Nonoccupational Postexposure Prophylaxis

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Section 1: Prevention of HIV
Topic 4: Nonoccupational Postexposure Prophylaxis

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Introduction and Background

In the mid 1990s, postexposure prophylaxis (PEP) was recognized as a safe and effective intervention to prevent the acquisition of HIV for healthcare workers exposed to HIV-contaminated blood or body fluids. In contrast, the use of nonoccupational PEP for a sexual or injection drug use HIV-related exposure has been more controversial. In 1997, the Centers for Disease Control and Prevention (CDC) concluded there was insufficient evidence regarding the efficacy of nonoccupational PEP to recommend either for or against its use, but in 2005 the CDC and the Department of Health and Human Services revised its position and issued nonoccupational PEP guidelines.[1] These guidelines were recently updated by the CDC and Department of Health and Human Services as the 2016 Nonoccupational PEP Guidelines.[2]

Preferred strategies for preventing HIV acquisition include daily use of HIV preexposure prophylaxis (PrEP), consistent and correct use of condoms, abstinence from injection drug use (or if this is not possible, then consistent use of sterile injection equipment).[2] Nevertheless, in some instances, the need for nonoccupational PEP arises, due to condom breakage, condomless sex with a person later identified as HIV-positive, sexual assault, or sharing injection equipment with a person who has HIV. Although the 2016 Nonoccupational PEP Guidelines and the 2005 version of these guidelines clearly recommend use of antiretroviral PEP for these type of nonoccupational exposures to HIV, the actual use of nonoccupational PEP has not achieved widespread acceptance and implementation.[3] Underutilization of nonoccupational PEP has multiple causes, but in some instances, medical mistrust rooted in experiences of HIV stigma, racism, homophobia or transphobia has impeded individuals from accessing these services.[4]

Clinicians should view the use of nonoccupational PEP as one of many HIV prevention options and utilize nonoccupational PEP when indicated. It is extremely important that clinicians evaluating individuals for nonoccupational PEP should recognize these individuals may be candidates to receive PrEP following completion of the nonoccupational PEP.[5]
Rationale for Providing Nonoccupational PEP

Due to ethical and logistical reasons, it is highly unlikely that a prospective randomized, placebo-controlled trial to evaluate nonoccupational PEP in humans will ever take place. In addition, human nonoccupational PEP studies are problematic because participants may have multiple exposures over the surveillance-testing period, making it difficult to discern the true benefit of nonoccupational PEP for a single exposure event.[6] Thus, the rationale for providing nonoccupational PEP is based on extrapolation from use of PEP in other settings, animal studies, retrospective reviews, and observational nonoccupational PEP reports.

Extrapolation from Occupational and Perinatal PEP Data

The rationale for nonoccupational PEP is based on the efficacy of PEP following occupational exposures to HIV.[2,7] Most notably, in 1997, investigators reported findings from a case-control study involving health care workers who sustained needlestick injuries from source individuals with HIV; this study demonstrated occupational PEP with oral zidovudine, taken within 4 hours by most of the participants, reduced the risk of HIV seroconversion by 81%.[7] In addition, several important perinatal transmission trials involving mothers with HIV have established the benefit of using PEP given to the mother during labor and to the baby following birth.[8] For example, a Ugandan study reported that administering single-dose nevirapine to mothers during labor and to their infants within 72 hours of birth reduced the vertical HIV transmission rate from 25.1 to 13.1%.[9] Multiple studies have also demonstrated the benefit of antepartum and intrapartum antiretroviral therapy to prevent mother-to-child HIV transmission, as well as the efficacy of extended antiretroviral prophylaxis for HIV-exposed infants to prevent breastfeeding transmission.[10,11]

Animal Studies

The rationale for the use of antiretroviral medications for nonoccupational exposures is also partially derived from animal PEP studies. One of the earliest studies showed that tenofovir reduced the rate of seroconversion among macaques inoculated intravenously with simian immunodeficiency virus (SIV), with the greatest reduction in transmission achieved when prophylaxis was initiated as early as possible and continued for 28 days (Figure 1) and (Figure 2).[12] A later study showed that tenofovir-based PEP is also effective in preventing HIV acquisition after intravaginal inoculation of female macaques with HIV-2: tenofovir prevented seroconversion in all 8 of the female macaques exposed to HIV-2 when initiated within 12 to 36 hours.[13] A systematic review and meta-analysis of PEP using pooled data of nonhuman primates across 18 studies (mostly involving intravenous inoculation with HIV) further substantiated the efficacy of PEP when initiated as soon as possible after HIV exposure.[14]

Nonoccupational PEP Data in Humans

Although the human studies on nonoccupational PEP are observational in nature and limited in sample size, available data involving men who have sex with men (MSM) suggest nonoccupational PEP reduces HIV transmission.[15,16] In addition, a feasibility study in San Francisco demonstrated medical providers could appropriately identify and provide recommended nonoccupational PEP to persons exposed to HIV via sexual contact or through injecting drugs.[17] In a separate San Francisco study, investigators reported HIV seroconversion among 7 of 702 (1%) persons who received nonoccupational PEP for a potential sexual or injection drug use exposures to HIV, but only 3 of the seroconversions likely represented true nonoccupational PEP “failure”.[6] Available data from other reports of HIV transmission in persons who received nonoccupational PEP suggest that most HIV transmissions resulted from poor medication adherence, or from exposures to HIV that occurred after completing the PEP regimen.[18,19,20,21] In addition, one failure occurred in a 40-year-old woman in France who started nonoccupational PEP more than 72 hours after a sexual exposure.[22] Multiple studies involving sexual assault survivors have demonstrated very low HIV...
transmission rates, even though persons receiving nonoccupational PEP following sexual assault often have poor medication adherence.[23,24,25,26,27]

### Rationale for Nonoccupational PEP in Persons Who Inject Drugs

New HIV infections among persons who inject drugs can occur directly from the injection drug use or it may involve coexisting sexual activities associated with increased HIV transmission risk, such as condomless sex, sex with multiple partners, and transactional sex.[28,29] Certain circumstances could arise whereby a person who injects drugs and normally uses safe injection practices has an HIV risk exposure. The use of nonoccupational PEP after an at-risk injection drug use exposure may have different efficacy compared with use after a sexual exposure, since the route and HIV inoculum differ in these two situations. Unfortunately, very limited data exist regarding the practical applications or efficacy of nonoccupational PEP among persons who inject drugs. Of interest, however, one case report described a patient who inadvertently received a large-volume transfusion of blood from a person with HIV, but early initiation of PEP prevented transmission of HIV.[30] Based on this, one might extrapolate that nonoccupational PEP could be effective following injection drug use-related exposures to HIV, particularly if started early.[31]
Evaluation for Nonoccupational PEP

