

LET'S TALK ABOUT HIV: INTEGRATING
HIV TESTING, TREATMENT, AND
PREVENTION IN PRIMARY CARE

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1

OBJECTIVES

- **After this training, participants will be able to:**
 - Identify Primary Care's role in the HIV Care Continuum
 - Explain HIV Basics, including HIV status neutral care, testing guidelines, treatment and prevention methods
 - Describe the importance of primary care in preventing, identifying, and treating HIV

2

DISCLOSURES

None

3

WHAT'S THE FIRST WORD THAT COMES TO MIND WHEN YOU HEAR "HIV"?

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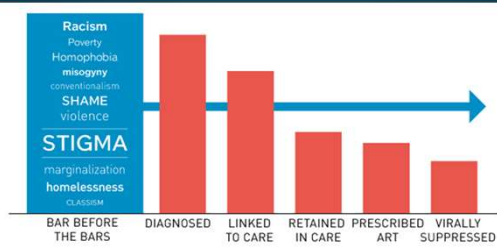


HIV CARE CONTINUUM:

The series of steps a person with HIV takes from initial diagnosis through their successful treatment with HIV medication.

6

Primary Care – the Key to Prevention



7

THERE IS NO SUCH THING AS A
SINGLE-ISSUE STRUGGLE BECAUSE WE DO
NOT LEAD SINGLE-ISSUE LIVES.

Audre Lorde

8



SOCIAL DETERMINANTS OF HEALTH

<https://www.cdc.gov/publichealthgateway/sdoh/index.html>

9



**THE SDOH THAT'S OFTEN OVERLOOKED:
SOCIAL AND COMMUNITY CONTEXT**

Stigma

- Discrimination
- Racism
- Safety
- Community Connections
- Relationships
 - Family, friends, community

10

“Why don’t they just....?”



Internal Stigma: Internal beliefs about public attitudes that result in negative consequences

External Stigma: Public beliefs about people with have devalued characteristics

Structural Stigma: Policies, decreased social supports, impacts on criminal justice system, hiring practices

STIGMA — A SET OF NEGATIVE BELIEFS CONNECTED TO A PARTICULAR PERSON, CIRCUMSTANCE, OR QUALITY DUE TO PERCEIVED CONTROL

Earnshaw VA, Am Psychol (2020)

11



Always remember...

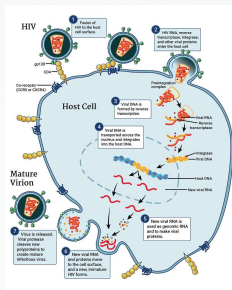
**HIV is a *medical diagnosis*,
NOT a character flaw.**

12

HIV BASICS

13

HUMAN IMMUNODEFICIENCYVIRUS (HIV)



- HIV is a virus that impacts the body's immune system
- It attacks the CD4 cells in a person's immune system – this makes it difficult for them to fight infections
- It is not generally curable, but it is treatable
- If not treated, it can lead to Acquired Immunodeficiency Syndrome (AIDS) – now called late-stage HIV

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14

HIV and AIDS: What's the Difference?

HIV

- HIV is the virus that causes HIV infection.
- HIV damages the immune system by killing CD4 cells.

CD4 Cells

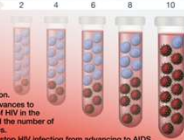
- CD4 cells are part of the immune system.
- HIV attacks and kills CD4 cells.
- Loss of CD4 cells makes it hard for the body to fight off infections.

AIDS

- AIDS is the last stage of HIV infection.
- As HIV infection advances to AIDS, the amount of HIV in the body increases and the number of CD4 cells decreases.
- HIV medicines can stop HIV infection from advancing to AIDS.
- Without HIV medicines, HIV advances to AIDS in about 10 years.
- ART (Antiretroviral treatment) can prevent HIV from spreading, and prevent HIV from advancing to AIDS.

For more information, visit HIVinfo.NIH.gov.

Years without HIV medicines



15

How is HIV Acquired?

SIMPLY PUT: Blood, semen and pre-semenal fluid, rectal fluids, vaginal fluids, breast milk

- Blood
- Semen and pre-semenal fluid
- Rectal fluids / anal sex
- Vaginal fluids / vaginal sex
- Oral sex
- IV drug use or sharing injection equipment
- Needle sticks
- Pregnancy
- Breast milk

Only **certain body fluids** from a person who has HIV **can transmit HIV**.

