharmacists at these clinics proactively reached out to patients on PrEP for upcoming (or overdue) clinical or laboratory visits, and supported patients to schedule and attend these visits. They also used clinic registries of patients who initiated PrEP to provide insurance support, adherence counseling, and laboratory and visit adherence support through in-person and telephone visits. Periods of PrEP use beginning after 11/1/2015 among patients at the four participating clinics were considered covered by panel management and patient navigation support. Patients in clinics without patient navigation received follow-up per provider and clinic procedures.

PrEP discontinuation was defined as either a documented discontinuation by a medical provider or a PrEP prescription gap of >90 days. Patients who were lost to follow-up were considered as discontinuing PrEP. Follow-up for discontinuation was censored at the end of July 2017. Kaplan-Meier plots were used to describe time from the beginning of each period of PrEP use to the first discontinuation, both overall and by race/ethnicity. Cox models with robust standard errors were used to assess correlates of earlier discontinuation. A total of 444 PrEP use periods among 348 patients were considered in the analysis reflecting that patients who discontinued PrEP could restart PrEP. All analyses were implemented using Stata Version 15.1 (Stata Corp, College Station, TX).

Results

Overall, 348 patients initiated PrEP during the observation period and had complete data for this analysis (Table 1). The majority were male sex at birth (84%), and the sample was ethnically/racially diverse with 39% White, 27% Latinx, and 12% Black patients. MSM comprised the most common PrEP indication (66%), although TGWSM accounted for 13%, 16% were in a sero-different relationship, and 4.9% were identified as high-risk heterosexual. Only two patients had a PrEP indication of PWID. Nearly one quarter of the sample had at least one gap in PrEP use after initiation, and the most common reason was being lost to follow-up (46%). Other reasons included missed visits or laboratory testing (44%), medication cost (12%), and decreased HIV risk (10%); there was no documented reason for 16% of gaps. More than three-fourths of the patients accessing PrEP had public insurance (Medicaid or Medicare) and 14% were uninsured; 12% had documented housing instability by a provider. Nearly half of the patients had a provider-documented mental health diagnosis, including 10% who had an SMI. Access to patient navigation services was available for approximately one quarter of the PrEP follow-up time in this analysis.

The median duration of PrEP use before discontinuation was 250 days (8.3 months) (Table 2 and Figure 1A). Black patients had much shorter durations of PrEP use compared with White patients (120 days vs 330 days) (Table 2 and Figure 1B). In adjusted analysis, older age was associated with a lower risk of having a gap in PrEP [adjusted Hazard Ratio (aHR) per decade of age 0.89; 95% Confidence Interval (CI): 0.80–0.99; $p=0.03$] (Table 3). Compared with White patients, Black patients were the only racial/ethnic group with a significantly higher risk of discontinuing PrEP (aHR 1.87; 95% CI: 1.27–2.74; $p=0.001$).

Patients who started PrEP with an indication of PWID (aHR 4.80; 95%CI 2.66–8.67, $p<0.001$), or who were TGWSM (aHR 1.94; 95%CI: 1.36–2.77; $p<0.001$) were more likely to discontinue PrEP compared with MSM. Patients who reported a history of illicit drug use were also more likely discontinue PrEP (aHR 1.55; 95% CI: 1.18–2.02; $p=0.001$) compared with patients with no history of illicit drug use. Panel management and PrEP navigation was not significantly associated with fewer PrEP discontinuations (aHR 1.23; 95% CI: 0.88–1.72; $p=0.23$).

Discussion

In this diverse cohort of primary care-based patients with multiple concomitant challenges such as housing instability, substance use and mental illness, PrEP discontinuation rates were higher than have been reported in research-based and specialty PrEP programs. PrEP discontinuation was higher for younger, Black, substance using, and TGWSM patients who are under-represented in published PrEP demonstration projects, but over-represented among new HIV diagnoses in the US. This is the first report demonstrating racial/ethnic differences in PrEP discontinuation among diverse patients in a safety-net primary care setting, where PrEP expansion has been a major focus for implementation. In our analysis, PrEP panel management with patient navigation was not associated with lower rates of PrEP discontinuation.