Multiple factors influence the risk of HIV transmission in nonoccupational exposures to HIV. The initial evaluation of persons seeking care after potential nonoccupational exposures to HIV requires gathering information to determine whether nonoccupational PEP is indicated (Figure 3).[2] The initial evaluation, as recommended in the 2016 Nonoccupational PEP Guidelines, should address the following: (1) the HIV status of the potentially exposed person, (2) information related to the source person’s HIV status, (3) details regarding the type of exposure involved, (4) timing and frequency of the exposure(s), and (5) any available information related to antiretroviral therapy taken by the source patient if they are known to be infected with HIV.[2] In addition, the initial history intake should include the following:

- **Information on HIV Status of Person Potentially Exposed to HIV:** The first step in evaluating the exposure is to determine the HIV status of the person seeking medical care. Persons with established HIV infection should receive long-term continuous antiretroviral therapy, not a 28-day course of nonoccupational PEP. In the setting of a nonoccupational exposure to HIV, the HIV status of the exposed person should be determined as soon as possible, although HIV testing should not necessarily delay nonoccupational PEP initiation.[2]

- **Information Related to Source Person’s HIV Status:** As part of the initial exposure evaluation, it is also important to determine whether the source person has HIV infection (or, if the status of the source is unknown, whether they are likely to have HIV infection). Often, the HIV status of the source person is not known (and not obtainable). The 2016 Nonoccupational PEP Guidelines recommend using nonoccupational PEP when the source person is known to have HIV infection; if the HIV status of the source person is not known, the recommendation is to evaluate on a case-by-case basis, ideally in consultation with an expert.[2] When asking about information related to the source, it is important to ask whether other recent exposures occurred with this source (and/or other sources), as recent additional HIV-related exposures could potentially confound management decisions. Nevertheless, repeated exposures do not negate the need to assess whether the most recent exposure warrants nonoccupational PEP.

- **Determination of Risk Related to Exposure:** The other key element of the initial nonoccupational PEP evaluation is to determine whether the exposure confers actual risk for HIV transmission (Table 1).[2,31] Nonoccupational PEP should only be used in the setting of “substantial risk for HIV acquisition”, defined as contact involving an area of the body known to be associated with HIV acquisition (vagina, rectum, eye, mouth or other mucous membranes, non-intact skin, or percutaneous needlestick injuries) with an infectious body fluid (e.g. blood, semen, vaginal secretions, rectal secretions, breast milk, or any body fluid visibly contaminated with blood). The risk of HIV transmission associated with nonoccupational exposures varies considerably by the type of sexual exposure, with receptive anal intercourse conveying the highest sexual risk.[2,31,32,33,34] Similarly, mucosal disruption in either the source person or the exposed person (as might occur with traumatic intercourse including sexual assault, or in the presence of ulcerative genital disease) increases risk of sexual HIV transmission: condom use markedly lowers the risk of transmission (Table 2).[31]

- **Timing of Risk Exposure:** It is important to determine the timing of exposure in persons seeking nonoccupational PEP. Available data suggest that PEP may not be effective if initiated beyond 72 hours after the exposure. Furthermore, PEP may not be the optimal long-term HIV prevention method for individuals who engage in activities involving frequent, recurrent HIV exposures, such as sex with an HIV-serodifferent sex partner without consistently using condoms or regularly sharing needles or equipment with injecting partners; these individuals should receive intensive risk reduction counseling and may instead be better suited for PrEP.[2]

- **Source Person Antiretroviral Treatment Information:** If a source person is known to have HIV and takes antiretroviral medications, the medical provider should determine what medication the source takes, the most recent HIV RNA level, and if the source person has
developed resistance to any antiretroviral medications. The risk of HIV transmission is higher if the source person has advanced HIV disease or high HIV RNA levels.[35]
Indications for Initiating Nonoccupational PEP

Based on the 2016 Nonoccupational PEP Guidelines, if the following criteria are met, a 28-day course of antiretroviral medications for nonoccupational PEP is recommended:[2]

- A person has had a nonoccupational exposure to blood, genital secretions, or other potentially infectious body fluids from a person known to be infected with HIV (if the HIV status of the source person is unknown, the recommendation is to evaluate on a case-by-case basis, ideally in consultation with an expert), and
- The exposure represents a substantial risk for HIV transmission, and
- The person seeks care within 72 hours of exposure

Note: Administering nonoccupational PEP with the goal of transitioning to PrEP can be considered beyond the 72-hour window in cases where multiple recent exposures to HIV have occurred more than 72 hours prior, but the most recent exposure occurred within the 72-hour window.

Challenges When Evaluating Whether to Initiate Nonoccupational PEP

A number of challenges may arise during the initial evaluation of persons for nonoccupational PEP after sexual and other nonoccupational exposures that complicate decisions regarding whether to initiate antiretroviral medications for nonoccupational PEP.

- With some exposures, the person seeking help may not actually know the HIV status (or any other information) of the source person.
- Individuals often present for care more than 72 hours after the exposure.
- The exposure that brings the person in for medical attention may not consist of an isolated event, but instead may be among multiple recent potential HIV exposure events; if this is the case, it is still necessary to assess the need for nonoccupational PEP for any significant exposures occurring within the last 72 hours.
- Certain sexual exposure events may involve concomitant exposures to other sexually transmitted pathogens, or hepatitis viruses with injection drug use (or percutaneous needlestick) exposures.
- Some nonoccupational PEP cases involve persons recently sexually assaulted, which can involve additional medicolegal concerns and complications.[6,36]
- From a practical perspective, insurers may not cover the cost of the 28-day course of nonoccupational PEP, or they may only partially cover the cost.
Recommended Therapy for Nonoccupational PEP

