



Post-Exposure Prophylaxis (PEP) to Prevent HIV Infection

November 2021

Table 2: Preferred Post-Exposure Prophylaxis Regimens for Patients Who Weigh ≥40 kg [a,c]	
Preferred Regimen	Notes
<ul style="list-style-type: none"> Tenofovir disoproxil fumarate 300 mg/ emtricitabine 200 mg (TDF/FTC; Truvada) once per day or TDF 300 mg/lamivudine (TDF/3TC; Cimduo) 300 mg once per day <p>plus</p> <ul style="list-style-type: none"> Raltegravir (RAL; Isentress) 400 mg twice per day or RAL HD 1200 mg once per day [b] or Dolutegravir (DTG; Tivicay) 50 mg once per day 	<ul style="list-style-type: none"> DTG: If prescribed, discuss with individuals of childbearing capacity the small risk of teratogenicity in the first trimester, and counsel about the need for birth control while completing the 28-day PEP regimen; there is no elevated risk beyond the first trimester (see <i>Box 2: Use of Dolutegravir in Individuals of Childbearing Capacity</i>). <ul style="list-style-type: none"> Metformin dosing should be limited to 1 g by mouth per day when an individual is taking DTG concurrently. DTG: Magnesium- or aluminum-containing antacids may be taken 2 hours before or 6 hours after DTG; calcium- containing antacids or iron supplements may be taken simultaneously if taken with food. RAL: Magnesium- or aluminum-containing antacids are contraindicated; coadministration of calcium-containing antacids is not recommended with RAL HD. TDF: Requires dose adjustment for creatinine clearance (CrCl) <50 mL/min. Alternatively, another agent can be considered, in which case consultation with an experienced HIV care provider is advised. TDF/FTC and TDF/3TC: Dosing should be adjusted in patients with baseline CrCl <50 mL/min.
<p>a. All medications are taken by mouth for 28 days.</p> <p>b. RAL HD: May be prescribed for patients who weigh >40 kg; RAL HD should not be prescribed for pregnant individuals.</p> <p>c. Available alternative formulations and methods of administration:</p> <ul style="list-style-type: none"> 3TC: Acceptable to crush or split. Available as an oral solution (10 mg/mL). DTG: Acceptable to crush. FTC: Acceptable to open and dissolve in water. Available as an oral solution (10 mg/mL). RAL: Available as a chewable tablet (25 mg, 100 mg) and oral powder for suspension (100 mg/packet); neither is bioequivalent to the 400 mg adult dose. TDF: Acceptable to dissolve in water. Available as an oral powder only (40 mg/1 g) that can be mixed with soft food. TDF/FTC: Acceptable to crush and dissolve. 	

Table 3: Alternative Post-Exposure Prophylaxis Regimens for Patients Who Weigh ≥40 kg [a,b]	
Alternative Regimens	Notes
Elvitegravir 150 mg/cobicistat 150 mg/emtricitabine 200 mg/tenofovir disoproxil fumarate 300 mg (EVG/COBI/FTC/TDF) as a fixed-dose single tablet once per day (Stribild) [c]	For individuals with creatinine clearance (CrCl) <70 mL/min: Fixed-dose single tablet EVG/COBI/TDF/FTC is <i>contraindicated</i> .
<ul style="list-style-type: none"> TDF 300 mg/FTC 200 mg (Truvada) <i>plus</i> ritonavir (RTV; Norvir) 100 mg <i>plus</i> darunavir (DRV; Prezista) 800 mg once per day [d] Substitutions: <ul style="list-style-type: none"> For FTC: Lamivudine (3TC; Epivir) 300 mg once per day. For DRV: Atazanavir (ATV; Reyataz) 300 mg once per day <i>or</i> fosamprenavir (FPV; Lexiva) 1400 mg once per day <i>plus</i> RTV 100 mg once per day. 	For individuals with baseline CrCl <50 mL/min: Adjust dosing of 3TC/FTC <i>plus</i> TDF.
<p>a. All medications are taken by mouth for 28 days.</p> <p>b. Available alternative formulations and methods of administration:</p> <ul style="list-style-type: none"> 3TC: Acceptable to crush or split. Available as an oral solution (10 mg/mL). ATV: Acceptable to open capsule and sprinkle contents. Oral dispersible powder (50 mg/packet). DRV: Probably acceptable to crush. Available as an oral suspension (100 mg/mL). DTG: Acceptable to crush. FTC: Acceptable to open and dissolve in water. Available as an oral solution (10 mg/mL). RAL: Available as a chewable tablet (25 mg, 100 mg) and oral powder for suspension (100 mg/packet); neither is bioequivalent to the 400 mg adult dose. RTV: Available as an oral solution (80 mg/mL). TDF: Can be dissolved in water. Available as an oral powder (40 mg/1 g) that can be mixed with soft food only. TDF/FTC: Acceptable to crush and dissolve. <p>c. Cobicistat-containing regimens should not be used during pregnancy.</p> <p>d. If DRV or ATV are prescribed during pregnancy, dose adjustments are required. See the guideline section <i>PEP During Pregnancy or Breastfeeding</i> or Clinicalinfo.HIV.gov > Table 10. <i>Antiretroviral Drug Use in Pregnant Women with HIV: Pharmacokinetic and Toxicity Data in Human Pregnancy and Recommendations for Use in Pregnancy</i>.</p>	


 Table 4: Post-Exposure Prophylaxis Regimens for Patients 2 to 12 Years Old Who Weigh <40 kg	
<p>See DHHS for dosing, administration and additional information about each medication. Each medication name below is linked to a page about that medication.</p>	
<ul style="list-style-type: none"> Preferred: Tenofovir disoproxil fumarate (TDF; Viread) <i>plus</i> emtricitabine (FTC; Emtriva) <i>plus</i> raltegravir (RAL; Isentress). TDF/FTC is available as the fixed-dose combination (Truvada). <ul style="list-style-type: none"> Substitutions: <ul style="list-style-type: none"> Lamivudine (3TC; Epivir) may be substituted for FTC. Dolutegravir (DTG; Tivicay) may be substituted for RAL. Alternatives: <ul style="list-style-type: none"> Age ≥2 years to 12 years: Zidovudine (ZDV; Retrovir) <i>plus</i> 3TC (Epivir) <i>plus</i> RAL (Isentress) or lopinavir/ritonavir (LPV/RTV; Kaletra). Age ≥3 years to <12 years: TDF (Viread) <i>plus</i> FTC (Emtriva) <i>plus</i> darunavir (DRV/Prezista) <i>plus</i> ritonavir (RTV; Norvir). <ul style="list-style-type: none"> Substitution: 3TC (Epivir) may be substituted for FTC. 	

Table 5: Antiretroviral Medications to Avoid for Post-Exposure Prophylaxis				
Antiretroviral Class	Agent	<40 kg	≥40 kg	Comments
First-generation protease inhibitors	<ul style="list-style-type: none"> Indinavir (IDV; Crixivan) Nelfinavir (NFV; Viracept) 	Avoid	Avoid	Poorly tolerated.
First-generation non-nucleoside reverse transcriptase inhibitors	<ul style="list-style-type: none"> Efavirenz (EFV; Sustiva) Nevirapine (NVP; Viramune) 	Avoid	Avoid	<ul style="list-style-type: none"> EFV: Potential for neuropsychiatric adverse effects. NVP: Associated with fulminant hepatic failure and risk of Stevens-Johnson syndrome [CDC 2001].
Nucleoside reverse transcriptase inhibitors	<ul style="list-style-type: none"> Abacavir (ABC; Ziagen) Didanosine (ddI; Videx) Stavudine (d4T; Zerit) Tenofovir alafenamide (TAF) Zidovudine (ZDV, AZT; Retrovir) 	Avoid d4T, ddI, ABC, TAF	Avoid all	<ul style="list-style-type: none"> ABC: Potential for serious, sometimes fatal hypersensitivity reaction. d4T, ddI, ZDV: Significant mitochondrial toxicities. TAF: Decreased vaginal, cervical, and rectal tissue concentrations of the active moiety of (tenofovir diphosphate) in healthy volunteers [Garrett, et al. 2016; Cottrell, et al. 2017].
CCR5 antagonist	Maraviroc (MVC; Selzentry)	Avoid	Avoid	Only shows activity against R5-tropic virus.

References

CDC. Serious adverse events attributed to nevirapine regimens for postexposure prophylaxis after HIV exposures-- worldwide, 1997-2000. *MMWR Morb Mortal Wkly Rep* 2001;49(51-52):1153-1156. [PMID: 11198946] <https://www.ncbi.nlm.nih.gov/pubmed/11198946>

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Garrett KL, Cottrell ML, Prince HM, et al. Concentrations of TFV and TFVdp in female mucosal tissues after a single dose of TAF. CROI; 2016 Feb 22-25; Boston, MA. <http://www.croiconference.org/sessions/concentrations-tfv-and-tfvdp-female-mucosal-tissues-after-single-dose-taf>