

HIV PrEP for Adolescents and Young Adults

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Module 2: [HIV PrEP In-Depth Topics](#)

Lesson 1: [HIV PrEP for Adolescents and Young Adults](#)

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Background and HIV PrEP Coverage in Youth

Background

Adolescence and young adulthood are stages of life that often encompass sexual debut, sexual exploration, and sexual experimentation.[1] When these periods intersect with activities such as substance use or decreased condom use, youth become increasingly prone to HIV acquisition.[1] In 2021, individuals aged 13 to 24 years of age accounted for roughly 1 in 5 persons who newly acquired HIV in the United States.[2] Sexual, gender and racial/ethnic minority youth, particularly those living in the Southern United States, are even more vulnerable to acquiring HIV. In this context, HIV preexposure prophylaxis (PrEP) is a known, effective, data-proven biomedical tool for HIV prevention.[3] In 2012, tenofovir DF-emtricitabine (TDF-FTC) was approved by the US Food and Drug Administration (FDA) for HIV PrEP in adults, but there was little initial uptake among young adults. In 2018, the FDA changed the criteria for HIV PrEP in youth to be based on weight, with approval for those weighing at least 35 kg (77 lb).[3]

HIV PrEP Coverage for Youth

In the United States, HIV PrEP has been underused as an HIV prevention tool in adolescents and young adults (Figure 1).[4] In 2022, only about 24% of youth 16-24 years of age who had an indication for PrEP actually received HIV PrEP (the percentage of people receiving HIV PrEP versus those with an HIV PrEP indication is often referred to as “PrEP coverage”).[4] Several studies have demonstrated barriers in the HIV PrEP care cascade for adolescents, including lack of HIV PrEP awareness among adolescents or their guardians, reluctance to use HIV PrEP, and reduced HIV PrEP prescriptions written by clinicians for adolescents who have a significant risk of acquiring HIV.[5,6] These studies underscore the importance of a multipronged approach to addressing deficits in the HIV PrEP care cascade—from awareness to prescribing to uptake to adherence, in order to successfully harness HIV PrEP to advance the goal of ending the HIV epidemic in the United States. Further, several issues continue to exist related to health professionals that contribute to low HIV PrEP prescribing for youth, including lack of knowledge, hesitancy to prescribe, and inability to effectively conduct sexual history discussions with youth.

Epidemiology of STIs and HIV in Youth

Youth Sexual Health Behaviors

The Youth Risk Behavior Surveillance System (YRBSS) is a population-based data tool used by the CDC to monitor different categories of health behaviors among students in grades 9 through 12 in the United States.[7,8] The activities evaluated include substance use, sexual health practices, dietary and exercise habits, and violent or injury-causing activities. Sexual risk behavior data from the 2021 YRBSS for high school students indicates that 29% of heterosexual students and 33% of lesbian, gay, and bisexual, and queer+ (LGBQ+) students indicated they had ever had sexual intercourse.[8] Among those students who reported being sexually active, only 52% reported using a condom the last time they had sex.[8] In this survey, only 6% of students had ever been tested for HIV; the HIV testing rate varied from 5% for heterosexual students to 7% for LGBQT students.[7]

STIs in Adolescents and Young Adults

The Centers for Disease Control and Prevention (CDC) annual surveillance data on sexually transmitted infections (STIs) includes data for adolescents and young adults.[9] For this surveillance report, the age range used to define adolescents and young adults is 15-24 years of age.[9] In recent years, STI rates have been very high among young people 15 to 24 years of age, with this age group accounting for 53% of newly reported cases of STIs for the year 2020 (Figure 2).[9] The STI rates in youth are notably higher in certain racial/ethnic minoritized groups, especially Black individuals. These disparities do not reflect sexual practices but rather other structural differences and social determinants, such as inequities in access to screening and treatment for STIs. In addition, it is well established that some STIs facilitate HIV acquisition and transmission through a variety of biological and nonbiological variables.[10,11,12]

HIV in Adolescents and Young Adults

According to the most recent CDC HIV surveillance data, 6,927 persons 13 to 24 years of age were newly diagnosed with HIV in the United States in 2021.[13] Overall, this age group accounted for 19% of all persons newly diagnosed with HIV diagnoses during 2021.[13] For persons 13 to 24 years of age who were newly diagnosed with HIV in 2021, most (79%) were young adults (20 to 24 years of age).[13] Among all persons 13 to 24 years of age newly diagnosed with HIV who reported gender, 84% were cisgender men, 12% were cisgender women, 4% were transgender women, and Figure 3)

Evidence for HIV PrEP in Youth

ATN 110

Adolescent Trials Network 110 (ATN 110) was an open-label study that examined adherence with oral daily HIV PrEP in young (18 to 22 years of age) men who have sex with men (MSM) ([Figure 4](#)).[\[14\]](#) Investigators enrolled 200 young men who reported risk for HIV acquisition in the prior 6 months. The study was conducted in 12 United States cities between March and September 2013.[\[14\]](#) All participants were prescribed daily oral TDF-FTC for 48 weeks.[\[14\]](#) The rates of STIs were high at baseline (22% of participants) and remained high throughout the study. There were four individuals who had HIV seroconversion during the study (3.29 per 100 person-years).[\[14\]](#) Investigators assessed medication adherence by determining tenofovir diphosphate levels in dried blood spots and found adherence declined as the study progressed, with a significant drop-off occurring at week 24.[\[14\]](#) Study participants reported different reasons for nonadherence, including being away from home, forgetting to take medication, and being busy with other activities.[\[14\]](#) In addition, racial disparities in adherence were also noted: Black and/or African American youth had lower adherence rates, which highlights the need for culturally tailored HIV PrEP education, delivery, and implementation.

ATN 113 (Project PrEPare)

The ATN 113 (Project PrEPare) study was conducted to assess the safety and adherence of TDF-FTC as HIV PrEP for adolescent MSM who were 15-17 years of age ([Figure 5](#)).[\[15\]](#) Further, the investigators also looked at sexual practices among the study participants. This was the first major adolescent-specific HIV PrEP demonstration project based in the United States, and it showed that daily oral TDF-FTC was well tolerated and safe when taken by adolescents, but adherence waned significantly during the study.[\[15\]](#) Although 95% of study subjects had detectable drug levels during the first 12 weeks of the study, only 17 (22%) had levels associated with taking four or more pills per week (by week 48 of the study).[\[15\]](#) Those individuals who did not have protective serum levels of tenofovir reported the following reasons for nonadherence: being too busy, forgetfulness, being away from home, and changes to their routine. Among the 78 enrolled persons, the HIV incidence rate was high (6.4% per 100 person-years), and the three participants who had HIV seroconversion had very low levels of tenofovir-diphosphate, as measured in dry blood spots, measured at the time of seroconversion. This study highlights the importance of addressing medication adherence and the challenges that are commonplace among youth who are taking HIV PrEP.[\[15\]](#)

HPTN083

The HIV Prevention Trials Network (HPTN) 083 study was a phase 2b/3, randomized clinical trial comparing the safety and efficacy of long-acting injectable cabotegravir (CAB-LA), administered every 8 weeks, to once-daily oral TDF-FTC as HIV PrEP, in cisgender men who have sex with men and transgender women who have sex with men.[\[16\]](#) The study enrolled approximately 4,500 participants, aged 18 years and older. Notably, 68% of the study participants were 29 years of age or younger.[\[16\]](#) By study week 153, there were 52 incident HIV infections, with 39 infections noted in the TDF-FTC arm versus 13 in the CAB-LA arm—representing a 67% risk reduction for acquiring HIV with CAB-LA compared to TDF-FTC.[\[16\]](#) This publication did not provide data on infections based on age group.[\[16\]](#)

