HIV Preexposure Prophylaxis With Emtricitabine and Tenofovir Disoproxil Fumarate Among Cisgender Women

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**IMPORTANCE** Emtricitabine and tenofovir disoproxil fumarate (F/TDF) for HIV preexposure prophylaxis (PrEP) is highly effective in cisgender men who have sex with men (MSM) when adherence is high (>4 doses/week). Real-world effectiveness and adherence with F/TDF for PrEP in cisgender women is less well characterized.

**OBJECTIVE** To characterize the effectiveness of F/TDF for PrEP and its relationship with adherence in cisgender women.

**DESIGN, SETTING, AND PARTICIPANTS** Data were pooled from 11 F/TDF PrEP postapproval studies conducted in 6 countries that included 6296 cisgender women aged 15 to 69 years conducted from 2012 to 2020. HIV incidence was evaluated according to adherence level measured objectively (tenofovir diphosphate concentration in dried blood spots or tenofovir concentration in plasma; n = 288) and subjectively (electronic pill cap monitoring, pill counts, self-report, and study-reported adherence scale; n = 2954) using group-based trajectory modeling.

**EXPOSURES** F/TDF prescribed orally once a day. HIV incidence was analyzed in subgroups based on adherence trajectory.

**MAIN OUTCOMES AND MEASURES** HIV incidence.

**RESULTS** Of the 6296 participants, 46% were from Kenya, 28% were from South Africa, 21% were from India, 2.9% were from Uganda, 1.6% were from Botswana, and 0.8% were from the US. The mean (SD) age at PrEP initiation across all studies was 25 (7) years, with 61% of participants being younger than 25 years. The overall HIV incidence was 0.72 per 100 person-years (95% CI, 0.51-1.01; 32 incident HIV diagnoses among 6296 participants). Four distinct groups of adherence trajectories were identified: consistently daily (7 doses/week), consistently high (4-6 doses/week), high but declining (from a mean of 4-6 doses/week and then declining), and consistently low (less than 2 doses/week). None of the 498 women with consistently daily adherence acquired HIV. Only 1 of the 658 women with consistently high adherence acquired HIV (incidence rate, 0.13/100 person-years [95% CI, 0.02-0.92]). The incidence rate was 0.49 per 100 person-years (95% CI, 0.22-1.08) in the high but declining adherence group (n = 1166) and 1.27 per 100 person-years (95% CI, 0.53-3.04) in the consistently low adherence group (n = 632).

**CONCLUSIONS AND RELEVANCE** In a pooled analysis of 11 postapproval studies of F/TDF for PrEP among cisgender women, overall HIV incidence was 0.72 per 100 person-years; individuals with consistently daily or consistently high adherence (4-6 doses/week) to PrEP experienced very low HIV incidence.
he World Health Organization recommends the inclusion of oral preexposure prophylaxis (PrEP) containing tenofovir disoproxil fumarate (TDF) as an option for HIV prevention in individuals at substantial risk of acquiring HIV as part of combination HIV prevention approaches. The combination of emtricitabine and TDF (F/TDF) was approved for HIV PrEP to prevent sexual acquisition of HIV infection in adults by the US Food and Drug Administration (FDA) in 2012 and was extended to include adolescents in 2018. However, the relative efficacy of F/TDF for PrEP in cisgender women compared with men who have sex with men (MSM) has been questioned due to negative results in some randomized clinical trials in women and because of reports of differential concentrations of tenofovir in vaginal vs rectal mucosa. Mucosal concentration data, in particular, have informed distinct approaches to the counseling of women compared with MSM, with women being advised that near-perfect daily adherence is required for optimal efficacy of F/TDF for PrEP.

Among cisgender MSM, the relationship between adherence (the number of F/TDF doses taken per week estimated from tenofovir diphosphate concentrations in dried blood spots (DBS)) and prevention efficacy has been described. Analysis of the iPrEx Open Label Extension study found no new HIV infections in participants with tenofovir diphosphate concentrations indicative of taking at least 4 doses per week. These findings were consistent with modeling data that estimated an HIV risk reduction of 76% for 2 doses per week, 96% for 4 doses per week, and 99% for 7 doses per week among cisgender MSM.