Source: <https://www.cdc.gov/hiv/causes/index.html>

16

ACUTE HIV INFECTION

fever
fatigue
headache
sore throat
swollen lymph glands
short-term nausea
muscular aches and pain
rash

Vervwellhealth.com

17

SYMPTOMS OF HIV

Many people are asymptomatic
- May stay this way for years

Early Infection: Fever, body aches, chills, night sweats, cough, swollen lymph nodes, rash. (Think Mono)

Late Infection: Weight loss, fatigue, frequent infections, certain cancers, all symptoms listed above, and

18

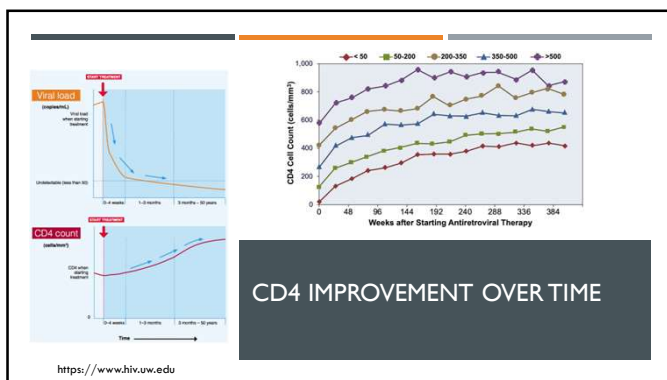
EXPLAIN "THE NUMBERS"

 **CD4** – How well your immune system is functioning (Normal 500 – 1200)

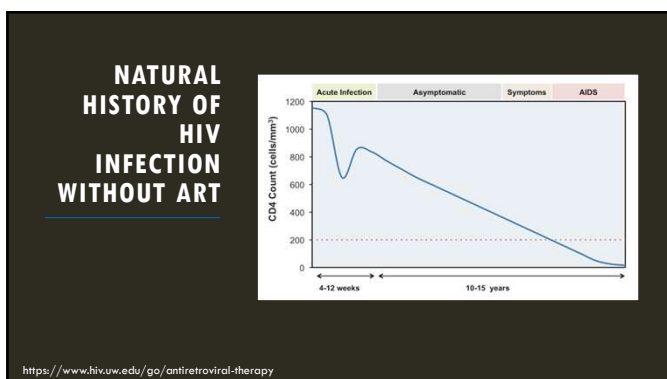
 **Viral Load** – How much HIV is circulating per ml of blood

 Typically, as the viral load increases, the CD4 decreases

19



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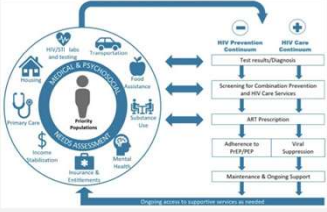
INITIATE ART AS SOON AS POSSIBLE AFTER DIAGNOSIS

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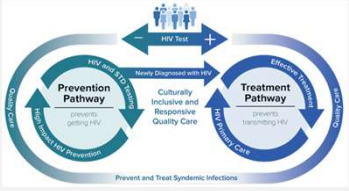
AIMS FOR STATUS NEUTRAL CARE

Follow CDC guidelines to test people for HIV

- Person-centered, "whole-person", respecting autonomy and person's health goals
- Improves accessibility for services
- Reduces stigma and biases
- Creates efficiencies
- Decreases new HIV infections



23



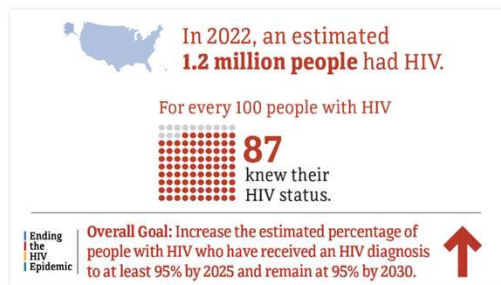
STATUS NEUTRAL

Follow CDC guidelines to test people for HIV. Regardless of HIV status, quality care is the foundation of HIV prevention and effective treatment. Both pathways provide people with the tools they need to stay healthy and stop HIV.

24

HIV TESTING

25



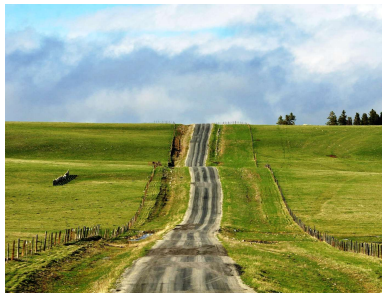
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South Dakota's HIV Disparities

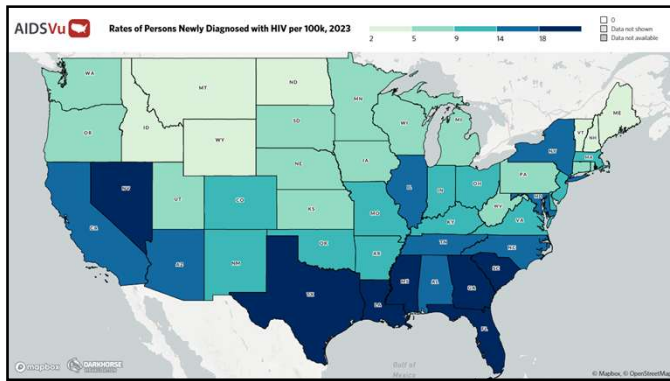
-Only about 30% of SD residents have been tested