HPTN084

The HPTN 084 study was a phase 3, double-blind, randomized clinical trial comparing the efficacy of CAB-LA as HIV PrEP to once-daily oral TDF-FTC in cisgender women.[\[17\]](#) A total of 3,224 cisgender women from sub-Saharan Africa, aged 18 years and older, were enrolled. Although all study participants were adults, approximately 50% were younger than 25 years of age.[\[17\]](#) For the 1,717 participants enrolled who were younger than 25 years of age, there were 20 incident HIV seroconversions in the TDF-FTC arm compared to three in the CAB-LA arm, representing an 85% risk reduction for HIV acquisition with CAB-LA.[\[17\]](#) This trial did not provide a specific, further age breakdown of participants younger than 25 years of age, but presumably,

this included a substantial number of young adults 18-21 years of age.[17]

Plus-Pills

Plus-Pills was an open-label, single-arm, phase 2 study conducted in South Africa that enrolled sexually active male and female adolescents, 15 to 19 years of age, to evaluate the safety and feasibility of daily oral TDF-FTC as HIV PrEP.[18] Similar to ATN 113, Plus-Pills validated that oral TDF-FTC was safe and tolerable in an adolescent population while also demonstrating medication adherence problems among youth.[18]

Partners PrEP Trial

The PARTNERS PrEP Trial enrolled HIV-serodifferent heterosexual couples from Kenya and Uganda and compared three oral daily HIV PrEP options to be taken by the partner without HIV: TDF alone, TDF-FTC, or placebo.[19] Although this trial was not designed specifically as an adolescent HIV PrEP study, among the 4,747 enrolled couples who were eligible, 533 of the couples had a seronegative partner younger than 25 years of age.[19] Overall, the study showed a reduced risk of HIV acquisition for HIV-seronegative participants (women and men) who took either TDF or TDF-FTC compared with those individuals who took placebo.[19] For the subgroup of persons younger than 25 years of age who were enrolled in the study, beneficial protection was observed in persons who received HIV PrEP when compared with those who received placebo: the HIV acquisition rates per 100 person-years were 1.07 in the TDF group, 2.34 in the TDF-FTC group, and 4.04 in the placebo group.[19]

IPREX Trial

The Preexposure Initiative (iPrEx) study was a phase 3, randomized, double-blind, placebo-controlled international trial that evaluated daily oral TDF-FTC as HIV PrEP in HIV-seronegative adult MSM and transgender women who have sex with men.[20] Among the 2,499 individuals who enrolled, 1,253 (50.1%) were 18 to 24 years of age.[20] In the subanalysis for those younger than 25 years of age, new HIV infections were diagnosed in 22 of 591 (3.7%) who were in the TDF-FTC group compared with 37 of 662 (5.6%) in the placebo group; this was a 34% reduction in HIV infections.[20]

Clinic Visits and Obtaining a Sexual History in Adolescents

Approach to Clinical Visit for Prevention Services

In order to provide an optimal clinical plan for adolescents and young adults, medical providers must be sensitive and flexible in their approach during this encounter. It is imperative that the provider forms a therapeutic partnership with the adolescent or young adult during the visit. To this end, every effort must be taken to ensure that the adolescent or young adult can talk alone with the clinician in the absence of parents or guardians. Typically, the American Association of Pediatrics recommends initiating one-on-one adolescent patient-provider visits at the age of 11. By collaborating with parents/guardians to establish these individual sessions as a part of routine health care, the clinical provider can then continue such visits into the patient's adolescent years. During those individualized clinic periods, the provider should build rapport and trust by being forthright in informing the adolescent of the patient-provider confidentiality and the instances when confidentiality may be breached.[\[21\]](#) A commonly used psychosocial framework for obtaining a clinical history is HEEADSSS: home, education and/or employment, eating, activities, drugs, sexuality, suicide and/or depression, and safety.

Obtaining a Sexual History

Eliciting a comprehensive history, including social and sexual history, should take place in a non-threatening environment, using non-judgmental, open-ended questions and without assumptions about the youth's sexual orientation or gender identity. In general, when obtaining a sexual history, the CDC recommends using the five "Ps" as a reminder: partners, practices, protection from STIs, past history of STIs, and prevention of pregnancy.[\[22\]](#) The "Taking a Sexual History" tool from the Sexual Information and Education Council of the United States (SIECUS) provides detailed guidance on how to take an inclusive sexual history in all youth, including lesbian, gay, bisexual, transgender, queer, and questioning (LGBTQ) youth.[\[23\]](#) When engaging in these discussions, it is extremely important for the provider to understand and contextualize other activities and circumstances that impact sexual activity, such as substance use, sexual or emotional abuse, peer pressure and bullying, mental illness, food insecurity, unstable housing, interactions with the juvenile justice system or transactional sexual encounters.

Assessing for HIV PrEP in Youth

The following summarizes indications for prescribing HIV PrEP to adolescents or young adults. These recommendations are based on the 2021 CDC HIV PrEP Guidelines.[\[3\]](#)

Sexually Active Adolescents and Young Adults

Adolescents and young adults who weigh at least 35 kg (77 lb) AND who have had anal or vaginal sex in the past 6 months may be eligible for HIV PrEP if they have any of the following ([Figure 6](#)):

- An HIV-seropositive sex partner (especially if a sex partner has an unknown or detectable HIV RNA level)
- One or more bacterial STIs in the prior 6 months: gonorrhea, chlamydia, and syphilis for MSM and transgender women who have sex with men, including those who inject drugs; gonorrhea and syphilis for heterosexual women and men, including persons who inject drugs
- History of inconsistent or no condom use with sex partner(s)

Adolescents and Young Adults who Inject Drugs

Adolescents and young adults, weighing at least 35 kg (77 lb) and who inject drugs or have injected in the past 6 months, should also be assessed for their sexual risk of HIV. These individuals should be prescribed PrEP if they have any of the following ([Figure 7](#)):

- Have an HIV-seropositive injecting partner
- Share injection equipment
- Engage in sexual activity that increases risk for acquiring HIV

US Preventive Services Task Force

The 2023 United States Preventive Services Task Force (USPSTF) has given an “A” rating for HIV PrEP for adolescents and adults who are at high risk of HIV acquisition and who weigh at least 35 kg (77 lb).[\[24\]](#)

Society for Adolescent Health and Medicine

The Society for Adolescent Health and Medicine (SAHM) issued a position paper in 2018 that provides context and guidance for pediatricians and adolescent medicine health care professionals to promote and enhance the use of HIV PrEP for youth to decrease the risk of acquiring HIV.[\[25\]](#) The SAHM position states that “adolescent and young adult health professionals should develop evidence-based, developmentally appropriate, culturally sensitive, and accessible PrEP service delivery models as part of routine care offered to adolescents and young adults.” In an effort to achieve this goal, medical providers are encouraged to counsel patients about PrEP, describe it as a safe and effective HIV prevention strategy in combination with other prevention strategies, and refer patients to available health care providers for initiation and monitoring of HIV PrEP.[\[25\]](#)

HIV PrEP Medications for Youth

FDA-Approved Medications for HIV PrEP

The FDA has approved three medications that can be used for HIV PrEP in young adults who weigh at least 35 kg (77 lb): daily oral tenofovir DF-emtricitabine (TDF-FTC), daily oral tenofovir alafenamide-emtricitabine (TAF-FTC), and long-acting injectable cabotegravir (CAB-LA). Note that TAF-FTC is not FDA-approved for cisgender women to prevent vaginal acquisition of HIV, regardless of age. Further, although no medication has been FDA-approved to prevent HIV acquisition associated with injection drug use, as noted below, the CDC has recommended TDF-FTC for HIV PrEP in people who inject drugs, based on published data in the Bangkok Tenofovir Trial.[26] The following summarizes potential medication options and CDC recommendations for HIV PrEP in adolescents and young adults.[3]