Although lower PrEP uptake and persistence has been previously reported among Black patients and youth in other PrEP demonstration projects and clinical settings in the US, these observations were all made in sexual health or specialty clinics.^[3, 19, 20] In the US PrEP Demonstration project, conducted at two sexual health clinics and a LGBT community health center, PrEP retention was also noted to be lower among Black compared with White participants during the 12 months of follow-up.^[12] Structural and social barriers such as stigma, medical mistrust, and perceived racism have been identified as important barriers for PrEP uptake, and likely impact PrEP continuation as well.^[21, 22] Even after adjusting for important individual factors in our analysis, disparities in PrEP discontinuations persist indicating social and structural barriers must better addressed to support Black patients continuing PrEP.

PWID had a high risk of PrEP discontinuation in this study, although only two patients were started on PrEP for this indication. PWID risk is a PrEP indication based on the Centers for Disease Control and Prevention PrEP guidelines; however there is little implementation data with this risk group. Our findings are similar to the data on lower HIV care engagement among PWID, suggesting that PWID need additional support across the HIV care and prevention continuum.^[23] Leveraging the experience from HIV treatment and prevention programs, integrating PrEP retention support into syringe access services, offering it in combination with methadone maintenance programs, or buprenorphine (in a primary care setting); and providing support for housing and other social needs have the potential to improve outcomes.^[24] Similarly, TGWSM also had a high of risk of PrEP discontinuation among our cohort of patients. Many of the prior clinical trials and demonstration projects enrolled a low proportion of trans women, limiting the ability to understand factors associated with PrEP uptake and discontinuation in this population. Community concerns

such as lack of targeting of PrEP for trans women, and the potential for drug-drug interactions with feminizing hormones, have been identified as potential barriers to PrEP uptake, while integration of PrEP within gender-affirming healthcare settings is a potential facilitator for PrEP engagement.^[25–27] These factors may also impact PrEP retention and discontinuation

Our analysis demonstrated that clinics with PrEP panel management or patient navigation did not have lower rates of PrEP discontinuations. Panel management, with associated registries, have been used as part of chronic disease management for several conditions including diabetes and cardiovascular disease.^[28] While the use of registries as part of quality improvement for diabetes management has been shown to be useful, one study suggests that impact is limited without attention to specific clinical actions.^[29] For example, in one study, improvements were made in diabetes screening, but without improvements in glycemic or blood pressure control. In the SFPCC, panel management was variably implemented within the different clinical settings by either patient navigators or a pharmacist, and further research into components of PrEP management that may lead to improved outcomes (eg., outreach to patients who missed appointments or follow-up laboratory testing) is needed. It is also possible that underlying systemic challenges at these clinics (e.g., not having integrated laboratory services or not having the ability to make follow-up appointments while in clinic), may also have contributed to the lack of improvement seen.

This analysis has several limitations. The number of patients accessing PrEP was relatively small in this clinical dataset compared with other clinical PrEP programs.^[12–14] This may be because of patients' ability to access sexual health services, including PrEP, through local municipal STD or a community-based sexual health clinic in San Francisco, allowing them to bypass receiving PrEP from their primary care provider. However, the population of PrEP patients in this study is more diverse (racially/ethnically and by PrEP indication) than reported in many other clinical settings, and is more reflective of the populations most at risk for HIV in the US. Another limitation was the use of electronic medical record review to identify patient cofactors. Because of variabilities in ascertainment and documentation by medical providers, reporting of these certain variables may be incomplete. Furthermore, a substantial proportion of patients were lost to follow-up over the study period. The causes for this lack of follow-up are likely variable and may include insurance change, relocation, seeking care elsewhere, and disengagement in care, although these reasons were rarely documented in the medical record. Because all patients lost to follow-up were classified as having discontinued PrEP, we may have biased our findings toward higher estimates of PrEP discontinuation if, in fact, patients continued PrEP outside of the SFPCC. Finally, we assumed that patients initiating PrEP were at ongoing risk for HIV throughout the evaluation period. Through the medical record review, we were not able to assess if PrEP discontinuations were secondary to decreases in HIV risk, and therefore appropriate discontinuations. SFPCC clinical guidelines encouraged providers to discuss HIV risk factors and assess ongoing need for PrEP through shared decision-making. However, many patients discontinued PrEP without consulting their providers and objective assessments of HIV risk were not always available or documented in the medical record.