-Nearly 30% have AIDS-defining condition at time of diagnosis

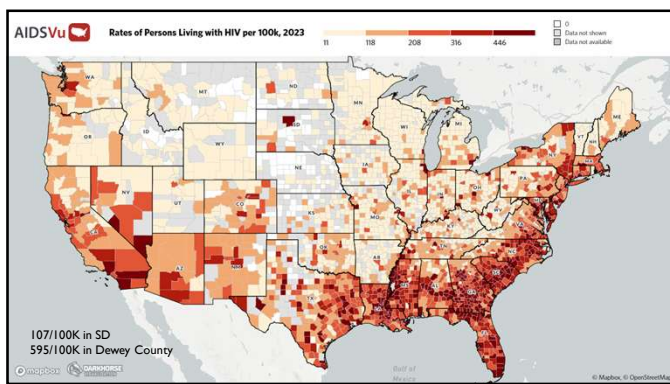
-The average person living with HIV spends more than \$550,000 on medical care in their lifetime



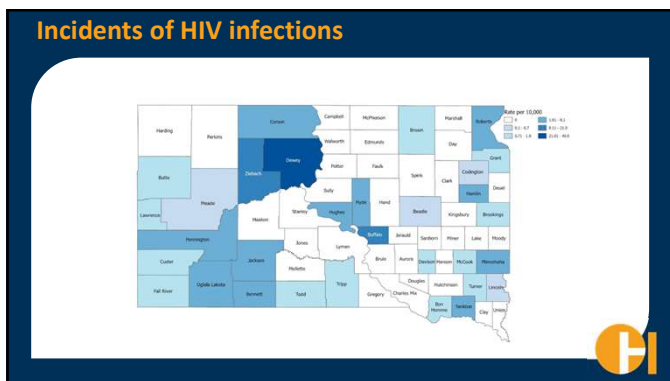
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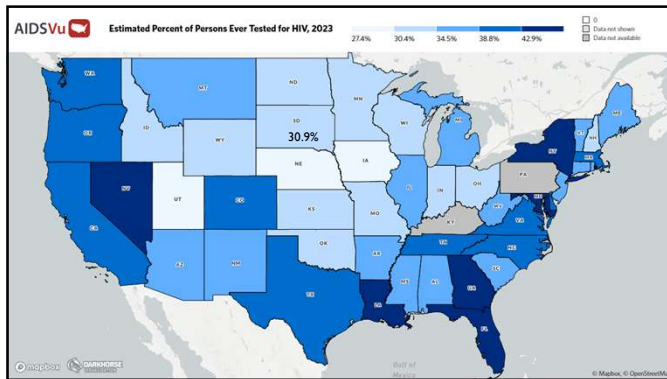
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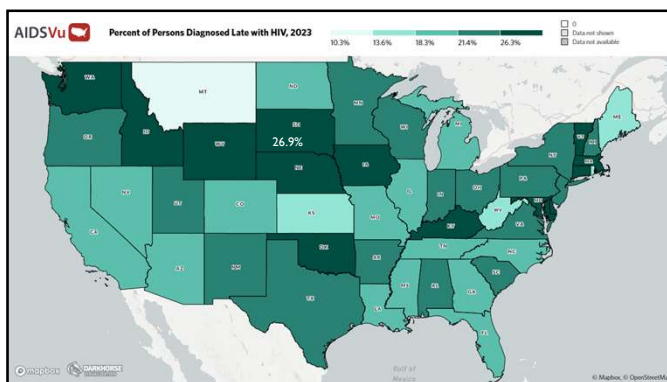
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31



32

Recommendations for HIV Testing

“Opt-out testing” ALWAYS

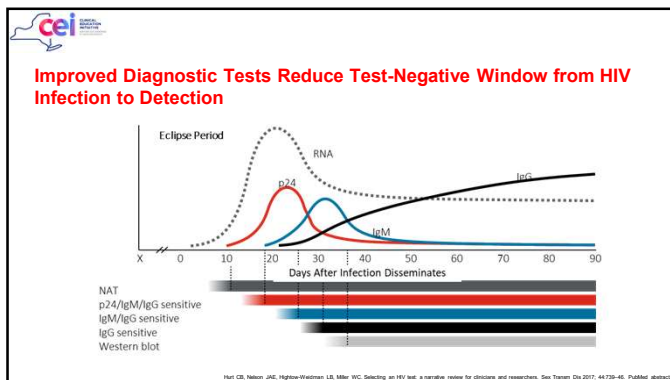
- HIV testing is the *STANDARD OF CARE* with any STI check
- One-time testing for low-risk patients – Everyone age 13-64 as part of routine care
 - Consider ending upper age limit
- Annual or more frequent testing for high-risk patients
 - People with history of IVDU
 - MSM (3-6 months)
 - Persons engaged in sex work or commercial sexual exploitation
 - Sex partners of HIV-positive individuals, IVDU
 - Pregnant Women
 - Recent treatment for an STI – Increases risk of HIV acquisition by 3-5x