Preferred and Alternative Antiretroviral Regimens

The 2016 Nonoccupational PEP Guidelines recommend using a three-drug combination in all cases when nonoccupational PEP is indicated.[2] The preferred nonoccupational PEP regimen for adults (and adolescents age 13 years and older) who have a baseline creatinine clearance of at least 60 mL/min consists of the fixed dose combination of tenofovir DF-emtricitabine combined with either raltegravir or dolutegravir (Table 3).[2] The recommended alternative regimen is darunavir boosted with ritonavir plus tenofovir DF-emtricitabine.[2] For adults and adolescents who have a baseline creatinine clearance less than 60 mL/min, the preferred and alternative regimens listed above should be modified by replacing tenofovir DF-emtricitabine with zidovudine and lamivudine; in this situation, the fixed-dose combination zidovudine-lamivudine should not be given so that doses of zidovudine and lamivudine can be adjusted individually based on renal impairment.[37]

Medications Not Recommended

The antiretroviral medication abacavir, which is commonly used to treat HIV infection, is not recommended for nonoccupational PEP. Prior to receiving abacavir, all patients should have HLA-B*5701 testing; the HLA-B*5701 test is used to predict abacavir hypersensitivity, a potentially fatal reaction. The need to immediately administer nonoccupational PEP antiretroviral medications does not allow sufficient time to obtain results from HLA-B*5701 testing. In addition, the use of tenofovir alafenamide-emtricitabine is also not recommended at this time as an alternative to tenofovir DF-emtricitabine, primarily due to lack of data with the use of tenofovir alafenamide-emtricitabine for PEP. Further, the advantages of tenofovir alafenamide-emtricitabine over tenofovir DF-emtricitabine when used for long-term treatment of persons with HIV (lower rates of nephrotoxicity and osteopenia) are not generally relevant when only prescribing a short 28-day regimen for nonoccupational PEP. The use of nevirapine for nonoccupational PEP is strongly contraindicated since the use of nevirapine in occupational PEP was associated with a significant risk of life-threatening hepatotoxicity.[38]

Nonoccupational PEP Medication Studies

The following summarizes major studies involving more contemporary three-drug antiretroviral regimens for nonoccupational PEP.

- **Dolutegravir plus Tenofovir DF-Emtricitabine (Sydney Study):** In an open-label, single-arm study, investigators from Sydney, Australia, investigators enrolled 100 gay and bisexual men to receive dolutegravir plus tenofovir DF-emtricitabine for 28 days for nonoccupational PEP; the regimen was well tolerated, adherence levels were very high (98%), completion rates (90%) were high, and no HIV seroconversions occurred; elevations in alanine aminotransferase occurred in 22% of the participants, but none developed clinical hepatitis.[39]

- **Elvitegravir-Cobicistat-Tenofovir DF-Emtricitabine (Fenway Health Study):** In an open-label, single-arm study, investigators from Fenway Health Clinic in Boston, Massachusetts enrolled adults (98% men) to receive the fixed-dose single tablet elvitegravir-cobicistat-tenofovir DF-emtricitabine for 28 days for nonoccupational PEP.[40] The 28-day nonoccupational PEP regimen was completed as prescribed by 71% of the participants and the regimen was moderately well tolerated, with the most common adverse effects reported as gastrointestinal discomfort (42%), diarrhea (38%), nausea/vomiting (28%), and fatigue (28%).[40] At the day 90 follow-up visit, none had HIV seroconversion.[40]

- **Raltegravir plus Tenofovir DF-Emtricitabine (Fenway Health Study):** The Fenway Health Clinic in Boston, Massachusetts reported their experience with a 28-day course of the three-drug regimen raltegravir plus tenofovir DF-emtricitabine as nonoccupational PEP in 100
adult men, most of whom were men who have sex with men (MSM); this regimen was completed as prescribed by 57% of the participants and was relatively well tolerated, with nausea/vomiting (27%) and diarrhea (21%) the most common reported side effects.[41] Among the 85 men who were evaluable at 3 months, none had acquired HIV.[41]

- **Raltegravir plus Tenofovir DF-Emtricitabine (Sydney Study):** Investigators from Sydney, Australia performed an open-label, prospective study that enrolled 86 MSM to receive a 28-day course of raltegravir plus tenofovir DF-emtricitabine for nonoccupational PEP; the regimen was well tolerated, adherence levels were high (89%), completion rates were very high (92%), and no HIV seroconversions occurred.[42] In this study, investigators also enrolled 34 men to receive the two-drug regimen tenofovir DF-emtricitabine and no HIV seroconversions occurred in this group as well.[42]

- **Rilpivirine-Tenofovir DF-Emtricitabine (Sydney Study):** Investigators from Sydney, Australia performed an open-label, single-arm, prospective study that enrolled 100 adult MSM to receive a 28-day course of the fixed-dose tablet rilpivirine-tenofovir DF-emtricitabine for nonoccupational PEP; the regimen was well tolerated, adherence rates were very high (98%), completion rates very high (92%), and no HIV seroconversions had occurred among the 70 men who had completed the 12-week follow-up visit.[43] The most commonly reported adverse effects were fatigue (34%) and nausea (23%).

### Duration of Therapy

The 2016 Nonoccupational PEP Guidelines recommend individuals who initiate antiretroviral therapy for nonoccupational PEP should complete a 28-day course.[2] Studies involving macaques have shown that PEP given for 28 days is more effective than 10 days, which is more effective than 3 days.[12] In addition, available data and experience with occupational postexposure prophylaxis support the use of a 28-day regimen.[7,44] From a conceptual standpoint, it is believed that PEP, in some instances, halts a very early and limited HIV infection, rather than truly preventing any cell in the body from becoming infected with HIV. Thus, early initiation of nonoccupational PEP, when combined with a 28-day duration of therapy, is believed adequate to minimize tissue involvement and contain any local HIV replication while allowing sufficient time for localized immune responses to clear out the limited HIV infection.

### Consideration of HIV Resistance in Source Person

If medical information is available regarding a source person known to be HIV-positive, the choice of nonoccupational PEP regimen should take into account the source person’s antiretroviral medication history, most recent HIV RNA level, and prior resistance testing results. In the event the source person has possible or known antiretroviral drug resistance, expert consultation should be obtained to determine the optimal nonoccupational PEP regimen for the individual exposed to HIV.