Centers for Disease Control and Prevention Recommendations

Regardless of gender, TDF-FTC and CAB-LA are recommended options for adolescents and young adults who weigh at least 35 kg (77 lb) to reduce the risk of acquiring HIV through sexual activity. Adolescents and young adults who weigh at least 35 kg (77 lb) can also use TAF-FTC to reduce the risk of HIV infection from sexual acquisition, but not individuals at risk of acquiring HIV through receptive vaginal sex.[3] For adolescents and young adults, TDF-FTC is the only medication recommended for preventing the acquisition of HIV via injection drug use.[3] The following summarizes potential medication options and CDC recommendations for HIV PrEP in adolescents and young adults.[3]

- **Tenofovir DF-Emtricitabine (TDF-FTC):** Daily oral TDF-FTC is recommended as an option for the prevention of sexual acquisition of HIV in adolescents and adults who weigh at least 35 kg (77 lb). In addition, daily oral TDF-FTC is the only medication recommended for use as HIV PrEP to prevent HIV acquisition in persons who inject drugs who weigh at least 35 kg (77 lb). The recommended dosing for oral TDF-FTC is one tablet daily. TDF-FTC should not be used in persons with a creatinine clearance of less than 60 mL/min. TDF-FTC is considered safe for use in pregnancy.
- **Tenofovir alafenamide-Emtricitabine (TAF-FTC):** Daily oral TAF-FTC is recommended as an option for prevention of sexual acquisition of HIV in adolescents and adult males and transgender females who weigh at least 35 kg. Note that TAF-FTC is not FDA-approved for cisgender women to prevent vaginal acquisition of HIV, regardless of age. In addition, this medication should not be prescribed if the estimated creatinine clearance is less than 30 mL/min. TAF-FTC is considered safe for use in pregnancy, but since this medication is not recommended for cisgender women to prevent vaginal acquisition of HIV, it should not be used during pregnancy.
- **Long-Acting Injectable Cabotegravir (CAB-LA):** CAB-LA is recommended for HIV PrEP to prevent the sexual acquisition of HIV in adolescents and young adults, regardless of gender, if they weigh at least 35 kg (77 lb). There are no restrictions on prescribing CAB-LA based on renal function. At this time, CAB-LA is not recommended for use in pregnancy due to lack of data. For detailed information about CAB-LA, including dosing for the optional oral lead-in, see the [Cabotegravir Guide](#) on this website.
- **On-Demand (2:1:1) HIV PrEP:** Oral TDF-FTC is the only medication that has been studied for the use of on-demand (2:1:1) HIV PrEP and this has been studied only for use in MSM who are 18 years of age and older. Based on available data, the CDC lists on-demand HIV PrEP with TDF-FTC as an alternative for HIV PrEP in young adult MSM who are 18 years of age and older. In addition, the use of on-demand HIV PrEP has not been studied in persons younger than 18 years of age, and therefore it is not recommended for this age group. Note that the on-demand dosing strategy has not been approved by the FDA for people of any age. For more detailed information, see the [On-Demand Dosing Guide](#) on this website.

Tolerability and Safety Data

For adolescents younger than 18 years of age, several studies suggest TDF-FTC is safe when used as HIV PrEP in adolescents and young adults, but there are no published safety data with TAF-FTC when used as HIV PrEP in this age group.[[14,15,18](#)] Substantial long-term HIV antiretroviral treatment experience with TDF-FTC and TAF-FTC—as components of antiretroviral therapy for the treatment of persons with HIV—has shown excellent tolerance and safety profiles.[[21,27,28](#)] There are no published data with CAB-LA when used for HIV PrEP in adolescents, but preliminary safety data from the MOCHA trial, which involved treatment of children with HIV using cabotegravir-rilpivirine as antiretroviral therapy, support use of both of these medications in children 12 years of age and older. The following summarizes major tolerability and safety concerns for adolescents and young adults.

- **Start-Up Syndrome:** Safety data in adults has shown that about 10% of persons starting oral TDF-FTC for HIV PrEP experience a “start-up syndrome” that consists of mild to moderate gastrointestinal symptoms, fatigue, and headache.[[3,29](#)] Less often, some will develop unintentional transient weight loss that may last for up to 24 weeks.[[3,29](#)]
- **Bone Mineral Density Changes:** Safety data from the ATN 110 study showed a modest decrease in bone mineral density of adolescents receiving TDF-FTC.[[14](#)] The clinical significance of these changes is unknown, and available data suggest these changes are reversible among persons younger than 25 years of age.[[14,30](#)] With regard to effects on the skeletal system, no medication-attributable fractures were reported in the ATN 113 study, but the increase in bone mineral density that is expected during this late adolescent age period was slightly blunted.[[15](#)] More research is necessary to ascertain the long-term clinical impact of TDF on bone density in adolescents who are receiving this medication for HIV PrEP, especially for those who take TDF for longer than 1 year.[[18](#)]
- **Nephrotoxicity:** The ATN 110 and 113 studies of TDF-FTC in adolescents demonstrated negligible renal adverse effects.[[14,15](#)]
- **Injection Site Reactions:** The major adverse effect of CAB-LA has been injection site reactions, which have resulted in the discontinuation of CAB-LA in about 2% of persons receiving this medication.[[16](#)] The most common symptoms associated with injection site reactions are pain and tenderness; if these symptoms occur, they typically begin about 1 day after the injection and last about 3 days.[[16](#)] There are no data on injection site reactions specific to adolescents or young adults with CAB-LA for HIV PrEP, but the rate and severity would be anticipated to be similar to that seen in adults.

Baseline Laboratory Evaluation

Baseline Evaluation

Adolescents and young adults who are interested in and willing to take HIV PrEP should undergo a baseline evaluation to obtain screening laboratory tests and to counsel the individual on taking HIV PrEP. As part of the baseline evaluation, it is essential to confirm the individual considering HIV PrEP (1) weighs at least 35 kg (77 lb) and (2) does not already have HIV. In addition, as part of the baseline evaluation for cisgender women (and all other individuals with a uterus), it is important to evaluate pregnancy status unless it is already known that the individual is pregnant. The routinely recommended baseline laboratory studies in the 2021 CDC HIV PrEP Guidelines are the same for youth as for older adults.[3] These baseline laboratory studies are summarized and discussed in detail in the *HIV PrEP Fundamentals* module lesson on [Baseline Evaluation and Starting HIV PrEP](#). In addition, summary tables for initial laboratory studies are available in the [Laboratory Monitoring Guide](#) on this website.