33

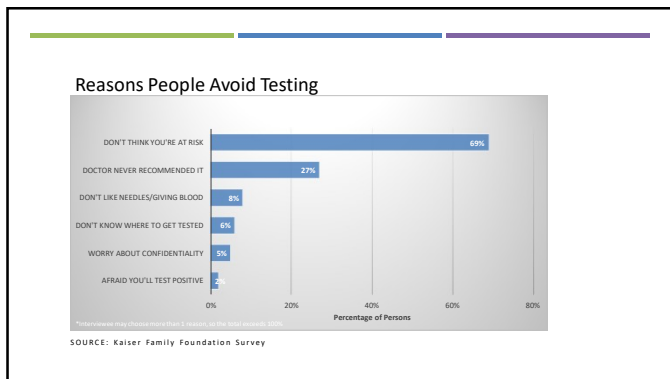
TYPES OF HIV TESTS

- Rapid HIV Ab** testing - detects HIV Ab, 6-8-week window period s/p exposure; some false positives (particularly in pregnant women); inexpensive; easy to run; results in 10-15 minutes; available OTC at the pharmacy
- p24 Ag/Ab** testing - detects p24 Ag and HIV Ab; 10-14-day window period; serum test; high sensitivity and specificity; increased cost; need a lab to run (usually back in 24 hours)
- HIV PCR** - detects HIV virus itself; better known as a VL; confirmatory test for all HIV screening tools; sensitive testing can detect HIV within days of exposure; very expensive; back in 72 hours (usually back in 24 hours)

34



35

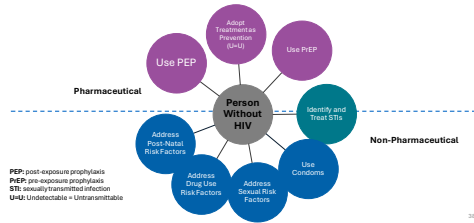


36

HIV PREVENTION

37

Current HIV Prevention Options (Slide credit: CDC)



38

Added benefit of HIV treatment – it works for prevention!



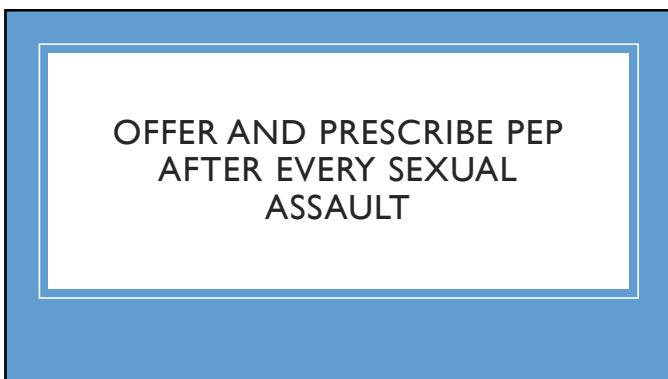
39



40

PrEP vs. PEP		
When you take steps to protect yourself against a disease, the HIV is called prophylaxis. PrEP and PEP are for protecting people who are HIV negative.		
PrEP stands for pre-exposure prophylaxis.	What's it called?	PEP stands for post-exposure prophylaxis.
Before HIV exposure: PrEP is taken before sex, drug use, or other HIV exposure.	When is it taken?	After HIV exposure: In emergency situations, PEP is started within 72 hours after possible exposure, and taken for a month thereafter.
PrEP is for people who don't have HIV and: • are at risk of getting HIV from sex • are at risk of getting HIV from injection drug use	Who's it for?	PEP is for people who don't have HIV but may have been exposed: • during sex • at work through a needlestick or other injury • during a sexual assault • by sharing injection drug equipment
Consistent use of PrEP can reduce the risk of getting HIV from sex by about 99%, and from injection drug use by at least 75%.	How effective is it?	PEP can prevent HIV when taken correctly, but it is not always effective. Start PEP as soon as possible to give it the best chance of working.
Ask your health care provider about a prescription for PrEP, or use HIVinfo.org to find a health care provider in your area who can prescribe PrEP.	How do you get it?	Within 72 hours after potential exposure to HIV, get a PEP prescription from your health care provider, urgent care, or an emergency room.
For more information, visit HIVinfo.nih.gov .		

41



42

Who Is PrEP For?

(Slide credit: CDC)

PrEP is for adults and adolescents without HIV who are at risk of getting HIV from sex or injection drug use.



CDC's PrEP guideline includes telling all sexually active adults and adolescents that PrEP can protect them from getting HIV

Giving patients information about PrEP:

- Increases the number of people who know about PrEP and equips them to share the information with their social networks and family members
- Helps patients overcome embarrassment and stigma so they can respond accurately to risk assessment questions

PrEP can be prescribed to any adult or adolescent patient who asks for it, even if they do not report HIV risk factors, as part of their comprehensive prevention plan

Centers for Disease Control and Prevention, US Public Health Service. Prescriptions principles for the prevention of HIV infection in the United States—2021 update: clinical practice guideline. Published December 2021. Accessed January 26, 2023. <https://www.cdc.gov/hiv/pdf/2021-principles-for-prep-guidelines-2021.pdf>