### Recommendation if Source Person Has an Undetectable HIV RNA Level

Multiple studies have shown that persons with HIV who consistently maintain undetectable serum HIV RNA levels do not sexually transmit HIV, even with condomless sex.[45,46,47] These studies led to the concept of undetectable equals untransmitable, which is commonly referred to as “U=U”.[48] The widespread support of the U=U concept has occurred only very recently and after the publication of the 2016 Nonoccupational PEP Guidelines. As might be expected, the 2016 Nonoccupational PEP Guidelines do not specifically address whether or not to initiate nonoccupational PEP if the exposure involves a source person with a recent undetectable HIV RNA level.[2] Thus, at the present time, there are no clear recommendations regarding use of nonoccupational PEP following sexual exposures to source persons with HIV who have consistently suppressed HIV RNA levels, but many experts would base their recommendation on the reliability of the information and documentation of persistently suppressed HIV RNA levels in the source persons with HIV. This issue will need to be addressed in future nonoccupational PEP guidelines.
Recommendation if Person Exposed to HIV is Receiving PrEP

Persons who are receiving HIV preexposure prophylaxis (PrEP) with daily tenofovir DF-emtricitabine and are adhering to a daily PrEP regimen do not need nonoccupational PEP following an exposure to HIV.[5] If an individual is receiving PrEP, but is not consistently taking PrEP on a daily basis, nonoccupational PEP might be indicated following an HIV exposure.[5] In this situation of sporadic PrEP use, the decision to initiate nonoccupational PEP should be made on a case-by-case basis, and ideally with the help of expert clinical consultation.
Expert Consultation for Nonoccupational PEP

Indications for Obtaining Expert Consultation

The 2016 Nonoccupational PEP Guidelines provide recommendations for scenarios that warrant expert consultation for nonoccupational PEP related to nonoccupational HIV exposure events.[2] Expert consultation is recommended in any of the following situations:

- The health care worker has limited experience with prescribing antiretroviral medications, or
- The individual exposed to HIV is a woman who is pregnant or breastfeeding, or
- The exposure event involves a child or adolescent, or
- The individual needing nonoccupational PEP has renal dysfunction, or
- The source person has known or suspected antiretroviral resistance.

PEPline Expert Consultation

Expert consultation for these issues and any other guidance on Nonoccupational PEP can be obtained by calling the National Clinician Consultation Center’s Post-Exposure Prophylaxis PEPline at 888-448-4911; this service is for health care professionals and is available 9 a.m.-8 p.m. Eastern Time (ET) Monday through Friday and 11 a.m.-8 p.m. ET on weekends and holidays. The National Clinician Consultation Center cannot accept calls from unknown numbers, so they ask that callers please unblock their phone prior to calling the PEPline.
Laboratory Testing for Source and Exposed Persons

The 2016 Nonoccupational PEP Guidelines provide recommendations for baseline laboratory studies (for the source and the person exposed to HIV), as well as a schedule of laboratory tests for monitoring the person exposed to HIV (Table 4).[2] Initial HIV testing of the individual exposed to HIV should consist of an HIV-1/2 antigen-antibody immunoassay, or HIV antibody testing if HIV antigen-antibody testing is not available.[2] Oral point-of-care HIV tests are not recommended, primarily due to their poor sensitivity in diagnosing acute or very recent HIV infection. Although it is very important to confirm the negative HIV status of the individual presenting for nonoccupational PEP, most experts do not advocate ordering an HIV RNA level (on the exposed person) unless they have signs or symptoms that suggest an acute HIV infection. In addition, persons undergoing evaluation for nonoccupational PEP should be instructed about the signs and symptoms associated with acute HIV infection and asked to return for evaluation if these occur during or after nonoccupational PEP.[2]

Baseline Laboratory Evaluation

The baseline laboratory evaluation should include the following tests for the individual exposed to HIV:[2]

- HIV-1/2 antigen-antibody immunoassay (or HIV antibody testing if the HIV-1/2 antigen-antibody immunoassay is not available)
- Serologic testing for hepatitis B virus (HBV) and hepatitis C virus (HCV)
- Testing for sexually transmitted infections, including serologic testing for syphilis and site-specific screening for chlamydia and gonorrhea
- Pregnancy test and emergency contraception, if indicated
- Serum creatinine (for calculating estimated creatinine clearance)
- Hepatic aminotransferase levels

Follow-Up Laboratory Studies

The 2016 Nonoccupational PEP Guidelines recommend the following laboratory tests for all persons who seek care for nonoccupational PEP.[2]

- **HIV Testing**: Follow-up HIV testing at 4 to 6 weeks after exposure and at 3 months after exposure to determine if HIV transmission has occurred.[2] Ideally, the HIV-1/2 antigen-antibody immunoassay should be used, but HIV antibody testing is acceptable if the HIV-1/2 antigen-antibody immunoassay is not available. If the exposed person becomes HCV antibody positive as a result of the original exposure, an additional HIV test should be conducted at 6 months after the exposure.[2]
- **HCV testing**: Follow-up HCV testing for susceptible exposed persons is recommended at baseline and at 6 months, but not at 4 to 6 weeks or 3 months.[2]
- **HBV Testing**: For exposed persons not immune to HBV at baseline, providers should ascertain HBV status of the source if possible and administer HBV postexposure prophylaxis as indicated. Follow-up HBV testing should then be conducted at 6 months.[2]
Nonoccupational PEP in Special Populations and Circumstances

The 2016 Nonoccupational PEP Guidelines identify additional considerations for certain special populations as outlined below.[2]

Pregnant Women and Women of Childbearing Potential

The recommended regimens for nonoccupational PEP are generally considered safe during pregnancy. Several antiretroviral medications are considered problematic for use in pregnancy.

- **Stavudine plus Didanosine**: The combination of stavudine and didanosine should not be used due to the risk of lactic acidosis.
- **Indinavir**: Indinavir should be avoided due to the risk of nephrolithiasis.
- **Nevirapine**: The medication nevirapine should never be used for PEP due to risk of severe hepatotoxicity. Furthermore, because the efficacy of hormonal contraception can be altered by some antiretroviral medications, women using such methods should be advised to use a secondary form of contraception (i.e. barrier methods) while taking nonoccupational PEP.
- **Darunavir plus Ritonavir**: Among the preferred and alternative recommended regimens for nonoccupational PEP, only darunavir boosted with ritonavir causes significant drug interactions with oral contraceptives.

Specific information is not provided in the guidelines regarding use of antiretroviral regimens for nonoccupational PEP in women who are breastfeeding. Expert consultation should be obtained for these cases.