Since the approval of F/TDF for PrEP, several global post-approval studies have been conducted to better define the effectiveness of F/TDF for PrEP in real-world settings and other contexts outside of randomized clinical trials, including in cisgender women. In the present analysis, data were pooled from 11 such studies in 6 countries (Botswana, India, Kenya, South Africa, Uganda, and the US) including 6296 women to evaluate F/TDF PrEP adherence and HIV incidence. Group-based trajectory modeling was employed, allowing leverage of a diverse range of quantitative and qualitative adherence data to examine the adherence-effectiveness relationship of F/TDF for PrEP in cisgender women. Using this approach, this study sought to determine the association between baseline demographics and other characteristics and adherence trajectories with HIV incidence in women in postapproval settings.

Methods

Study Selection

Data were analyzed from prospective studies involving oral F/TDF for PrEP in cisgender women and adolescents conducted between 2012, the year F/TDF for PrEP was approved for use in the US, and 2020. These studies were performed as part of the postmarketing commitment to the FDA and were either demonstration projects (n = 4) or open-label studies (n = 7), with 7 conducted in sub-Saharan Africa, 2 in India, and 2 in the US (Table). Study populations varied, but all included individuals who could benefit from PrEP and persons in serodiscordant (1 partner living with HIV and 1 partner without HIV) relationships, adolescent females and young women, and female sex workers.

Incidence

Between November 2012 and December 2020, a total of 6296 cisgender women aged 15 to 69 years initiated F/TDF for PrEP as participants in 11 prospective studies (Table). Participants were followed up regularly (follow-up visit frequencies across studies ranged from a mean of every 1.1-4.5 months) and underwent HIV testing using protocols in place at each study site. The “on-PrEP” HIV incidence rate was calculated by dividing the number of new HIV infection diagnoses by cumulative person-years of follow-up, determined as the time between PrEP initiation and the earliest date of HIV diagnosis, PrEP discontinuation, study end, or 96 weeks. The HIV incidence rate was calculated overall and in groups defined by adherence trajectories, as detailed below, to estimate effectiveness of F/TDF in preventing HIV acquisition using Poisson regression.

Adherence Measures

Adherence data were available for a subset (n = 2955 [47%]) of participants. Adherence was assessed by a variety of methods, which were dictated by the protocol for each independent study (Table). The majority were subjective measures (92%), including electronic pill cap monitoring, pill counts, self-report, or a study-reported adherence scales. Objective adherence measures included quantification of tenofovir diphosphate in DBS in 209 participants, reflecting adherence over the past 8 to 12 weeks, or of tenofovir in plasma in 46 participants, reflecting adherence over the past 2 to 7 days. Of the 46 participants with plasma tenofovir data, 18 also had tenofovir diphosphate in DBS measured. Adherence was categorized as daily (7 doses/week), high (4-6 doses/week), moderate (2-3 doses/week), or low (less than 2 doses/week) according to the following established cutoffs for tenofovir diphosphate in DBS: greater than or equal to 1249 fmol/punch, 699 to 1250 fmol/punch, 350 to 700 fmol/punch, and less than 350 fmol/punch, respectively. For the 46 participants who had tenofovir plasma data that were used in the group-based trajectory modeling (ie, not including those with concurrent tenofovir diphosphate measurements in

Key Points

Question What is the HIV incidence among cisgender women with different adherence trajectories to preexposure prophylaxis (PrEP) with emtricitabine and tenofovir disoproxil fumarate?

Findings In this pooled analysis of 6296 participants from 11 postapproval studies of PrEP in cisgender women, the overall HIV incidence was 0.72 per 100 person-years. The HIV incidence rates per 100 person-years for different PrEP adherence trajectories were 0 for consistently daily (7 doses/week), 0.13 for consistently high (4-6 doses/week), 0.49 for high but declining (from a mean of 4-6 doses/week and then declining), and 1.27 for consistently low (less than 2 doses/week) adherence.

