

Evidence-Based Clinical Pathway: Non-occupational Post HIV Exposure Antiviral Prophylaxis (nPEP) Prepared by: Gibson D, Gentges J

11/27/16 replaces no older

Primary Sources: 1. 1) Updated Guidelines for Antiretroviral Postexposure Prophylaxis After Sexual, Injection Drug Use, or Other Nonoccupational Exposure to HIV - United States, 2016 from https://aidsinfo.nih.gov/guidelines. 2) Recommendations for Use of Antiretroviral Drugs in Pregnant HIV-1-Infected Women for Maternal Health and Interventions to Reduce Perinatal HIV Transmission in the United States from https://aids.nih.gov/guidelines. 3) Liang SY et al. Update on Emerging Infections: News From the Centers for Disease Control and Prevention. Annals of Emergency Medicine 2016; 68: 335-338

*Substantial Risk for HIV Acquisition

Exposure to blood, semen, vaginal secretions, rectal secretions, breast milk, or any body fluid that is visibly contaminated with blood if the source is known to be HIV positive

*Negligible Risk for HIV Acquisition

Exposure to Urine, nasal secretions, saliva, sweat, or tears if not visibly contaminated with blood

Additional Recommendations:

- All persons discharged with nPEP should have close follow up for HIV testing, monitoring for
- side-effects of treatment, further lab testing and treatment/vaccinations
- All persons considered for nPEP for sexual exposure should be evaluated for STIs as indicated.
- All persons considered for nPEP should be evaluated for HBV vaccination/treatment
- Provide counseling related to HIV prevention strategies, as appropriate

Pathway applicability: Patients with potential *exposure (Vagina, rectum, eye, mouth, or other mucous membrane, nonintact skin, or percutaneous contact) to HIV Is there substantial risk of

HIV acquisition based on Hepatitis C (same exposure? bodily fluids as HIV) serology recommended Yes Yes HIV antibody test (Combined Ag/Ab test preferred) Serum Creatinine >72 hours **LFTs** since **Pregnancy Test** exposure? Hepatitis B serology (HBsAg, HBsAb, HBcAb) Hepatitis C Ab Νo

Age < 28 days

Consult Pediatric

ID/HIV Specialist

Preferred Regimen Tenofovir DF 300

Age >13, normal CrCl

mg/ emtricitabine 200 mg (Truvada) QD with

(raltegravir 400 mg **BID** OR dolutegravir 50

Alternative Regimen

mq QD)

Tenofovir DF 300 mg/ emtricitabine 200 mg (Truvada) QD with

(Darunivir 800 mg [as 2, 400 mg tabs] QD) AND ritonavir 100 mg QD

Preferred Regimen

Age >13, CrCl < 60

Zidovudine and lamivudine, doses adjusted to CrCl with

(raltegravir 400 mg

OR dolutegravir 50 mg QD)

Alternative Regimen

Zidovudine and lamivudine, doses adjusted to CrCl with

(Darunivir 800 mg [as 2, 400 mg tabs] QD) AND ritonavir 100 mg QD

Preferred Regimen

Age 2-12**

Tenofovir DF, emtricitabine, and raltegravir

Alternative Regimen

Zidovudine and lamivudine

(raltegravir OR lopinavir/ritonavir)

Alternative Regimen

Tenofovir DF, emtricitabine, and lopinavir/ritonavir

Preferred Regimen

Age 28 days - 2 years**

Tenofovir DF, emtricitabine, and raltegravir

Alternative Regimen

Zidovudine and lamivudine (raltegravir OR lopinavir/ritonavir)

Alternative Regimen

Tenofovir DF, emtricitabine, and lopinavir/ritonavir

Lamuvidine Renal Dosing CrCl (mL/min) Dose

nPEP not recommended

Hepatitis B (any bodily fluid

exposure) and

30-49 150 mg q24h 15-29 150 mg, then 100 mg q24h 5-14 150 mg, then 50

mg q24h 1 x 50 mg, then <5 or on HDc

25 mg q24h

Zidovudine Renal Dosing

CrCl (mL/min) Dose <15 or HDc 100 TID or 300mg daily

^{**} Table 6 of http://www.cdc.gov/hiv/pdf/programresources/cdc-hiv-npep-guidelines.pdf has dosing for pediatric patients (click here)