Children and Adolescents

In many pediatric/adolescent nonoccupational PEP cases, expert consultation will be necessary. In the 2016 Nonoccupational PEP Guidelines, the preferred regimen for children aged 4 weeks to less than 2 years of age is zidovudine and lamivudine plus either raltegravir or lopinavir-ritonavir, with all medications given orally and dose-adjusted for age and weight. For children aged 2 to 12 years, the preferred regimen is tenofovir DF plus emtricitabine plus raltegravir (dosed according to age and weight).[2] For children age 13 and older with normal renal function, the adult and adolescent preferred and alternative nonoccupational PEP regimens can be used.[2]

Sexual Assault Survivors

Survivors of sexual assault may be less likely to seek care for nonoccupational PEP and may require additional psychological support, clinical follow-up, and adherence counseling. Testing and treatment for other sexually transmitted infections, emergency contraception evaluation, and supportive counseling are also highly recommended in cases of sexual assault.

Inmates

The 2016 Nonoccupational PEP Guidelines recommend that correctional facilities establish HIV prevention programs. Elements should include confidential and voluntary HIV testing, risk reduction services, and nonoccupational PEP protocols. The Federal Bureau of Prisons published a clinical practice guideline based on the 2005 CDC Nonoccupational PEP guidelines and recommends that each facility develop its own protocol, but the CDC recommends that the most updated guidelines be used whenever possible.[2]

HIV Postexposure Prophylaxis in Mass Casualty Events

In response to concerns for a potential mass casualty event within the United States, the Centers for
Disease Control and Prevention convened a working group to address management of blood-borne pathogen exposure in persons who are injured in bombings and other mass-casualty events, as well as for emergency responders in these catastrophic events. This particular situation does not fall neatly under the guidelines for either occupational or nonoccupational PEP. Accordingly, a separate document was published in 2008, which specifically addresses postexposure prophylaxis for HIV, HBV, HCV, and tetanus in the setting of mass casualties.[49] Postexposure prophylaxis for HIV is not routinely indicated for persons exposed to blood or tissue in bombings or mass-casualty events. In certain situations, however, nonoccupational PEP might be indicated if the risk for HIV exposure was determined to be high, such as with bombing of a research facility that has blood specimens obtained from persons with HIV, or culture vials growing HIV. If postexposure prophylaxis is indicated, the same principles of timing, laboratory testing, and antiretroviral medication selection should apply.
Initial Medication Prescription and Follow-up after Evaluation

Initial Medication Prescription and Timely Follow-Up

The 2016 Nonoccupational PEP Guidelines recommend that medical providers should consider giving an initial prescription for 3 to 7 days of antiretroviral medication (i.e. a starter pack) or provide a prescription for an entire 28-day course. Ideally, at the initial visit, the facility would supply the starter pack medication or the full 28-day supply of medication, both to minimize any delay in receiving the first dose and to address any barriers that could prohibit the patient from filling the prescription. In addition, prior to leaving the facility, the person evaluated for nonoccupational PEP should have an early follow-up visit scheduled to assess adherence to nonoccupational PEP, monitor for toxicity, and provide any additional counseling or education that might be needed.[2] If the individual receives only a 3- to 7-day supply of antiretroviral medication, coordination of timely follow-up is essential to ensure they do not run out of medication.

Challenges to Follow-Up

One study reported significant patient attrition between initial emergency department visit for nonoccupational PEP and first follow-up clinic appointment.[36] Only about half of nonoccupational PEP patients attended their follow-up appointment, and less than a quarter of those initially started on nonoccupational PEP completed the full 28-day regimen.[36] Older age and self-pay status predicted lower rates of follow-up and poor adherence; women were less likely than men to complete the full course of nonoccupational PEP.

HIV Prevention Counseling

Patients who are evaluated for nonoccupational PEP should receive HIV prevention counseling. This includes counseling on risk-reduction behaviors (e.g. using a barrier method with sex partners, not sharing equipment used to inject drugs) and referral to local community resources, if available). At follow-up visits, health care providers should assess for ongoing risk behaviors, provide additional counseling, and connect patients with services as needed.
Transitioning to Preexposure Prophylaxis

Preexposure Prophylaxis (PrEP)

Multiple studies conducted in recent years have demonstrated that daily use of PrEP (most commonly given as a fixed-dose combination of tenofovir DF-emtricitabine) protects against HIV infection among heterosexuals at high risk for HIV acquisition,[50,51,52,53] MSM and transgender women,[51,54,55] and persons who inject drugs.[56] These studies show that PrEP is a feasible and effective HIV prevention strategy that may be more protective than repeated courses of nonoccupational PEP.

Transitioning from Nonoccupational PEP to PrEP

Many individuals who present for nonoccupational PEP following a sexual or injection drug use exposure may be excellent candidates for PrEP, particularly if they report ongoing activities associated with increased risk for HIV acquisition. The PrEP Clinical Practice Guideline recommends that persons receiving a 28-day course of nonoccupational PEP should be evaluated for transitioning to PrEP if (1) they have repeatedly sought nonoccupational PEP, and/or (2) they have frequent, recurrent exposures to HIV that would require sequential or near-continuous courses of nonoccupational PEP.[5]

Persons taking nonoccupational PEP who are deemed appropriate candidates to receive PrEP can immediately transition to PrEP after completing the 28-day course of nonoccupational PEP.[5] The transition from nonoccupational PEP to PrEP should occur without a gap, but prior to the transition, documentation of HIV-negative status should be obtained, preferably with an HIV-1/2 antigen-antibody test.[5] To practically achieve a transition without allowing a gap, the recommended 4- to 6-week follow-up HIV testing should be performed several days earlier than normal (several days prior to the end of the 28-day course of nonoccupational PEP). The approach of immediate transition will diminish the risk of HIV acquisition among individuals who have significant ongoing risk. There is no definitive evidence that receipt of nonoccupational PEP delays HIV seroconversion.[2]

Expert Consultation for Transitioning from PEP to PrEP

In July 2012, the FDA approved tenofovir DF-emtricitabine for use as PrEP in individuals with ongoing high risk of HIV acquisition. The PrEP Clinical Practice Guideline recommend that individuals with frequent and multiple exposures to HIV, such as MSM who report condomless anal sex with a partner who has HIV infection, should not be managed with repeated courses of nonoccupational PEP, but instead strongly considered for PrEP. Clinicians with questions about PrEP can call the National Clinician Consultation Center's Pre-Exposure Prophylaxis PrEPline at 855-448-7737 for expert consultation. The PrEPline is available Monday – Friday, 9 a.m.–8 p.m. ET.
Concerns with Nonoccupational Postexposure Prophylaxis

Toxicity of Antiretroviral Therapy

Initial concerns about severe side effects and pharmacological toxicities in otherwise healthy persons have been ameliorated by the use of less toxic, well-tolerated antiretroviral agents.\[57]\n
Use of Nonoccupational PEP and Drug Resistance Mutations

Selection of drug-resistant HIV can potentially result from the use of nonoccupational PEP. The development of HIV drug resistance during receipt of nonoccupational PEP is highly unlikely because few individuals receiving nonoccupational PEP will develop HIV infection and emergence of new resistant strains would be uncommon following a 28-day course of a potent three-drug antiretroviral nonoccupational PEP regimen. Nonoccupational PEP should not be withheld due to theoretical concerns about the potential selection of drug-resistant HIV.