Medication Adherence

Reduced Adherence

In order for HIV PrEP to be effective in preventing the acquisition of HIV, adequate medication adherence is essential. For adolescents and young adults, medication adherence is often challenging and typically more problematic than with adults ([Figure 8](#)).[\[14,15,18,31\]](#) Problems with adherence in younger persons taking HIV PrEP may result from a variety of factors, including susceptibility to peer influence, an underdeveloped cognitive ability for organization and long-term planning, socioeconomic or familial stressors, mental illness, or substance use issues.[\[21,29\]](#) For many HIV PrEP studies, investigators have estimated adherence based on tenofovir diphosphate levels in dried blood spots. Most studies have used a tenofovir diphosphate level of at least 700 fmol/punch to be consistent with taking at least four pills of TDF-FTC per week, which is the level most experts believe is required to provide adequate protection for HIV acquisition when taking daily oral TDF-FTC.[\[32,33\]](#)

- **ATN 110:** In the open-label ATN 110 study, young MSM 18 to 22 years of age were offered oral daily TDF-FTC for HIV PrEP. Investigators estimated adherence by determining tenofovir diphosphate levels in dried blood spots.[\[14\]](#) The estimated adequate adherence was 56% at week 4 and declined to 34% by week 48.[\[14\]](#) The most common reasons reported for missing pill doses were forgetting, being away from home, and too busy with other things.[\[14\]](#)
- **ATN 113:** In this open-label study, young MSM 15-17 years of age were offered oral daily TDF-FTC for HIV PrEP.[\[15\]](#) Similar to ATN 110, adherence was estimated by determining tenofovir diphosphate levels in dried blood spots.[\[15\]](#) The estimated adequate adherence was 54% at week 4 and declined to only 22% at week 48.[\[15\]](#) The most common reported reasons for missing doses were being away from home, being too busy, and forgetting.
- **Plus Pills:** In this open-label study conducted in South Africa, young females and males 15-19 years of age who were at risk of acquiring HIV were offered HIV PrEP with daily oral TDF-FTC.[\[18\]](#) Two-thirds of the participants were female.[\[18\]](#) Similar to ATN 110 and ATN 113, adherence was estimated by determining tenofovir diphosphate levels in dried blood spots.[\[18\]](#) The estimated adequate adherence was 55% at week 4 and declined to 16% at week 48.[\[18\]](#) Major reasons identified for adherence problems were forgetting, tablet size, and decreased perception of risk.[\[18\]](#)
- **RADAR Cohort:** In a longitudinal cohort study involving young MSM 16 to 29 years of age living in the Chicago metropolitan area who reported use of HIV PrEP in the prior 6-month period, investigators evaluated HIV PrEP discontinuation.[\[34\]](#) Overall, for these participants who used HIV PrEP in the prior 6 months, 65 of 197 (33%) discontinued taking HIV PrEP, and 79% of those who discontinued HIV PrEP did not notify their medical provider prior to stopping HIV PrEP.[\[34\]](#) The main reasons identified for HIV PrEP discontinuation included trouble getting to medical appointments, issues related to insurance coverage or loss, and the belief they were not at risk of acquiring HIV anymore.[\[34\]](#)

Strategies for Improving Adherence

Some adolescent HIV PrEP clinical trials demonstrated that having monthly clinic visits early on during HIV PrEP start-up resulted in improved medication adherence.[\[14,15\]](#) This suggests that frequent interactions between adolescents and their medical providers may enhance HIV PrEP adherence. These visits should be used as an opportunity to routinely screen for potential barriers to adherence, as well as provide advice, support, and/or resources to overcome these obstacles in order to retain these individuals within the HIV PrEP continuum of care. The use of technology-focused interventions and other strategies to increase medication adherence is an evolving field. In addition, mobile technologies have been used successfully to improve antiretroviral regimen adherence in youth with HIV, and these could be leveraged for the HIV PrEP arena as well.[\[35,36,37,38,39,40,41\]](#)

Follow-Up and Laboratory Monitoring

Clinical Follow-Up for Persons Receiving HIV PrEP

For adolescents and young adults who start on oral HIV PrEP, clinical follow-up should occur every 3 months.[3] At each clinical follow-up visit, the clinician should ask about medication adherence and tolerance, as well as any changes in routines and schedules. With adolescents, it is particularly important to include these discussions regarding adherence at every clinic visit. For those receiving CAB-LA for HIV PrEP, the first clinical visit should occur 1 month after the first initiation injection and then every 2 months thereafter in conjunction with the maintenance CAB-LA injections.[3] Regardless of what type of medication is used for HIV PrEP, each clinical follow-up visit should include a discussion of the need for ongoing HIV PrEP, a review of any symptoms that would suggest acute HIV infection, and an assessment of any symptoms that suggest a current STI.[3] With each clinical visit, the clinician should provide an opportunity for the adolescent/young adult taking HIV PrEP to ask any questions or raise any concerns they have. Any individual who has symptoms consistent with an STI should undergo prompt STI testing and receive appropriate treatment as clinically indicated; this evaluation should occur regardless of when the next routine STI screening is due.

Laboratory Monitoring While on PrEP

The recommended follow-up laboratory studies in the 2021 CDC HIV PrEP Guidelines are the same for youth as for older adults.[3] These monitoring laboratory studies are summarized and discussed in detail in the *HIV PrEP Fundamentals* module lesson on [Follow-Up Care and Monitoring on HIV PrEP](#). In addition, summary tables for initial laboratory studies are available in the [Laboratory Monitoring Guide](#) on this website.

Legal and Insurance Issues

Medical providers who care for adolescents, especially for sexual health services, must be well informed regarding legal issues, especially pertaining to consent for clinical care and confidentiality. A CDC review of the statutes and regulations related to minors' informed consent to receive STI and HIV services concluded that as of 2020, all states and the District of Columbia had explicit laws permitting minors of a specific minimum age to provide informed consent to access STI diagnosis and treatment.[42] The age threshold is uniquely defined by each state or jurisdiction. Further, nuances exist in the interpretation and implementation of these laws with regard to HIV services, including HIV PrEP. As an example, in certain areas, HIV services could be included under general STI services, since the law may not explicitly distinguish between HIV and other STIs.[42] In other cases, certain statutes or regulations may allow for HIV prevention (hence HIV PrEP) to be considered under general preventive health care. The key point is that no state or jurisdiction in the United States bans a minor from providing their autonomous informed consent to receive HIV PrEP.[21,42] It is also important to note that consenting to HIV PrEP does not guarantee the confidentiality of care and limitations to confidentiality exist and vary by state (e.g., billing documentation from commercial health insurance plans, mandatory child abuse reporting, or neglect reporting laws). Therefore, an adolescent care provider must scrutinize the HIV PrEP prescription landscape in the context of local laws on autonomous health care decision-making by minors. Ideally, medical providers can serve as key sexual health advocates for their young patients by forging a constructive, therapeutic partnership with parents or guardians while also balancing the need for adolescent autonomy to the extent that is permissible by law.[21]

The following sites provide information related to HIV and STI consent and minors:

- Centers for Disease Control and Prevention (CDC): [Minor's Consent Law](#)
- Guttmacher Institute: [Minor's Access to STI Services](#)
- National Alliance of State and Territorial AIDS Directors (NASTAD): [Minor Consent and Confidentiality Laws for PrEP and HIV Treatment](#)

Insurance Issues Pertaining to Prescribing PrEP to Adolescents

Barring feasibility due to logistical or patient safety reasons (e.g., concerns for abuse), an adolescent health provider must make every effort to engage both adolescents and their parent(s) in mutual discussion, prior to HIV PrEP initiation. Ideally, the provider should first obtain consent from the adolescent to engage parent(s) followed by parental consent. The latter is important because parents/guardians could be made aware of such information from other sources, such as routine health plan communication or insurance plan documentation (e.g., explanation of benefits letter). Uninsured minors may be eligible for medication access support for HIV PrEP, but this may also need parental/guardian consent. If circumstances warrant absolute confidentiality for the minor, then adolescent health providers may need to pursue creative strategies, such as appealing to medication access programs.

Summary Points

- Approximately 20% of new HIV diagnoses in the United States in 2021 were among individuals 13 to 24 years of age.
- HIV PrEP is an important and underutilized tool for HIV prevention among adolescents and young adults who are at high risk for HIV acquisition.
- HIV PrEP is indicated for adolescents and young adults who have sexual risk factors or injection drug use risk factors that make them more susceptible to HIV acquisition.
- Three different HIV PrEP regimens are approved for adolescents and young adults weighing at least 35 kg (77 lb). They are TDF-FTC, TAF-FTC, and CAB-LA.
- TDF-FTC is not indicated for HIV PrEP in cisgender women (of any age) to prevent HIV acquisition via receptive vaginal sex.
- Risk assessment and baseline laboratory evaluation are required prior to prescribing HIV PrEP, including documentation that the person to receive HIV PrEP has a negative baseline HIV test.
- On-demand (2-1-1) dosing of TDF-FTC as HIV PrEP is only recommended as an option for MSM adolescents and young adults older than 18 years of age.
- Adherence challenges exist with youth taking oral HIV PrEP and these should be discussed on a regular basis at clinic visits and efforts made to address any adherence issues.
- The need for parental consent to prescribe HIV PrEP to adolescents varies from state to state. In general, there is no state or jurisdiction in the United States that bans a minor from providing their autonomous informed consent to receive HIV PrEP.
- Ideally, providers must make an effort to obtain parental or guardian consent prior to prescribing HIV PrEP to a minor, but the inability to do this should not preclude the adolescent's access to HIV PrEP.

Citations

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Figures

Figure 1 HIV PrEP Coverage for Youth

Source: Centers for Disease Control and Prevention. Core indicators for monitoring the Ending the HIV Epidemic initiative (preliminary data): National HIV Surveillance System data reported through September 2022; and preexposure prophylaxis (PrEP) data reported through June 2023. HIV Surveillance Data Tables 2023;4(No. 4). Published December 2023.

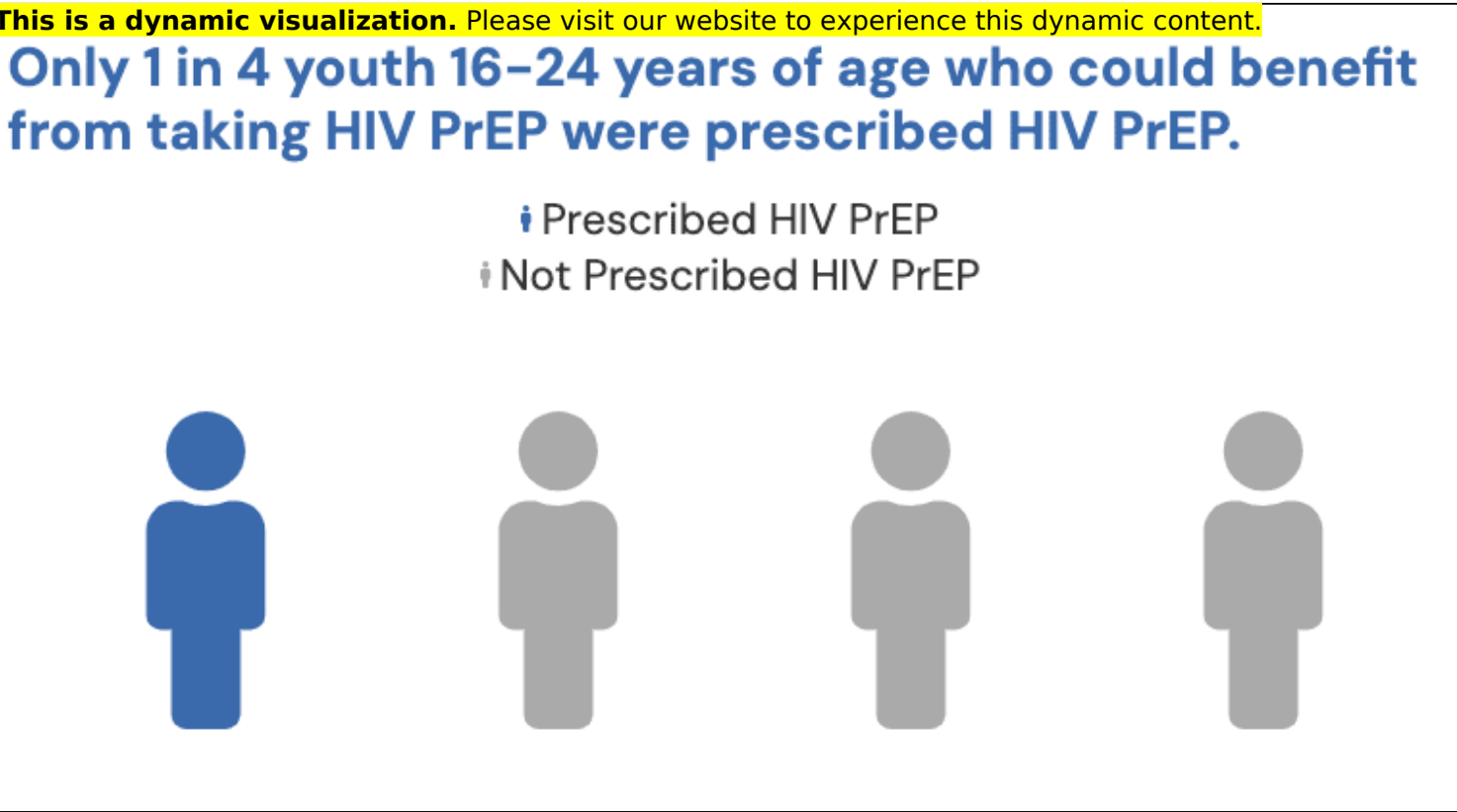


Figure 2 New Diagnoses of STIs Among Adolescents and Young Adults in the United States, 2021

Source: Centers for Disease Control and Prevention. Sexually Transmitted Disease Surveillance 2021. Atlanta: U.S. Department of Health and Human Services; April, 2023.

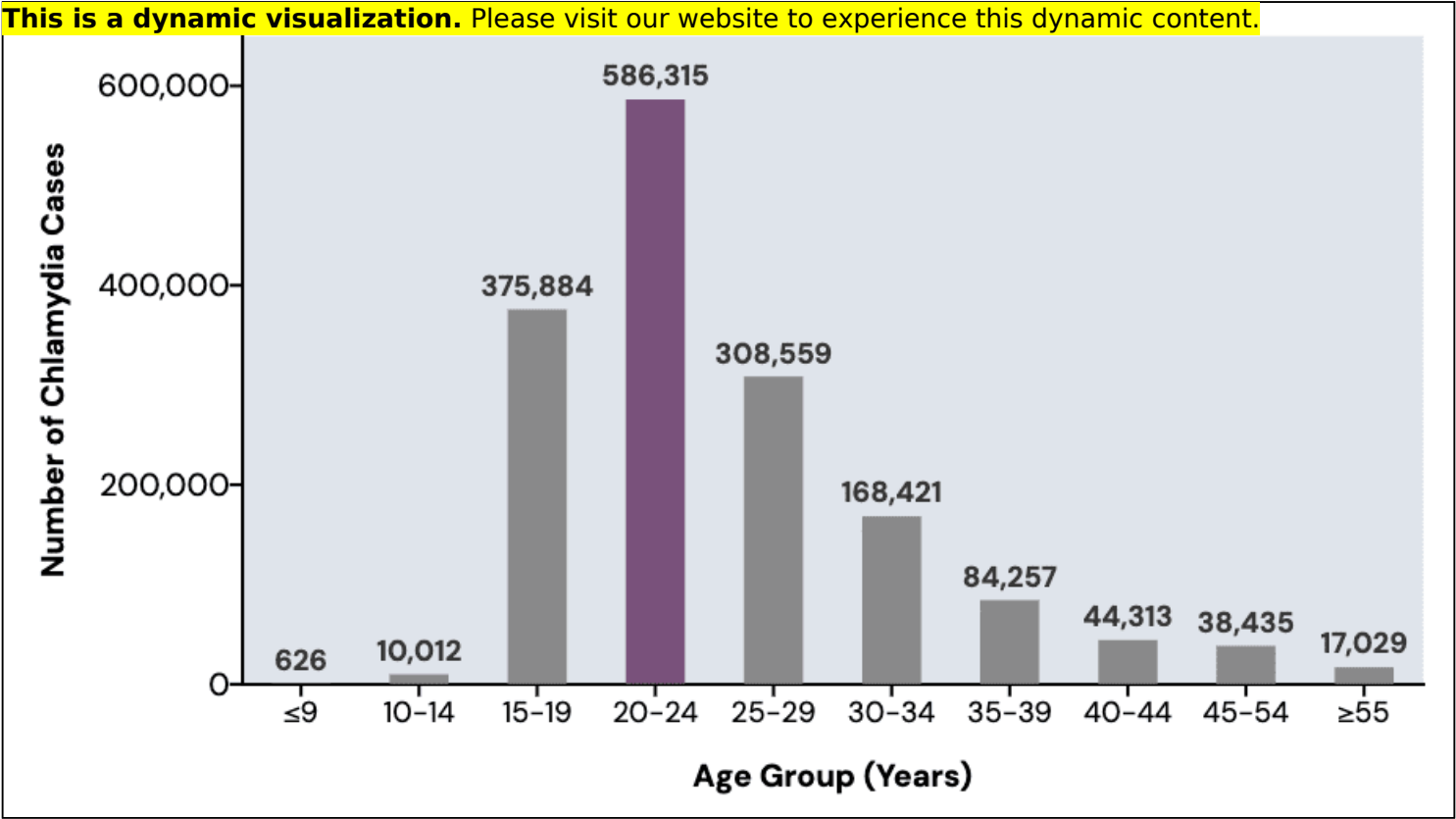


Figure 3 New HIV Diagnoses among Adolescents in the United States

Source: Centers for Disease Control and Prevention. Diagnoses of HIV infection in the United States and dependent areas, 2021. HIV Surveillance Report, 2021; vol. 34. Published May 2023.

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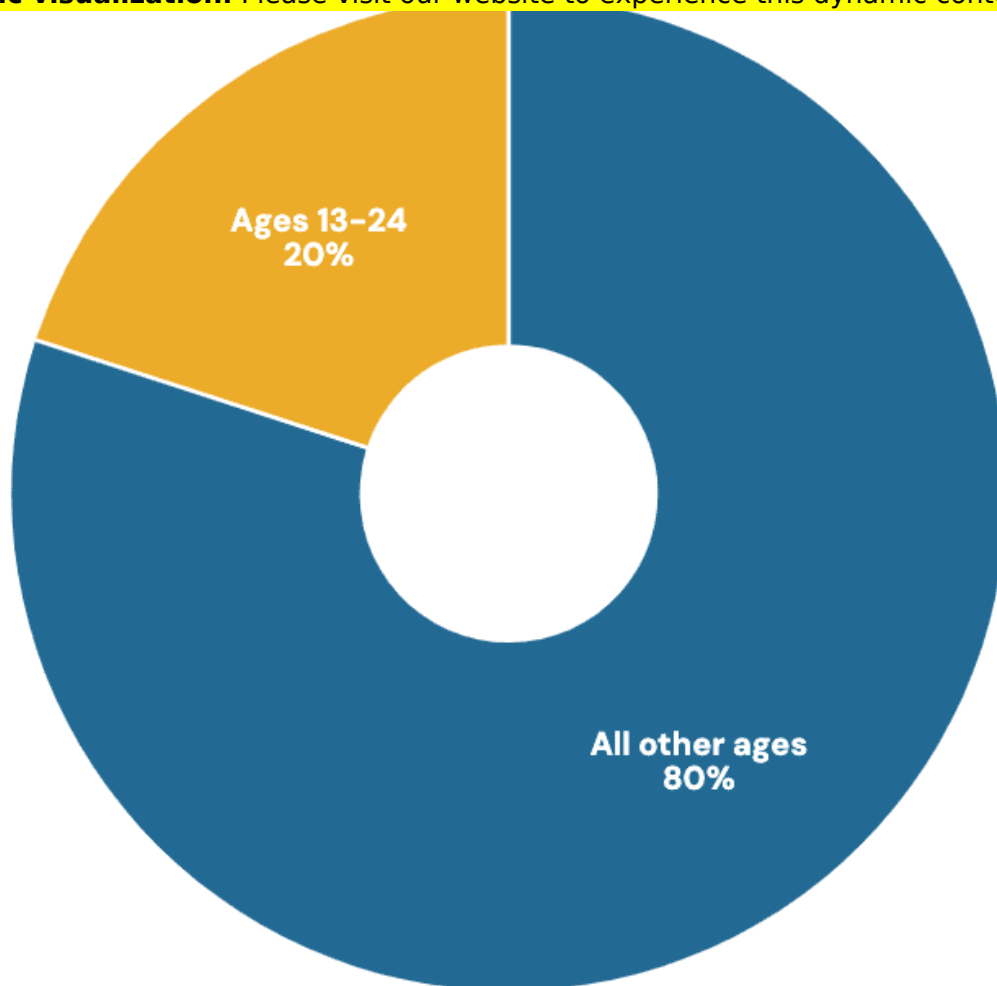


Figure 4 ATN 110 - HIV PrEP Demonstration Project and Safety Study for Young MSM

Source: Hosek SG, Rudy B, Landovitz R, et al. An HIV Preexposure Prophylaxis Demonstration Project and Safety Study for Young MSM. J Acquir Immune Defic Syndr. 2017;74:21-9.

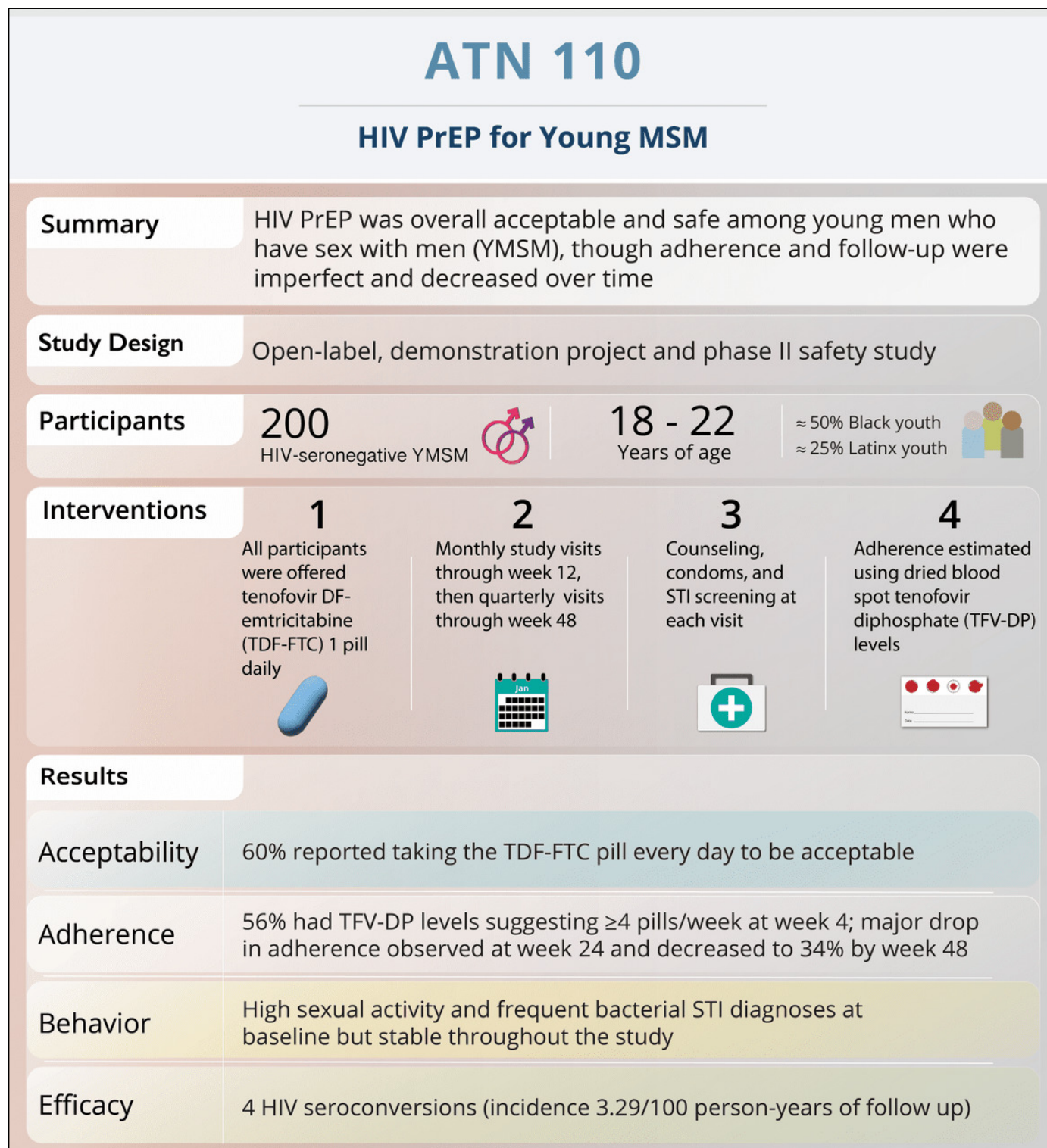


Figure 5 ATN 113 - Safety and Feasibility of HIV PrEP for Adolescent Men Who Have Sex with Men

Source: Hosek SG, Landovitz RJ, Kapogiannis B, et al. Safety and Feasibility of Antiretroviral Preexposure Prophylaxis for Adolescent Men Who Have Sex With Men Aged 15 to 17 Years in the United States. JAMA Pediatr. 2017;171:1063-71.

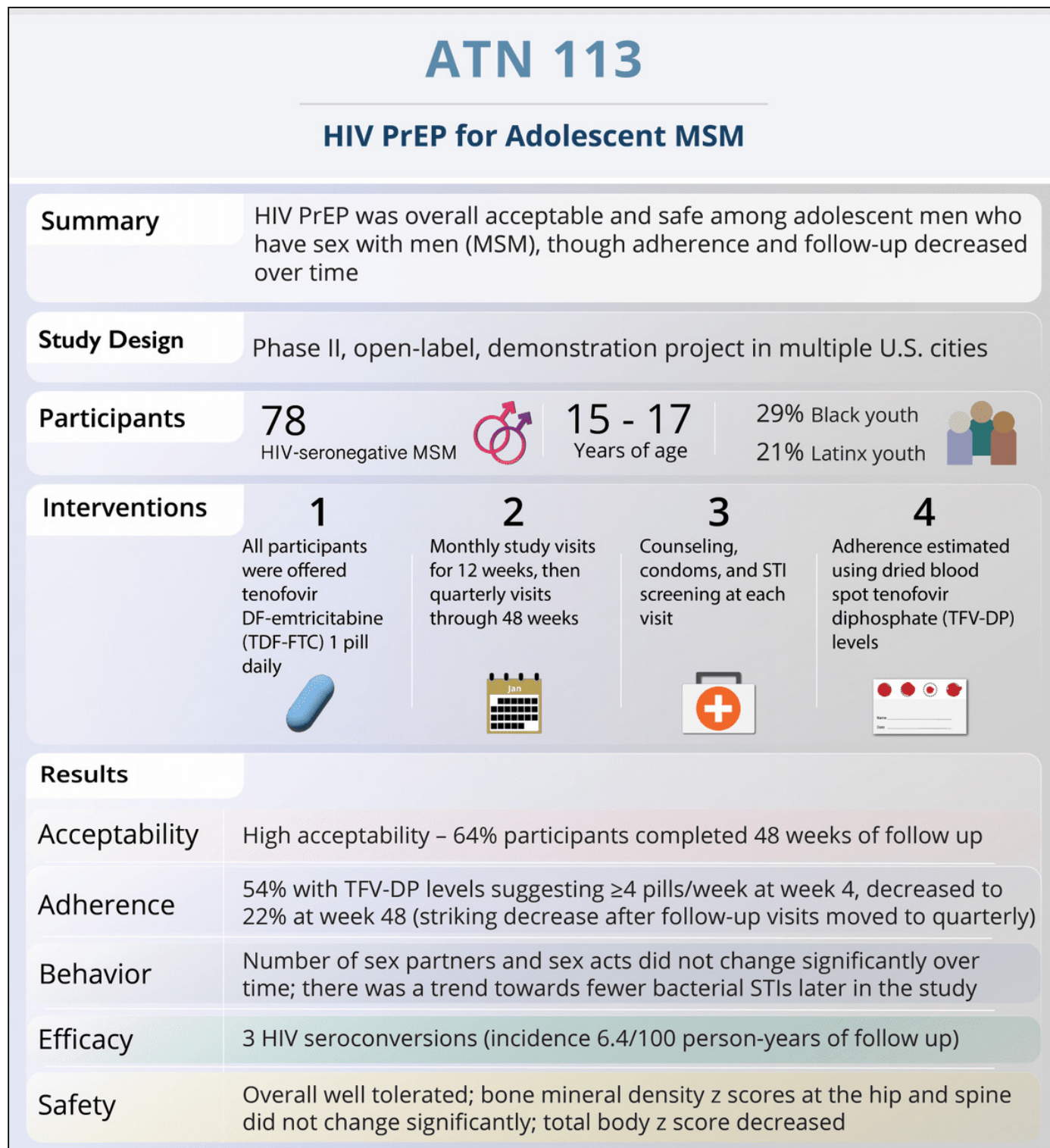


Figure 6 Assessing Indications for PrEP in Sexually Active Adolescents and Young Adults

Source: Centers for Disease Control and Prevention: US Public Health Service: Preexposure prophylaxis for the prevention of HIV infection in the United States—2021 Update: a clinical practice guideline. December 2021:1-108.

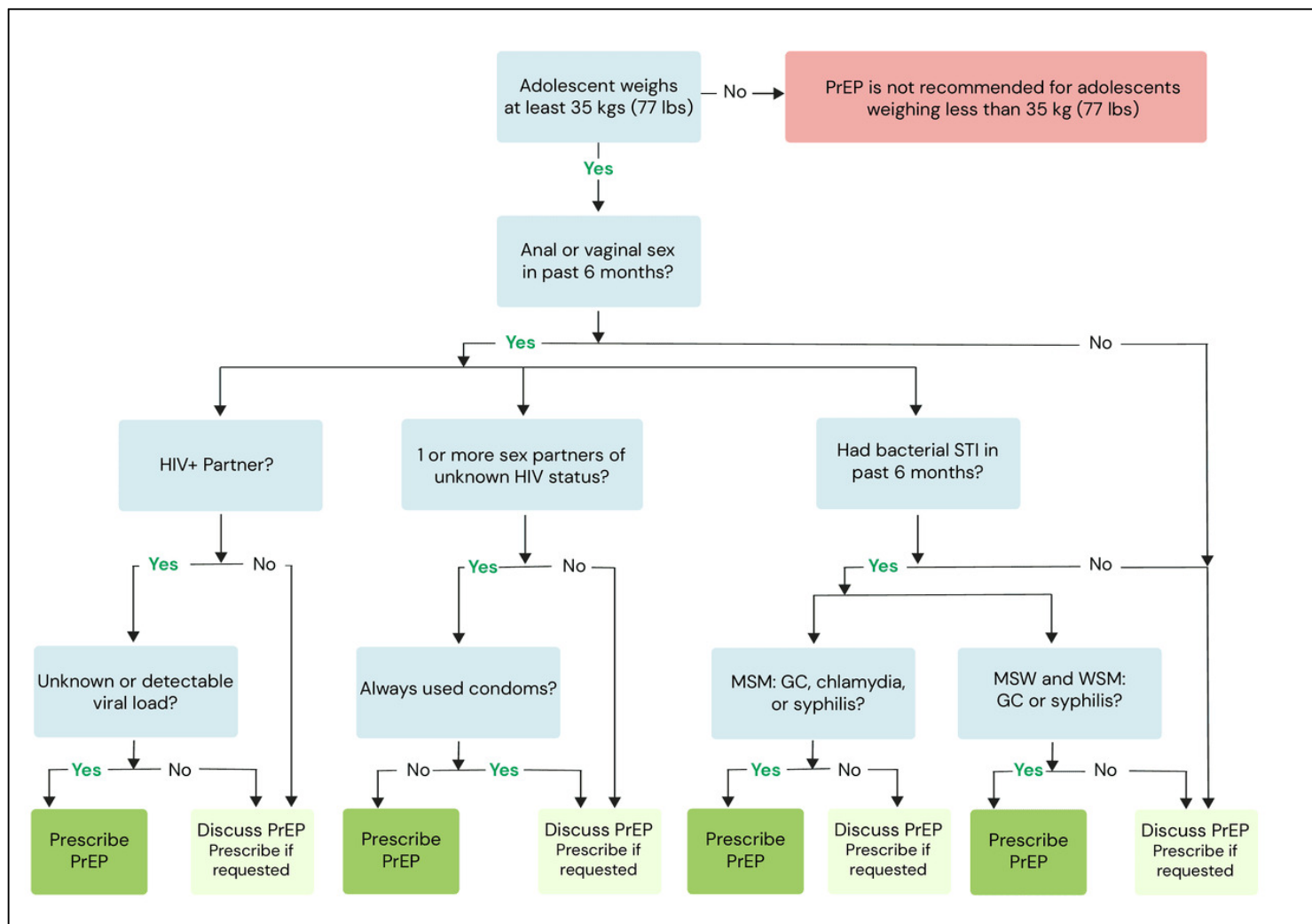


Figure 7 Assessing Indications for PrEP in Young Adults Who Inject Drugs

Source: Centers for Disease Control and Prevention: US Public Health Service: Preexposure prophylaxis for the prevention of HIV infection in the United States—2021 Update: a clinical practice guideline. December 2021:1-108.

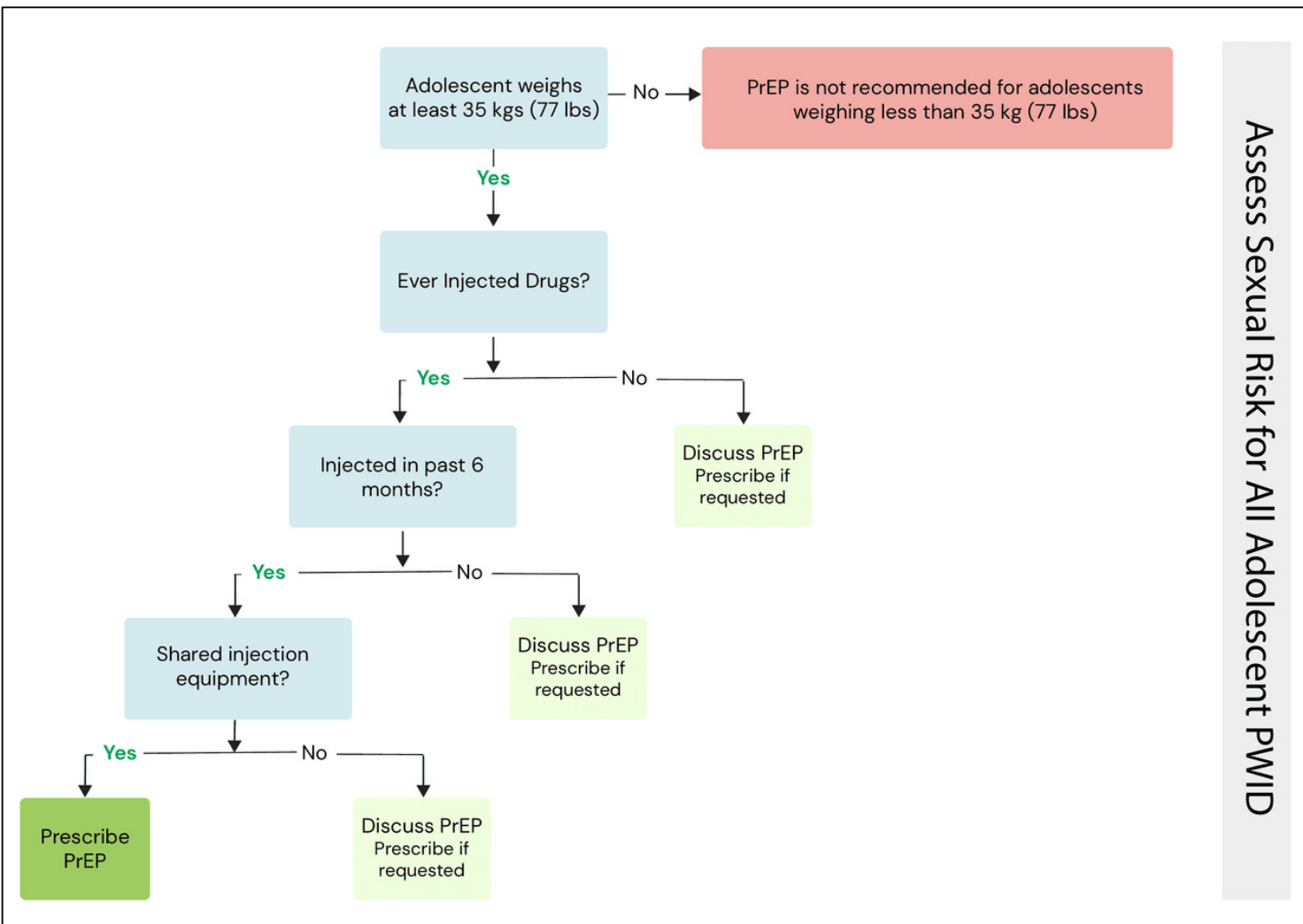


Figure 8 Adolescent PrEP Adherence

Source: (1) ATN-110: Hosek SG, Rudy B, Landovitz R, et al. An HIV Preexposure Prophylaxis Demonstration Project and Safety Study for Young MSM. *J Acquir Immune Defic Syndr*. 2017;74:21-9.(2) ATN-113: Hosek SG, Landovitz RJ, Kapogiannis B, et al. Safety and Feasibility of Antiretroviral Preexposure Prophylaxis for Adolescent Men Who Have Sex With Men Aged 15 to 17 Years in the United States. *JAMA Pediatr*. 2017;171:1063-71. (3) Gill K, Johnson L, Dietrich J, et al. Acceptability, safety, and patterns of use of oral tenofovir disoproxil fumarate and emtricitabine for HIV pre-exposure prophylaxis in South African adolescents: an open-label single-arm phase 2 trial. *Lancet Child Adolesc Health*. 2020;4:875-83.

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Adherence in Adolescent PrEP

Protective level of tenofovir diphosphate levels (>700 fmol/punch of drug) detected consistent with taking ≥ 4 pills/week