Meaning In a pooled analysis of postapproval studies, cisgender women with daily or consistently high adherence to emtricitabine and tenofovir disoproxil fumarate for PrEP experienced very low HIV incidence.
participants with plasma concentrations greater than or equal to 40 ng/mL were assigned to the high adherence group. This threshold was selected based on prior studies suggesting that 40 ng/mL reflects recent adherence32,33 and that participants with pill coverage of at least 80% had plasma tenofovir concentration greater than 40 ng/mL.31 Participants with concentrations less than 40 ng/mL were included in the low adherence group. If more than 1 type of adherence monitoring variable was available, a single measure was used based on the following priority: (1) tenofovir diphosphate concentration in DBS, (2) TFV concentration in plasma, (3) physician- or study-reported 5-level summary (very good, good, fair, poor, very poor), (4) physician- or study-reported 2-level summary (yes, no), (5) electronic adherence monitoring device, (6) self-report, and (7) pill count.

**Group-Based Trajectory Modeling**

Group-based trajectory modeling14 was used to identify patterns of PrEP adherence over 96 weeks. This approach has been applied to PrEP adherence studies previously.15-17 Data were available to support trajectory modeling in 2954 women. Linear, quadratic, or cubic models were used to allow the data to cluster into 2 to 6 groups of adherence trajectories based on ordinal adherence metrics. An ordinal number was assigned indicating overall adherence to denote the cutoffs in doses per week across all adherence measurements. For example, if adherence

<table>
<thead>
<tr>
<th>Source</th>
<th>Country</th>
<th>Study population</th>
<th>No. of participants</th>
<th>Measures used for adherence trajectory</th>
<th>No. of participants with adherence data (No. with objective measures*)</th>
<th>Age range, y</th>
<th>New HIV diagnosis, No.</th>
<th>Start date</th>
<th>End date</th>
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<tr>
<td>Partners Demo²¹</td>
<td>Kenya, Uganda</td>
<td>Serodiscordant couples</td>
<td>334</td>
<td>Tenofovir diphosphate concentration in DBS, plasma tenofovir concentration, and electronic monitoring</td>
<td>329 (88)</td>
<td>18-54</td>
<td>8</td>
<td>11/30/12</td>
<td>5/2/16</td>
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<td>TDF2 OLE²²</td>
<td>Botswana</td>
<td>Young heterosexual adults</td>
<td>102</td>
<td>Tenofovir diphosphate concentration in DBS</td>
<td>45 (45)</td>
<td>23-44</td>
<td>0</td>
<td>3/6/13</td>
<td>7/17/14</td>
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<td>PrePception²⁶</td>
<td>US</td>
<td>HIV-negative women and male partners living with HIV</td>
<td>24</td>
<td>Tenofovir diphosphate concentration in DBS</td>
<td>16 (16)</td>
<td>22-47</td>
<td>0</td>
<td>2/25/14</td>
<td>4/7/18</td>
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<td>Kolkata PrEP Demo²⁷</td>
<td>India</td>
<td>Female sex workers</td>
<td>678</td>
<td>5-Level physician-reported summary</td>
<td>678</td>
<td>19-50</td>
<td>0</td>
<td>1/18/15</td>
<td>9/21/19</td>
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<td>India</td>
<td>Female sex workers</td>
<td>647</td>
<td>Self-report</td>
<td>646</td>
<td>19-51</td>
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<td>12/15/18</td>
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<td>Electronic monitoring</td>
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<td>3/23/16</td>
<td>2/21/18</td>
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<td>Tenofovir diphosphate concentration in DBS</td>
<td>7 (7)</td>
<td>22-69</td>
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<td>5/13/16</td>
<td>8/30/17</td>
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<td>Young women (aged 15-24 y)</td>
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<td>Tenofovir diphosphate concentration in DBS and electronic monitoring</td>
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<td>Young women (aged 16-25 y)</td>
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<td>Tenofovir diphosphate concentration in DBS and 5-level physician-reported summary</td>
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<tr>
<td>Total</td>
<td></td>
<td></td>
<td>6296</td>
<td></td>
<td>2954 (237)</td>
<td>32</td>
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Abbreviations: CRUSH, Connecting Resources for Urban Sexual Health; MPYA, Monitoring PrEP Among Young Adult Women; POWER, Prevention Options for Women Evaluation Research.