Use of Nonoccupational PEP and Changes in Sexual Activities

Some clinicians have expressed concern that availability of nonoccupational PEP could theoretically result in a change in sexual activity that could increase risk for acquiring HIV.\[3]\ Multiple studies involving MSM have shown that sexual risk activity does not significantly change among individuals who receive nonoccupational PEP.\[17, 58, 59]\ In one study in San Francisco, 72% of nonoccupational PEP recipients reported a decrease in sexual risk behavior over the next 12 months, 13% reported no change, and 14% reported an increase in sexual risk behavior; in that study, 17% of participants requested a repeat course of nonoccupational PEP within the following year.\[58]\ In a separate study, investigators reported that 12% of individuals requested a second course of nonoccupational PEP within 6 months of the initial nonoccupational PEP course.\[17]\ In a study that specifically examined whether the knowledge of nonoccupational PEP availability would lead to an increase in risk behavior, there was no difference in risk behavior among men who have sex with men exclusively and who knew about nonoccupational PEP versus those who had never heard of it.\[59]\ Taken together, the available data suggest that receipt of nonoccupational PEP (consisting of both antiretroviral prophylaxis and counseling) does not increase HIV risk behaviors nor has it been linked to an increase in HIV prevalence. Of note, no studies have been published that have examined changes in injection drug use patterns among persons who inject drugs who have received a course of nonoccupational PEP.
Summary Points

- Nonoccupational PEP represents an important tool in the arsenal of HIV prevention but remains an underutilized strategy.
- As a whole, there is a body of data that supports use of nonoccupational PEP and indicates it may reduce the risk of HIV infection after nonoccupational exposures to HIV.
- The use of antiretroviral medications for nonoccupational PEP is recommended for HIV-negative persons following an exposure that has a substantial risk for HIV acquisition, if started within 72 hours of the exposure.
- A 28-day course of three-drug antiretroviral medications is recommended for nonoccupational PEP.
- The preferred nonoccupational PEP regimen for adult and adolescents aged 13 years and older who have a baseline creatinine clearance of at least 60 mL/min consists of the fixed-dose combination of tenofovir DF-emtricitabine plus either raltegravir or dolutegravir.
- The person exposed to HIV should have baseline laboratory studies that include HIV testing, screening for sexually transmitted infections, serum creatinine, aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels, testing for viral hepatitis, and a pregnancy test (if indicated).
- Expert consultation should be sought for all situations that fall outside the scope of the guidelines, including situations when exposure to drug-resistant HIV has occurred. Consultation can be obtained through local expertise (if available) or by calling the National Clinician Consultation Center’s Post-Exposure Prophylaxis Hotline (PEPline) at 888-448-4911.
- All persons seeking care for nonoccupational PEP should have follow-up HIV testing at 4 to 6 weeks and again at 3 months to determine if HIV infection has occurred. The HIV-1/2 antigen-antibody immunoassay is preferred for HIV testing. Additional HIV testing should be conducted at 6 months if the exposure resulted in HCV transmission.
- Nonoccupational PEP has not been linked to high rates of adverse side effects, selection of resistant HIV virus, or increases in higher risk sexual activity behaviors.
- Most individuals who seek nonoccupational PEP should be evaluated as potential candidates to receive PrEP following completion of nonoccupational PEP. The transition from occupational PEP to PrEP, if warranted, should occur without a gap.
Citations


References


- Centers for Disease Control and Prevention, U.S. Department of Health and Human Service. Update: Interim Statement Regarding Potential Fetal Harm from Exposure to Dolutegravir—Implications for HIV Post-exposure Prophylaxis (PEP). [CDC] -


- Donnell D, Mimiaga MJ, Mayer K, Chesney M, Koblin B, Coates T. Use of non-occupational post-exposure prophylaxis does not lead to an increase in high risk sex behaviors in men who
have sex with men participating in the EXPLORE trial. AIDS Behav. 2010;14:1182-9. [PubMed Abstract]


- Panel on Antiretroviral Therapy and Medical Management of Children Living with HIV. Guidelines for the use of antiretroviral agents in pediatric HIV infection. September 12, 2019 [AIDSinfo].


Figures

Figure 1 Tenofovir for Postexposure Prophylaxis following SIV-1 Inoculation of Macaques

In this study, investigators inoculated 24 macaques with simian immunodeficiency virus (SIV) and then instituted various postexposure prophylaxis regimens with tenofovir (PMPA), which is (R)-9-(2-phosphonylmethoxypropyl)adenine.


<table>
<thead>
<tr>
<th>Study Hour</th>
<th>0</th>
<th>24</th>
<th>48</th>
<th>72</th>
</tr>
</thead>
<tbody>
<tr>
<td>SIV Inoculation</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Start PEP</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Study Features**
- N = 24 macaques
- Randomized to 6 treatment arms
- SIV inoculated intravenously
- SIV dose 10bx 50% infective dose
- PEP started at 24, 48, or 72 hours
- PEP duration: 3, 10, or 28 days
- PEP regimen: Tenofovir (PMPA) SQ
- Analyzed for antibody and viremia

Tenofovir (PMPA) = (R)-9-(2-phosphonylmethoxypropyl)adenine
Figure 2 SIV Transmission Based on Timing of Initiation and Duration of PEP

Among animals that received placebo, all became infected with simian immunodeficiency virus (SIV). The most effective tenofovir prevention regimen consisted of early initiation of tenofovir (at 24 hours) and long duration of postexposure prophylaxis (28 days).