We found that PrEP discontinuation was high among a diverse primary care-based cohort of patients receiving PrEP, with significant disparities among populations disproportionately impacted by HIV. Additional data are needed to understand patient, provider, and structural factors associated with PrEP discontinuation and interventions to support PrEP use for patients at risk for HIV.

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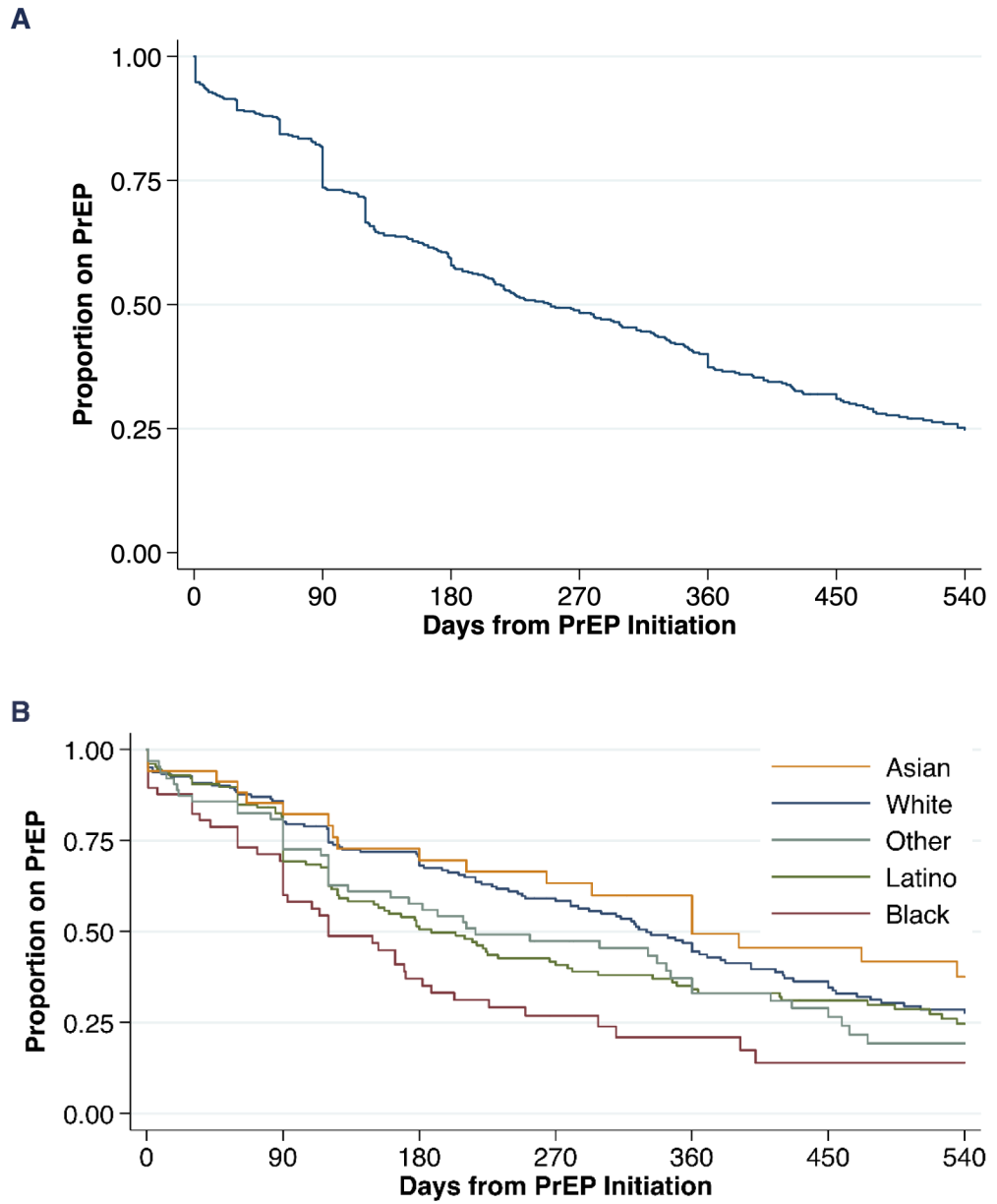