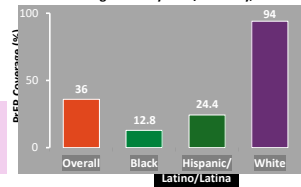
43

43

The PrEP Gap in the United States

- 1.2 million people are likely to benefit from PrEP
- But in 2022, among those who could benefit, few were prescribed PrEP
 - Only 41.0% of men
 - Only 14.5% of women
 - Only 23.5% of those aged 16-24 yr
- Significant inequities in PrEP coverage exist based on race/ethnicity

PrEP Coverage in US by Race/Ethnicity, 2022



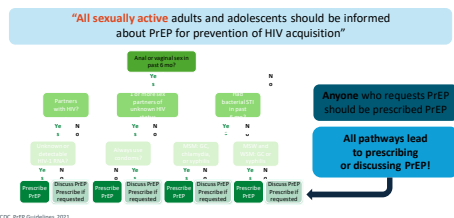
American Indians/Alaska Natives: Compared with White counterparts, HIV infection rates $\geq 2\times$ higher; 42% have heard of PrEP, only 3.5% were currently taking PrEP or had used PrEP in the past 12 mo

[cdc.gov/hiv/library/reports/surveillance-data-tables/vol-4-no-3/index.html](https://www.cdc.gov/hiv/library/reports/surveillance-data-tables/vol-4-no-3/index.html)
[minorityhealth.hhs.gov/health-and-american-indians-alaska-natives](https://www.minorityhealth.hhs.gov/health-and-american-indians-alaska-natives)
[cdc.gov/hiv/pdf/funding/announcements/ps18-1802/cdc-hiv-ps18-1802-monitoring-evaluation-report-2021.pdf](https://www.cdc.gov/hiv/pdf/funding/announcements/ps18-1802/cdc-hiv-ps18-1802-monitoring-evaluation-report-2021.pdf)

Slide credit: [clinicaloptions.com](https://www.clinicaloptions.com)

44

PrEP Guidance: CDC



45

Expanding PrEP Uptake by Expanding HCPs Who Prescribe It

- **Does not require HIV expertise**
 - PrEP uptake will remain stagnant if few prescribe

Lesson from contraception care in the United States

Expansion of contraception care services (beyond OB/GYN and midwives) increased access to and prescription of contraception

Other prescribing HCPs: family medicine physicians, internal medicine physicians, pediatricians, and APPs

May be especially important in more rural areas

Chen. Am J Obstet Gynecol. 2022;226:232.e1.

46

Mechanisms to Improve Conversations About PrEP

Normalize PrEP as a prevention option for *any* person who is (or plans to be) sexually active

- Initiate the conversation
- Build rapport and trust with your patients
- Provide person-centered care
- **Shared decision-making**
 - Promotes cultural competence and humility
 - Builds trust within patient-provider relationship
- **Avoid stigmatizing language**
- **Set up safe clinic spaces**
 - Promote representation of racial and ethnic minority and transgender people
 - Display signals of acceptance (eg, wear pride pin on clothing)
 - Pronouns on ID tag
- Integrate sexual health and harm reduction in your practice

McKulthly. Cult Health Sex. 2022;24:1033. Gollub. Curr HIV/AIDS Rep. 2018;15:190. cdc.gov/research/action/stories/detail-county-talk.html.

Slide credit: clinicaloptions.com



47

PrEP Formulations

- 2012: Oral FTC/TDF (Truvada)
- 2019: Oral FTC/TAF (Descovy)
- 2020: Generic PrEP
- 2021: Long Acting (LA) Cabotegravir Injectable (Apretude)
- 2025: Lenacapavir (Yeztugo)



48

ORAL PREP

49

TENOFOVIR DF-EMTRICITABINE (TRUVADA)

Approved Indication for HIV PrEP: Tenofovir DF-emtricitabine is indicated for HIV PrEP to reduce the risk of sexually acquired HIV in adults and adolescents (weighing at least 35 kg).

Formulation/Dosing: Tenofovir DF-emtricitabine is a fixed-dose combination that contains 300 mg of tenofovir DF and 200 mg of emtricitabine. The recommended dosing of tenofovir DF-emtricitabine when used for HIV PrEP is one tablet once daily, with or without food. Alternative dosing, such as on-demand (2-1-1) dosing, is not included in the FDA indication but can be considered "off-label" for select men who have sex with men.

Use in Persons with Renal Impairment: Tenofovir DF-emtricitabine is not recommended for HIV PrEP in persons with an estimated creatinine clearance of less than 60 mL/min.

<https://www.hiv.uw.edu/>

50

TENOFOVIR ALAFENAMIDE- EMTRICITABINE (DESCOVY)

Approved Indication for HIV PrEP: Tenofovir alafenamide-emtricitabine is indicated for HIV PrEP in adults and adolescents (weighing at least 35 kg) at risk of acquiring HIV sexually, excluding women at risk from receptive vaginal sex.¹ Tenofovir alafenamide-emtricitabine is not indicated for receptive vaginal sex because its effectiveness in this population has not been established, although it is currently under investigation.