Figure 3 Algorithm for Evaluation and Treatment of possible nonoccupational HIV exposures


**Substantial risk for HIV acquisition**

- ≤72 hours since exposure
  - Source patient known to be HIV positive: nPEP recommended
  - Source patient of unknown HIV status: Case-by-case determination

**Negligible risk for HIV acquisition**

- ≥73 hours since exposure
  - nPEP Not recommended

---

**Substantial Risk for HIV Acquisition**

*Exposure of*
- Vagina, rectum, eye, mouth or other mucous membrane, nonintact skin, or precutaneous contact

*With*
- Blood, semen, vaginal secretions, rectal secretions, breast milk, or any body fluid that is visibly contaminated with blood

*When*
- The source is known to be HIV-positive

**Negligible Risk for HIV Acquisition**

*Exposure of*
- Vagina, rectum, eye, mouth or other mucous membrane, intact or nonintact skin, or percutaneous contact

*With*
- Urine, nasal secretions, saliva, sweat, or tears if not visibility contaminated with blood

*Regardless*
- Of the known or suspected HIV status of the source
Table 1.

Estimated Per-Act Probability of Acquiring HIV from an Infected Source, by Exposure Act

<table>
<thead>
<tr>
<th>Exposure Type</th>
<th>Rate for HIV Acquisition per 10,000 Exposures</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Parenteral</strong></td>
<td></td>
</tr>
<tr>
<td>Blood transfusion</td>
<td>9,250</td>
</tr>
<tr>
<td>Needle sharing during injection drug use</td>
<td>63</td>
</tr>
<tr>
<td>Percutaneous (needlestick)</td>
<td>23</td>
</tr>
<tr>
<td><strong>Sexual</strong></td>
<td></td>
</tr>
<tr>
<td>Receptive anal intercourse</td>
<td>138</td>
</tr>
<tr>
<td>Insertive anal intercourse</td>
<td>11</td>
</tr>
<tr>
<td>Receptive penile-vaginal intercourse</td>
<td>8</td>
</tr>
<tr>
<td>Insertive penile-vaginal intercourse</td>
<td>4</td>
</tr>
<tr>
<td>Receptive oral intercourse</td>
<td>Low</td>
</tr>
<tr>
<td>Insertive oral intercourse</td>
<td>Low</td>
</tr>
<tr>
<td><strong>Other</strong></td>
<td></td>
</tr>
<tr>
<td>Biting</td>
<td>Negligible</td>
</tr>
<tr>
<td>Spitting</td>
<td>Negligible</td>
</tr>
<tr>
<td>Throwing body fluids (including semen or saliva)</td>
<td>Negligible</td>
</tr>
<tr>
<td>Sharing sex toys</td>
<td>Negligible</td>
</tr>
</tbody>
</table>

*Factors that may increase the risk of HIV transmission include sexually transmitted diseases, acute and late-stage HIV infection, and high viral load. Factors that may decrease the risk include condom use, male circumcision, antiretroviral treatment, and preexposure prophylaxis (PrEP). None of these factors are accounted for in the estimates presented in the table.

^HIV transmission through these exposure routes is technically possible but unlikely and not well documented.

Source:

Table 2.

**Relative Risks of Factors that Alter Per-Act HIV Transmission Risk for Sexual Exposures**

<table>
<thead>
<tr>
<th>Cofactor</th>
<th>Relative Risk</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Factors that Increase Transmission Probability</strong></td>
<td></td>
</tr>
<tr>
<td>High plasma viral load (log_{10} copies/mL)</td>
<td>2.89</td>
</tr>
<tr>
<td>Genital ulcer disease</td>
<td>2.65</td>
</tr>
<tr>
<td>Acute versus asymptomatic stage of disease</td>
<td>7.25</td>
</tr>
<tr>
<td>Late versus asymptomatic stage of disease</td>
<td>5.81</td>
</tr>
<tr>
<td><strong>Factors that Decrease Transmission Probability</strong></td>
<td></td>
</tr>
<tr>
<td>Use of antiretroviral medications by HIV-infected partner</td>
<td></td>
</tr>
<tr>
<td>Early versus delayed treatment</td>
<td>0.04</td>
</tr>
<tr>
<td>Received treatment versus no treatment</td>
<td>0.08</td>
</tr>
<tr>
<td>Preexposure Prophylaxis of HIV-uninfected partner</td>
<td></td>
</tr>
<tr>
<td>Among heterosexual couples</td>
<td>0.29</td>
</tr>
<tr>
<td>Among men who have sex with men</td>
<td>0.56</td>
</tr>
<tr>
<td>Among injection drug users</td>
<td>0.52</td>
</tr>
<tr>
<td>Condom use</td>
<td>0.20</td>
</tr>
<tr>
<td><strong>Male Circumcision (heterosexual partners)</strong></td>
<td></td>
</tr>
<tr>
<td>HIV-uninfected partner is male</td>
<td>0.50</td>
</tr>
<tr>
<td>HIV-uninfected partner is female</td>
<td>0.80</td>
</tr>
<tr>
<td><strong>Male circumcision (men who have sex with men)</strong></td>
<td></td>
</tr>
<tr>
<td>Insertive partner is HIV-uninfected partner</td>
<td>0.27</td>
</tr>
<tr>
<td>Receptive partner is HIV-uninfected</td>
<td>1.2</td>
</tr>
</tbody>
</table>

*For a detailed description of data and factors used to generate estimates, see original table in article referenced below.

Source:

### Table 3. 2016 CDC and HHS Guidelines for Nonoccupational Exposure to HIV

**Preferred and Alternative 28-Day Regimens for Nonoccupational PEP<sup>a,b</sup>**

<table>
<thead>
<tr>
<th>Adults and adolescents aged ≥13 years, including pregnant women, with normal renal function (creatinine clearance ≥60 mL/min)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Preferred Regimens:</strong></td>
</tr>
<tr>
<td>• Raltegravir (400 mg twice daily) plus tenofovir DF-emtricitabine (300-200 mg once daily)</td>
</tr>
<tr>
<td>• Dolutegravir (50 mg once daily) plus tenofovir DF-emtricitabine (300-200 mg once daily)</td>
</tr>
<tr>
<td><strong>Alternative Regimen:</strong></td>
</tr>
<tr>
<td>• Darunavir (800 mg once daily) plus ritonavir (100 mg once daily) plus tenofovir DF-emtricitabine (300-200 mg once daily)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Adults and adolescents aged ≥13 years with renal dysfunction (creatinine clearance ≤59 mL/min)&lt;sup&gt;+&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Preferred Regimens:</strong></td>
</tr>
<tr>
<td>• Raltegravir (400 mg twice daily) plus zidovudine (dose adjusted) plus lamivudine (dose adjusted)</td>
</tr>
<tr>
<td>• Dolutegravir (50 mg once daily) plus zidovudine (dose adjusted) plus lamivudine (dose adjusted)</td>
</tr>
<tr>
<td><strong>Alternative Regimen:</strong></td>
</tr>
<tr>
<td>• Darunavir (800 mg once daily) plus ritonavir (100 mg once daily) plus zidovudine (dose adjusted) plus lamivudine (dose adjusted)</td>
</tr>
</tbody>
</table>

<sup>a</sup>These recommendations do not reflect current Food and Drug Administration-approved labeling for antiretroviral medications listed in this table.