Figure 1: Overall Pre-exposure prophylaxis use over time (A) and by race/ethnicity (B) within San Francisco Department of Public Health primary care clinics.

Table 1:

Demographic, patient, clinic, and provider characteristics of patients initiating Pre-exposure prophylaxis (PrEP) within the San Francisco Department of Public Health primary care clinics.

Variable	N (%) N=348
<i>Patient</i>	
Birth Sex	
Male	293 (84)
Female	55 (16)
Age (median, IQR)	35 (28–45)
Race/Ethnicity	
White	135 (39)
Black	42 (12)
Latino	93 (27)
Asian	28 (8)
Other	50 (14)
PrEP Indication	
Men who have sex with men	230 (66)
Sero-different couple	55 (16)
Transgender women who have sex with men	44 (13)
Injection Drug Use	2 (1)
High-risk heterosexual	17 (5)
Insurance	
Private	27 (8)
Public (Medicaid or Medicare)	273 (78)
Uninsured	48 (14)
Housing instability	43 (12)
Mental Health Diagnosis ¹	
None	183 (53)
Anxiety	31 (9)
Depression	77 (22)
Post-traumatic Stress Disorder	23 (7)
Severe mental illness (Bipolar or Schizophrenia)	34 (10)
Heavy Alcohol Use ¹	61 (18)
Illicit substance use (heroin, cocaine, or meth) ¹	55 (16)
<i>Clinic and Provider</i>	
PrEP Panel Management or patient navigation	114 ² (26)
Number of PrEP patients seen by provider	
1	66 (19)

Variable	N (%) N=348
2-5	116 (33)
>5	169 (48)
<i>PrEP Initiation and Persistence</i>	
PrEP Initiation Year	
2012-2014	66 (19)
2015	114 (33)
2016	164 (47)
Number of gaps in PrEP use ³	
0	264 (76)
1	72 (21)
2	12 (3)

¹Based on provider assessment as documented in the patient medical record.

²Of 444 PrEP use periods

³Gaps were defined as interruptions of > 90 days in PrEP use.

Table 2:

Median days of PrEP use prior to discontinuation by demographics, PrEP indication, substance use, and PrEP panel management or patient navigation within San Francisco Department of Public Health primary care clinics.*

Variable	Median Days (95% Confidence Interval)
Overall	250 (210 – 310)
Age	
<25	90 (82 – 222)
25–39	226 (188 – 292)
40–54	360 (231 – 391)
55+	382 (179 – 450)
Race/Ethnicity	
White	330 (280 – 379)
Black	120 (90 – 171)
Latino	188 (127 – 270)
Asian	360 (211 – --)
Other	217 (120 – 346)
PrEP Indication	
Men who have sex with men	292 (222 – 347)
Sero-different couple	331 (183 – 391)
Transgender women who have sex with men	120 (69 – 178)
Injection Drug Use	30 (30 – --)
High-risk heterosexual	350 (85 – --)
Illicit substance use	
Yes	178 (120 – 239)
No	285 (221 – 343)
PrEP Panel Management or patient navigation	
Yes	161 (114 – 222)
No	298 (228 – 347)

* Discontinuation was defined as either a documentation of discontinuation by a medical provider or a PrEP prescription gap of >90 days.

Table 3:

Unadjusted and adjusted analysis of variables associated with discontinuations in Pre-exposure Prophylaxis use within San Francisco Department of Public Health primary care clinics.*

Variable	Unadjusted		Adjusted	
	Hazard Ratio	p-value	Hazard Ratio	p-value
<i>Patient</i>				
Birth Sex				
Male	(ref)		-	
Female	1.18 (0.86–1.62)	0.30	-	
Age (per 10 years)	0.85 (0.77–0.95)	0.004	0.89 (0.80–0.99)	0.03
Race/Ethnicity				
White	(ref)		(ref)	
Black	2.00 (1.38–2.89)	<0.001	1.87 (1.27–2.74)	0.001
Latino	1.26 (0.94–1.70)	0.12	1.08 (0.79–1.48)	0.62
Asian	0.80 (0.53–1.21)	0.30	0.73 (0.40–1.18)	0.20
Other	1.31 (0.92–1.85)	0.13	1.20 (0.84–1.72)	0.32
PrEP Indication				
Men who have sex with men	(ref)		(ref)	
Sero-different couple	1.02 (0.76–1.36)	0.92	0.92 (0.68–1.25)	0.60
Transgender women who have sex with men	2.04 (1.44–2.91)	<0.001	1.94 (1.36–2.77)	<0.001
Injection Drug Use	5.21 (2.55–10.6)	<0.001	4.80 (2.66–8.67)	<0.001
High-risk heterosexual	1.14 (0.61–2.14)	0.68	0.84 (0.40–1.74)	0.64
Insurance				
Private	(ref)		-	
Public (Medicaid or Medicare)	0.99 (0.66–1.48)	0.95	-	
Uninsured	1.18 (0.73–1.92)	0.50	-	
Housing instability	1.16 (0.83–1.60)	0.39	-	
Mental Health Diagnosis ¹				
None	(ref)		-	
Anxiety	0.92 (0.59–1.44)	0.73	-	
Depression	0.97 (0.73–1.28)	0.81	-	
Post-traumatic Stress Disorder	0.94 (0.57–1.54)	0.79	-	
Severe mental illness (Bipolar or Schizophrenia)	1.38 (0.96–1.98)	0.08	-	
Heavy Alcohol Use ¹	1.07 (0.81–1.43)	0.62	-	
Illicit substance use (heroin, cocaine, or meth) ¹	1.59 (1.23–2.05)	<0.001	1.55 (1.18–2.02)	0.001
<i>Clinic and Provider</i>				
PrEP Panel Management or patient navigation	1.48 (1.14–1.92)	0.003	1.23 (0.88–1.72)	0.23
Number of patients seen by provider				
1	(ref)		(ref)	

Variable	Unadjusted		Adjusted	
	Hazard Ratio	p-value	Hazard Ratio	p-value
2–4	1.08 (0.74–1.56)	0.69	1.00 (0.68–1.47)	0.93
>4	1.35 (0.98–1.86)	0.07	1.05 (0.74–1.50)	0.77
<i>PrEP Initiation and Persistence</i>				
PrEP Initiation Year				
2012–2014	(ref)		(ref)	
2015	1.35 (0.99–1.84)	0.06	1.15 (0.83–1.59)	0.40
2016	1.63 (1.19–2.24)	0.003	1.19 (0.81–1.75)	0.36

* Discontinuation was defined as either a documentation of discontinuation by a medical provider or a PrEP prescription gap of >90 days. PrEP discontinuation was evaluated using repeated measures.

¹Based on provider assessment as documented in the patient medical record.

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