Formulation/Dosing: Tenofovir alafenamide-emtricitabine is a two-drug, fixed-dose combination that contains 25 mg of tenofovir alafenamide and 200 mg of emtricitabine. Oral tenofovir alafenamide-emtricitabine should be taken as one tablet once daily, with or without food. Alternative dosing, such as on-demand (2-1-1) dosing, is not recommended.

Use in Persons with Renal Impairment: For HIV PrEP, tenofovir alafenamide-emtricitabine is not recommended for persons who have an estimated creatinine clearance of less than 30 mL/min, unless they are on dialysis.

<https://www.hiv.uw.edu/>

51

INJECTABLE PREP

52

CABOTEGRAVIR-IM (LONG-ACTING INJECTABLE) BRAND NAME: APRETUDE

FDA-Approved Indication for HIV PrEP: Long-acting injectable cabotegravir administered as an intramuscular dose (cabotegravir-IM), is indicated for HIV PrEP in adults and adolescents (weighing at least 35 kg) who are at risk of sexual acquisition of HIV. Cabotegravir-IM has not been studied as a prevention measure for people who are at risk of acquiring HIV from injecting drugs.

Formulations/Dosing: Cabotegravir-IM is available as a 200 mg/mL solution and is administered as a 3 mL intramuscular injection in the gluteal region. Oral cabotegravir is a 30 mg tablet that is taken once daily, with or without food. Cabotegravir-IM is administered as a 600 mg (3 mL) injection, repeated 1 month after the first injection, and then administered every 2 months thereafter. An optional lead-in with oral cabotegravir 30 mg once daily may be used for approximately 1 month to assess the tolerability of cabotegravir. If the oral cabotegravir lead-in is used, the first injection of cabotegravir-IM should be given on the last day of the oral lead-in (or within 3 days of completing the oral lead-in).

Use in Persons with Renal Impairment: For HIV PrEP cabotegravir has no renal restrictions. For persons who have a creatinine clearance less than 30 mL/min, increased monitoring for cabotegravir toxicity is recommended. Hemodialysis is not expected to impact cabotegravir levels.

<https://www.hivuw.edu/>

53

LENACAPAVIR-SQ (LONG-ACTING INJECTABLE) BRAND NAME: YEZTUGO

FDA-Approved Indication for HIV PrEP: Long-acting injectable lenacapavir administered as a subcutaneous injection (lenacapavir-SQ), is indicated for HIV PrEP to reduce the risk of sexually acquired HIV-1 for adults and adolescents weighing at least 35 kg (77 lbs). Two lenacapavir-SQ medications are available for commercial use: lenacapavir-SQ (Yeztugo) for HIV PrEP and lenacapavir-SQ (Sunlenca) for HIV treatment. Although these two lenacapavir-SQ medications contain the same recommended dose and dosing frequency, it is important to designate the correct brand, based on whether the indication for use is HIV PrEP or HIV treatment. In addition, there are also two different oral brand preparations: oral lenacapavir (Yeztugo) for HIV PrEP and oral lenacapavir (Sunlenca) for HIV treatment.

Formulations/Dosing: Lenacapavir-SQ is available in 1.5 mL vials that contain 463.5 mg of lenacapavir. Lenacapavir is also available as a 300 mg tablet that can be taken with or without food. Use of lenacapavir-SQ for HIV PrEP requires a 2-day initiation phase that uses both oral and injection doses. Day 1 of the initiation phase requires a 600 mg (2 x 300 mg tablets) dose of oral lenacapavir and a 927 mg (2 injections, each with 463.5 mg) of lenacapavir-SQ. Day 2 of the initiation phase requires a 600 mg (2 x 300 mg tablets) dose of oral lenacapavir. After the 2-day initiation phase, lenacapavir-SQ 927 mg (2 injections, each with 463.5 mg injections) is administered every 26 weeks.

Use in Persons with Renal Impairment: There are no dosage adjustments of lenacapavir-SQ or oral lenacapavir in persons with mild, moderate, or severe renal insufficiency. There are insufficient data on the use of lenacapavir-SQ or oral lenacapavir in persons who have a creatinine clearance less than 15 mL/min. In addition, there are insufficient data on the use of lenacapavir-SQ or oral lenacapavir in persons receiving hemodialysis. Since lenacapavir is highly protein bound, hemodialysis is not likely to significantly impact lenacapavir levels.

54

PREP MONITORING

• Baseline

- HIV test – within 7 days
- Renal Function
- STI Testing
- Lipid Panel
- Hep B
- Hep C
- Pregnancy

<https://www.hiv.uw.edu/>

• Monitoring

- HIV test
 - Q3 months orals
 - Q 4 months Apretude
 - Q 6 months Yeztugo
- Renal Function – every 6 months if over 50, yearly if under 50
- STI Testing – MSM every 3-6 months
- Lipid Panel – every 12 months
- Hep C every 12 months in MSM, IDU
- Pregnancy – every 3-6 months

55



National HIV PrEP Curriculum

www.hivprep.uw.edu

APP: HIV PREP TOOLS FOR CLINICIANS

This app supports health care professionals to:

- Evaluate persons for HIV PrEP
- Identify appropriate HIV PrEP medications
- View recommended laboratory tests
- Assess, initiate, and monitor HIV PrEP

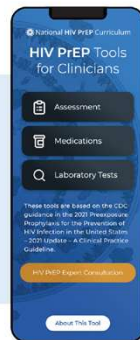
Access the app on a web browser (hivprep.uw.edu/page/tools/app), via the CDC STI Tx Guide app, or download the free app



APPLE APP STORE



GOOGLE PLAY

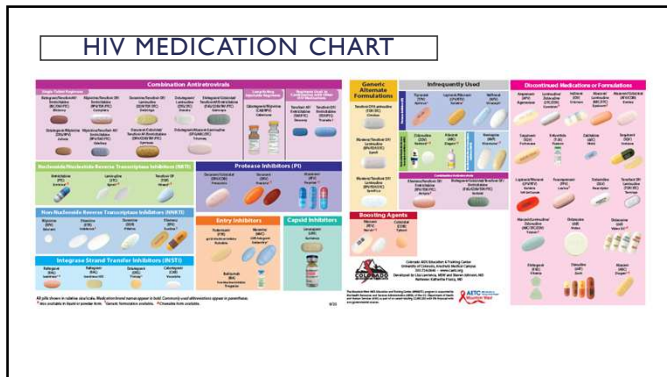


University of Washington (UW) Professor of Medicine Dr. David Spach is Editor-in-Chief of this site and four other curricula about HIV, STDs, HCV, and HBV. All sites offer FREE CE. Created and managed by the UW Infectious Diseases Education & Assessment (IDEA) Program (idea.medicine.uw.edu).

56

HIV TREATMENT

57



58

RAPID ART INITIATION

Candidate Criteria:

- New reactive HIV test, confirmed HIV or suspected acute HIV
- ART-naïve or limited prior ART
- No conditions requiring ART deferral (e.g., cryptococcal, TB meningitis, DMV, retinitis)
- Perform baseline labs; ART can start while awaiting results (A3)

59

RECOMMENDATIONS:

Rationale for Rapid ART Initiation:

- Recommend ART for all patients diagnosed with HIV (A1)
- Offer rapid ART – same day preferred, within 72 hours if eligible (A1)
 - Confirmed HIV diagnosis (A1)
 - Reactive screening, confirmatory pending (A2)
 - Acute HIV infection: Ab-negative, RNA-positive (A2)
- Counsel patients with HIV-negative partners on reduced transmission risk once virally suppressed; strongly recommend ART (A1)
- Involve patients in ART decisions; patient decides timing/initiation (A3)
- If patient declines rapid ART, revisit as soon as possible
- Initiate ART in advanced HIV/AIDS even with adherence barriers; refer to specialized support (A2)
- Monitor response or consult HIV specialist after initiation (A2)

Rapid ART initiation, the standard of care, is safe, effective, and widely accepted; even patients facing substance use, mental health, or housing should be offered immediate ART, with counseling and support services addressing potential adherence barriers.

60

PROTOCOL FOR RAPID ART INITIATION

ART can be started while awaiting laboratory test results

Identify Rapid ART Candidates	Assessment, Counseling, and Referrals	Baseline Lab Testing	Payment Assistance	Initiate ART	Follow-up
<ul style="list-style-type: none"> Candidates have: A new reactive POC HIV test result, new HIV diagnosis, known or suspected acute HIV, or known HIV, and No or limited prior ART use (except PEP/PrEP), and No known medical conditions or OIs that require deferral of ART initiation 	<ul style="list-style-type: none"> Assess health literacy Discuss HIV diagnosis and disclosure Discuss ART benefits, efficacy, and management Identify and address medical and psychosocial barriers to treatment and adherence Refer for substance use treatment, behavioral health services, and housing assistance, as needed 	<ul style="list-style-type: none"> HIV Ag/Ab test Viral load Resistance testing CD4 count HAV, HBV, and HCV testing STI testing (including syphilis) Urinalysis Pregnancy test for individuals of childbearing potential 	<ul style="list-style-type: none"> Assess need for payment assistance Refer patients with no insurance to a Certified Application Counselor or Navigator Provide resources for sliding fee and/or payment assistance 	<ul style="list-style-type: none"> Choose among preferred regimens based on patient characteristics and preference Initiate ART immediately – preferably on the same day – or within 72 hours Administer the first dose on-site if possible 	<ul style="list-style-type: none"> Contact the patient within 24 to 48 hours by phone (or other preferred method) Assess medication tolerance and adherence If feasible, schedule in-person visit with medical care provider within 7 days Reinforce adherence Change or adjust the initial ART regimen as needed based on results of initial lab and/or resistance testing

61

MEDICAL HISTORY CHECKLIST

Before Rapid ART Initiation, Ask About:

- Date and result of last HIV test
- Serostatus of sex partners and their ART regimens (if known)
- Prior use of antiretrovirals, including PrEP or PEP (with dates)
- Comorbidities, especially renal or liver disease and hepatitis B
- Current prescribed and OTC medications
- Drug allergies
- Substance use
- Symptoms concerning for – cryptococcal meningitis, TB meningitis, and CMV retinitis
 - Guidelines for the Prevention and Treatment of Opportunistic Infections in Adults and Adolescents with HIV
- Psychiatric history, including depression, psychosis, or suicidality
- Possible pregnancy and reproductive plans (if applicable)

A Guide for Health Care Providers



62

INITIAL WORK-UP

Baseline

- CBC
- CMP
- Hgb A1c
- TSH
- Hepatitis Panel | Hep B serology
- Vaccine titers – Varicella, MMR, Hep A, Hep B
- TB Gold
- EBV, CMV, Toxoplasmosis
- Lipid Panel
- Vitamin D
- Chlamydia, Gonorrhea, Syphilis
- Pregnancy Test

HIV Labs

- CD4
- Viral Load
- Genotype
- HLA B5701
- Tropism Testing

Health Maintenance

- Pap smear, Pelvic Exam
- Mammogram, CBE
- Colonoscopy
- Low Dose CT Scan
- Vaccine Updates
- Bone Density
- STI testing
- Mental Health Screening
- Substance Use Screening

63

MY PATIENT IS ON OTHER MEDS – NOW WHAT DO I DO?

HIV and HCV Drug Interactions: Quick Guides for Clinicians

University of Liverpool: HIV Drug Interactions Checker

64

MEDICATIONS – KEY CONCEPTS

STERIODS

- HIV medications can increase concentrations
 - Cushing's syndrome adrenal suppression
- More common with "boosters"
- Adjust dosing for:
 - Most inhaled steroids
 - Prednisone
- AVOID Flonase

OVER THE COUNTER (OTC) MEDS

- St. John's Wort – CYP450 3A4
 - May reduce concentration of PIs and NNRTIs by as much as 82%
- Garlic – topic of debate
- PPIs and H2 receptor antagonists
- Antacids – magnesium and/or aluminum
 - Neutralizes stomach acids and may interfere with absorption of ART (space out the dosing if necessary)

65

HOW DO I MONITOR MY PATIENTS' VIRAL LOAD

The frequency of viral load testing depends on several factors. Current guidelines recommend viral load monitoring as follows:

- With initiation of ART (before initiation and within 2 to 4 weeks after treatment initiation, followed by 4- to 8-week intervals until the levels become undetectable)
- After ART modification because of suboptimal response (within 2 to 4 weeks after treatment modification, followed by 4- to 8-week intervals until the levels become undetectable)
- After ART modification because of toxicity or need for regimen simplification (within 4 to 8 weeks after changing therapy)
- In patients on a stable, suppressive ART regimen (every 3 to 6 months, or every 6 months if virally suppressed for more than 2 years, to confirm durable viral suppression)
- In patients with suboptimal response (frequency depends on clinical circumstances)

A patient's plasma HIV RNA viral load should be measured regularly to confirm initial and sustained response to ART. Most patients taking ART as prescribed achieve viral suppression within 6 months.

66

RECOMMENDED FREQUENCY OF CD4 MONITORING

September 2025 HHS Guidelines Update:

Clinical Scenario	CD4 Monitoring
After initiating ART	3 months after initiation
During first 1-2 years of ART with viral suppression and CD4 count ≥ 300 cells/mm ³	Every 6 months
After 1-2 years of ART with viral suppression and CD4 count ≥ 300 cells/mm ³	Optional
If CD4 count < 300 cells/mm ³	Every 3-4 months
After modifying ART due to virologic failure	Every 3 to 6 months

67

RESOURCES

68

HIV Symptom Evaluation Guides

Each HIV Symptom Evaluation Guide provides clinicians with a framework for evaluating certain common symptoms that individuals with HIV may experience. These guides offer a summary of important clinical questions and considerations, as well as decision trees to help with the evaluation process.

Download All Guides

Share



Cough and Dyspnea

Odynophagia

Diarrhea

Headache

Memory Changes

<https://www.hiv.uw.edu/page/symptom-evaluation/guides>

Developed by
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UNIVERSITY OF
WASHINGTON

An AIDS Program
AETC

Part of
IDEA

69



National HIV Curriculum

www.hiv.uw.edu

Six modules with 35 lessons and corresponding Question Bank topics address:

SCREENING AND DIAGNOSIS	CO-OCCURRING CONDITIONS
BASIC HIV PRIMARY CARE	PREVENTION OF HIV
ANTIRETROVIRAL THERAPY	KEY POPULATIONS

CREATE FREE ONLINE LEARNING GROUPS TO:

- Invite, train, and onboard staff and residents by assigning select content
- Review group progress reports to track CE and shape future training
- Augment medical, nursing, pharmacy, dental and other healthcare professionals training programs



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70

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71

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72