* The number of participants with objective adherence measures (at least 1 measurement of tenofovir diphosphate concentration in dried blood spots [DBS] or tenofovir concentration in plasma) in the trajectory modeling (ie, 237 individuals had objective adherence measure data and 277 participants were assessed using exclusively subjective measurements in the trajectories). Adherence was categorized as consistently daily (7 doses/wk), high (4-6 doses/wk), moderate (2-3 doses/wk), and low (<2 doses/wk) as described in the Methods. The optimal (most stable) model from group-based trajectory modeling resulted in 4 groups with distinct patterns of adherence. Three groups had stable adherence over time, regardless of the model (3 or 4 groups) used: consistently daily (7 doses/wk), consistently high (4-6 doses/wk), and consistently low (<2 doses/wk). The fourth group had dynamic adherence over time, which went from initially high to declining (from 4-6 to 2-3 doses/wk).
wascategorizedas7dosesperweek,thearrangementforthatmeasurementwas4onascaleof1to4.Using
a specified number of trajectories and various shapes (eg, linear, quadratic), the model estimated the posterior probability
of an individual belonging to each group and the individual was classified based on the highest posterior probability. The final
models were selected and groups were defined based on bayesian or Akaike information criterion. Individual trajectories were
further examined using spaghetti plots.

Multinomial ordinal logistic regression models were used to calculate odds ratios (ORs) and associated 95% CIs in evaluating
the association between baseline participant characteristics and group-based adherence trajectory categories. Mixed
models with random effect by each individual study were used to account for clustering within the study. Incidence rate ratios (IRRs) were calculated to compare HIV incidence rates across different adherence trajectories, using the low adherence group as the reference.

All statistical analyses were performed using SAS software version 9.3 (SAS Institute) except for the trajectory group
modeling, which was conducted using the R version 3.00 with the gbmt package (R Foundation; https://cran.r-project.org/
web/packages/gbmt/index.html).

All studies were reviewed and approved by appropriate institutional review boards or similar authorities prior to implementa-
tion. Data used in this analysis were previously deidentified and members of the study team had no access to any
identifiable participant information.

Results

Participant Characteristics
Of the 6296 participants, 2886 (46%) were from Kenya, 1751 (28%) were from South Africa, 1325 (21%) were from India,
183 (2.9%) were from Uganda, 102 (1.6%) were from Botswana, and 49 (0.8%) were from the US. The mean (SD) age at PrEP
initiation across all 11 studies was 25 (7) years, with 61% of participants being younger than 25 years. The majority of participants were married (52%) and nulliparous (54%), and 21% reported engaging in transactional sex (eTable 1 in Supplement 1).

HIV Incidence
There were 32 incident HIV diagnoses among the 6296 participants, which is an overall HIV incidence rate of 0.72 per 100
person-years (95% CI, 0.51-1.01). HIV incidence was higher in younger (<25 years), married, and nulliparous participants (Figure 1). HIV incidence in other subgroups is shown in eTable 2 in Supplement 1.

Adherence Measures
Measures of adherence were available for 2955 participants and are reported by visit. The number of participants with objective adherence data declined from 147 at week 16 to 22 at week 96 (Figure 2A). Among those with objective adherence data, the percentage with high or daily adherence varied between 20% and 33%. Of the 2887 participants with subjective adherence data, a similar decline in the number of participants with reported adherence data was observed, with the percentage reporting high or daily adherence varying between 67% and 82% until week 80, then dropping to 13% at week 96 (Figure 2B). Subjectively reported adherence was substantially higher compared with objective measures at all points throughout participation, with the exception of the final visit.

Group-Based Adherence Trajectories
The various adherence measures used in modeling the adherence trajectory groups are summarized in eTable 3 in
The optimal (most stable) model from group-based trajectory modeling resulted in 4 groups with distinct patterns of adherence. Three groups had stable adherence over time, regardless of the model (3 or 4 groups) used: consistently daily (7 doses/week), consistently high (4-6 doses/week), and consistently low (less than 2 doses/week). The fourth group had dynamic adherence over time: initially high then declining (from a mean of 4-6 to 2-3 doses/week; Figure 3). Linear regression was applied to the adherence scores (including subjective and objective measurements) for the 1166 individuals who were clustered into the high but declining group. The linear regression trend is shown in Figure 3A. The regression trend reveals a linear decline, with “high” representing the Y-intercept of the regression line, signifying a group-based starting point with a mean of 4 to 6 doses per week. The negative slope of the regression line indicates the downward trend after this initial starting point. Therefore, the mean of 4 to 6 doses per week is not based on individual data or a specific period, rather it reflects a group-level overall starting point. Not all individuals initiated from the intercept and the declines did not occur within a specific or consistent timeframe.

Baseline participant characteristics associated with adherence trajectory categories are summarized in eTable 4 in Supplement 1. Participants younger than 25 years had lower odds of having consistently daily (OR, 0.42 [95% CI, 0.25-0.71], consistently high (OR, 0.82 [95% CI, 0.53-1.27]), or high but declining (OR, 0.51 [95% CI, 0.37-0.69]) (vs consistently low) adherence trajectory compared with women 25 years or older.
Adherence was categorized as consistently daily (7 doses/week), high (4-6 doses/week), moderate (2-3 doses/week), and low (less than 2 doses/week), as described in the Methods. The optimal (most stable) model from group-based trajectory modeling resulted in 4 groups with distinct patterns of adherence. Three groups had stable adherence over time, regardless of the model (3 or 4 groups) used: consistently daily (7 doses/week), consistently high (4-6 doses/week), and consistently low (less than 2 doses/week). The fourth group had dynamic adherence over time: initially high then declining (from 4-6 to 2-3 doses/week).

HIV Incidence Rates Among Women by Adherence Trajectory

A total of 12 incident HIV infections were observed among the 2955 participants with adherence data, compared with 20 diagnoses among 3341 participants without adherence data. None of the 498 participants with consistently high adherence by group-based trajectory acquired HIV (Figure 4); because there were no cases of infection, an IRR could not be calculated for this group. Only 1 of the 658 participants with consistently high adherence (4-6 doses/week) acquired HIV, resulting in an HIV incidence rate estimate of 0.13 per 100 person-years (95% CI, 0.02-0.92) for this group. The incidence rate was 0.49 per 100 person-years (95% CI, 0.22-1.08) in the high but declining adherence group (n = 1166) and 1.27 per 100 person-years (95% CI, 0.53-3.04) in the consistently low adherence group (n = 632). Individuals with consistently high adherence (IRR, 0.10 [95% CI, 0.01-0.88]) and individuals with high but declining adherence (IRR, 0.38 [95% CI, 0.12-1.26]) had lower HIV incidence compared with individuals with consistently low adherence.

Discussion

This pooled analysis of more than 6000 cisgender women using F/TDF for PrEP showed higher HIV incidence among younger, married, and nulliparous women, which is consistent with previous reports. Among the nearly 3000 individuals with adherence data, the effectiveness of F/TDF in preventing HIV infection was very high in women who demonstrated consistently high (4-6 doses/week) or consistently daily (7 doses/week) adherence. This interpretation is comparable to the established relationship between effectiveness and adherence observed in cisgender MSM. Even with the relatively low number of incident HIV diagnoses, a clear relationship was observed between evidence (both objective and subjective) of higher adherence with lower risk of HIV infection.

Current guidelines for PrEP use among cisgender women highlight the more rigid recommendations relative to those for MSM. In its 2021 PrEP Clinical Guideline Update, the US Centers for Disease Control and Prevention advised clinicians to counsel patients on the importance of daily dosing, noting that there is less “forgiveness” for missed PrEP doses for cisgender women than for MSM. These recommendations are largely based on pharmacokinetic modeling that suggests that cervicovaginal tissue concentrations of tenofovir disoproxil are lower than those observed in rectal tissue with similar dosing, although a correlation between tissue tenofovir disoproxil concentration and PrEP efficacy has not been definitively demonstrated. However, even with low overall adherence to oral F/TDF in some randomized clinical trials among cisgender women, a direct relationship between objective measures of adherence and efficacy was observed.

A 2023 analysis evaluating the F/TDF adherence-efﬁcacy relationship in the HPTN 083 and 084 trials demonstrated that, although there are differences in the pharmacologic forgiveness proﬁle between cisgender men and women, the protective efﬁcacy of F/TDF against HIV remained high (>80%) in cisgender women, despite less than daily dosing. Indeed, participants with tenofovir disoproxil concentrations indicative of taking 4 to 6 doses per week had a nearly 90% HIV risk reduction in the HPTN 084 trial, consistent with the results observed in the present study. Modeling studies leveraging F/TDF clinical trial data in cisgender women have demonstrated similar ﬁndings, underscoring how differences in efﬁcacy between cisgender men and women that were previously observed were likely driven by adherence rather than biological differences.

Women may be motivated to use oral F/TDF for reasons that relate directly to the timing of HIV risk or to perceived beneﬁt. For example, high uptake of F/TDF use was recently reported among HIV-exposed women with plans for pregnancy in rural Uganda. Sexual behaviors that expose women to HIV acquisition also change over time, as demonstrated in the HPTN 064 study, in which social factors were found to influence sustained high-risk behavior, and should be integrated into decisions about approach to PrEP use. These findings highlight how the broader context surrounding individuals’ HIV prevention needs should be integrated into decisions about PrEP use and that cisgender women need not be restricted to a rigid daily regimen, similar to the adherence forgiveness that has been reported in cisgender MSM.

Limitations

This analysis has several limitations. First, data were pooled from heterogeneous postapproval studies that differed in important ways, including (1) approaches to collecting the...
objective and subjective measures of adherence, (2) varied geographies with different HIV incidence, (3) distinct inclusion and exclusion criteria reflecting diverse behaviors associated with HIV acquisition among the participants, (4) approaches to mode of PrEP delivery (eg, home PrEP delivery, use of mobile clinic), and (5) differences in sociodemographics related to country setting. Second, adherence data, especially from objective measures, were only available for a subset of participants, limiting generalizability. The use of plasma tenofovir concentrations to categorize participants into adherence categories may be susceptible to misclassification because plasma tenofovir has a short window of detection. However, no changes were observed in sensitivity analyses excluding the 46 participants for whom plasma tenofovir measures were used in group-based trajectory modeling (data not shown). Importantly, objective adherence data were available only for a minority of participants and were largely representative of the total population. As previously reported, it was also observed that patients’ self-reported adherence was substantially overestimated compared with objective measures.\textsuperscript{4,5,40} This reinforces the notion that considerable caution should be applied when interpreting self-reported adherence to PrEP. Third, the accuracy of the group-based trajectory methodology used could be improved with a greater sample size and longer duration of follow-up.

**Conclusion**

This analysis of the effectiveness of F/TDF for PrEP in cisgender women in real-world settings supports that HIV prevention effectiveness is high with adherence to greater than 4 doses per week, which is comparable to the adherence-efficacy relationship for PrEP use in cisgender MSM. This finding may inform provision of F/TDF for PrEP to, and counseling for, cisgender women, emphasizing that although daily adherence is optimal, a minimum of 4 doses per week of F/TDF is expected to provide effective protection for most females.