<sup>b</sup>Ritonavir is used in clinical practice as a pharmacokinetic enhancer to increase the trough concentration and prolong the half-life of darunavir, lopinavir, and other protease inhibitors. Ritonavir is not counted as a drug directly active against HIV in the above “3-drug” regimens.

<sup>+</sup>The dose adjustments for zidovudine and lamivudine are made based on degree of renal function.

Source:

Table 4.

Nonoccupational PEP (nPEP)
Recommended Laboratory Monitoring of Source and Exposed Persons

<table>
<thead>
<tr>
<th>Test</th>
<th>Source</th>
<th>Baseline</th>
<th>4-6 Weeks after exposure</th>
<th>3 Months after exposure</th>
<th>6 months after exposure</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIV-1/2 Ag/Ab (or antibody testing if Ag/Ab test unavailable)</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>√</td>
<td>√</td>
</tr>
<tr>
<td>HIV-1/2 Ag/Ab (or antibody testing if Ag/Ab test unavailable)</td>
<td>4-6 Weeks</td>
<td>√</td>
<td>√</td>
<td>√</td>
<td>√</td>
</tr>
<tr>
<td>HIV-1/2 Ag/Ab (or antibody testing if Ag/Ab test unavailable)</td>
<td>3 Months</td>
<td>√</td>
<td>√</td>
<td>√</td>
<td>√</td>
</tr>
<tr>
<td>HIV-1/2 Ag/Ab (or antibody testing if Ag/Ab test unavailable)</td>
<td>6 months</td>
<td>√</td>
<td>√</td>
<td>√</td>
<td>√</td>
</tr>
<tr>
<td>Hepatitis B serology, including: hepatitis B surface antigen (HBsAg)</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis B surface antibody (anti-HBs)</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis B surface antibody (anti-HBs)</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis B surface antibody (anti-HBs)</td>
<td>6 months</td>
<td>√</td>
<td>√</td>
<td>√</td>
<td>√</td>
</tr>
<tr>
<td>Hepatitis C antibody test</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis C antibody test</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis C antibody test</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Hepatitis C antibody test</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Syphilis serology</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Syphilis serology</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Syphilis serology</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Syphilis serology</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Gonorrhea</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Gonorrhea</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Gonorrhea</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Gonorrhea</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Chlamydia</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Chlamydia</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Chlamydia</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Chlamydia</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Pregnancy</td>
<td>Baseline</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Pregnancy</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Pregnancy</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Pregnancy</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Serum creatinine (for calculating estimated creatinine clearance)</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Serum creatinine (for calculating estimated creatinine clearance)</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Serum creatinine (for calculating estimated creatinine clearance)</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Serum creatinine (for calculating estimated creatinine clearance)</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Alanine aminotransferase, aspartate aminotransferase</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Alanine aminotransferase, aspartate aminotransferase</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Alanine aminotransferase, aspartate aminotransferase</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>Alanine aminotransferase, aspartate aminotransferase</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>HIV RNA level</td>
<td>Baseline</td>
<td>√</td>
<td>√</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>HIV RNA level</td>
<td>4-6 Weeks</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>HIV RNA level</td>
<td>3 Months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
<tr>
<td>HIV RNA level</td>
<td>6 months</td>
<td>_</td>
<td>_</td>
<td>_</td>
<td>_</td>
</tr>
</tbody>
</table>

For all persons considered for or prescribed nPEP for any exposure

For all persons considered for or prescribed nPEP for sexual exposure

For persons prescribed:
- Tenofovir DF-emtricitabine + raltegravir
- Tenofovir DF-emtricitabine + dolutegravir

For all persons with HIV infection confirmed at any visit
<table>
<thead>
<tr>
<th>Test</th>
<th>Source</th>
<th>Exposed</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Baseline</td>
</tr>
<tr>
<td>HIV genotypic drug resistance test</td>
<td></td>
<td>Baseline</td>
</tr>
</tbody>
</table>

For all persons considered for or prescribed nPEP for any exposure

Abbreviations: Ag/Ab, antigen/antibody combination test; HIV, human immunodeficiency virus; nPEP, nonoccupational postexposure prophylaxis; tenofovir DF, tenofovir disoproxil fumarate.

- Any positive or indeterminate HIV antibody test should undergo confirmatory testing of HIV status.
- Only if hepatitis C infection was acquired during the original exposure; delayed HIV seroconversion has been seen in persons who simultaneously acquire HIV and hepatitis C infection.
- If exposed person susceptible to hepatitis B at baseline.
- If exposed person susceptible to hepatitis C at baseline.
- If determined to be infected with syphilis and treated, should undergo serologic syphilis testing 6 months after treatment.
- Testing for chlamydia and gonorrhea should be performed using nucleic acid amplification tests. For patients diagnosed with a chlamydia or gonorrhea infection, retesting 3 months after treatment is recommended.
  - For men reporting insertive vaginal, anal, or oral sex, a urine specimen should be tested for chlamydia and gonorrhea.
  - For women reporting receptive vaginal sex, a vaginal (preferred) or endocervical swab or urine specimen should be tested for chlamydia and gonorrhea.
  - For men and women reporting receptive anal sex, a rectal swab specimen should be tested for chlamydia and gonorrhea.
  - For men and women reporting receptive oral sex, an oropharyngeal swab should be tested for gonorrhea.
- If not provided presumptive treatment at baseline, or if symptomatic at follow-up visit.
- If woman of reproductive age, not using effective contraception, and with vaginal exposure to semen.

\[ e\text{CrCl} = \frac{(140 - \text{age}) \times \text{ideal body weight}}{\text{serum creatinine} \times 72} \times 0.85 \text{ for females}. \]

At first visit where determined to have HIV infection.

Source: