

OCCUPATIONAL POST-EXPOSURE PROPHYLAXIS (PEP) AGAINST HIV IN ADULTS

Any possible exposure to body fluids from potentially HIV-infected patients, particularly needlestick injuries, should be reported immediately and prompt action taken. The incident should be reported to the senior nurse (the PNO or Night Superintendent) and Patient Safety Nurse and the member of staff involved should be assessed by a Medical Officer or Specialist.

The Lead Consultant for HIV services is Dr Josephine Mugume.

Kisiizi follows the national guidelines recommended by the Uganda Ministry of Health and the steps outlined in the information below should be followed.

Kisiizi Pharmacy stocks anti-retroviral medication.

Reference: December 2016 Consolidated Guidelines for Prevention and Treatment of HIV in Uganda



Ministry of Health

See extracted section below on PEP:

CONSOLIDATED GUIDELINES FOR PREVENTION AND TREATMENT OF HIV IN UGANDA

3.3.2. POST-EXPOSURE PROPHYLAXIS

Definition: Post-exposure prophylaxis (PEP) is the short-term use of ARVs to reduce the likelihood of acquiring HIV infection after potential occupational or non-occupational exposure.

Types of exposure:

- Occupational exposures occur in the health care or laboratory setting and include sharps and needlestick injuries or splashes of body fluids to the skin and mucous membranes.
- Non-occupational exposures include unprotected sex, exposure following assault like in rape and defilement, and road traffic accidents.

Steps in assessing a potential PEP recipient

Health facilities providing PEP must have trained health care workers on infection prevention and control, and management of PEP. The health care workers should use the steps in Table 7 to assess clients for PEP eligibility and provide PEP.

Table 7: Steps for providing post-exposure prophylaxis (PEP)

Step	Description					
Step 1: Clinical	Conduct a rapid assessment of the client to assess exposure and risk and provide					
assessment and	immediate care.					
providing first	Occupational exposure:					
aid	After a needle stick or sharp injury					
	Do not squeeze or rub the injury site					
	 Wash the site immediately with soap or mild disinfectant (chlorhexidine gluconate 					
	solution)					
	Use antiseptic hand rub/gel if no running water					
	 Don't use strong, irritating antiseptics (like bleach or iodine) 					
	After a splash of blood or body fluids in contact with intact skin					
	Wash the area immediately					
	Use antiseptic hand rub/gel if no running water					
	 Don't use strong, irritating antiseptics (like bleach or iodine) 					
	For exposure-specific injuries, refer to the PEP Guidelines					
Step 2:	Provide PEP when:					
Eligibility	Exposure occurred within the past 72 hours; and					
assessment	The exposed individual is not infected with HIV; and					
	The 'source' is HIV-infected, has unknown HIV status or is high risk					
	Do not provide PEP when:					
	The exposed individual is already HIV-positive					
	The source is established to be HIV-negative					
	 Individual was exposed to bodily fluids that do not pose a significant risk (e.g. tears, 					
	non-blood-stained saliva, urine, sweat)					
	Exposed individual declines an HIV test					

Step 3:	Counsel on:				
Counseling and	The risk of HIV from the exposure				
support	Risks and benefits of PEP				
	Side effects of ARVs (see Table 52)				
	Enhanced adherence if PEP is prescribed				
	Importance of linkage for further support for sexual assault cases				
Step 4:	PEP should be started as early as possible, not beyond 72 hours of exposure				
Prescription	Recommended regimens include:				
	o Adults: TDF+3TC+ATV/r				
	o Children: ABC+3TC+LPV/r				
	A complete course of PEP should run for 28 days				
	Do not delay the first doses because of lack of baseline HIV test				
	Document the event and patient management in the PEP register (ensure)				
	confidentiality of patient data)				
Step 5: Provide	Discontinue PEP after 28 days				
follow-up	Perform follow-up HIV testing three months after exposure				
	 Counsel and link to HIV clinic for care and treatment if HIV-positive 				
	 Provide prevention and education/risk reduction counseling if HIV-negative 				

TOXICITIES AND SIDE EFFECTS FO COMMONLY USED ARV'S

Table~52: Toxicities/side~effects~of~commonly~used~ARVs~and~recommended~substitutions

Age category	Regimen	Major toxicity events	Responsible ARV	Suggested management
Adults, adolescents, pregnant and lactating women	TDF+3TC+EFV	Persistent central nervous system toxicity (such as dizziness, insomnia, abnormal dreams) or mental symptoms (anxiety, depression, mental confusion) Convulsions Hepatotoxicity Severe skin and hypersensitivity reactions Gynecomastia Chronic kidney disease	EFV	Reassure, lower the dose of EFV to 400 mg. If persists substitute EFV with DTG and use regimen TDF+3TC+DTG. Substitute with DTG Use regimen TDF+3TC+DTG Substitute with ABC
		Acute kidney injury and Fanconi syndrome Decreased bone mineral density Lactic acidosis or severe hepatomegaly with steatosis	TDF	Use regimen ABC+3TC+EFV
	TDF+3TC+DTG	Chronic kidney disease Acute kidney injury and Fanconi syndrome Decreased bone mineral density Lactic acidosis or severe hepatomegaly with steatosis	TDF	Substitute with ABC Use regimen ABC+3TC+DTG
		Hepatotoxicity Hypersensitivity reactions	DTG	Substitute with EFV Give TDF+3TC+EFV If EFV is contraindicated: use TDF+3TC+ATV/r
	ABC+3TC+DTG	Hypersensitivity reaction	ABC	Stop and substitute with TDF Use regimen: TDF+3TC+DTG If TDF is contraindicated: use AZT+3TC+DTG
		Hepatotoxicity Hypersensitivity reactions	DTG	Substitute with EFV Give TDF+3TC+EFV If EFV is contraindicated: use TDF+3TC+ATV/r

Age category	Regimen	Major toxicity events	Responsible ARV	Suggested management
Adults, adolescents, pregnant and lactating women	AZT+3TC+NVP	Severe anemia, neutropenia Lactic acidosis or severe hepatomegaly with steatosis lipoatrophy, lipodystrophy, myopathy Severe vomiting	AZT	Substitute with TDF Use regimen: TDF+3TC+NVP If TDF is contraindicated: use ABC+3TC+NVP
		Acute symptomatic hepatitis Severe skin rash Hypersensitivity reaction, Stevens-Johnson Syndrome (severe or life-threatening rash, mucosal involvement)	NVP	Substitute with DTG Use regimen: AZT+3TC+DTG
	ATV/r-based regimen	Electrocardiographic abnormalities (PR and QRS interval prolongation)		Use with caution in people with pre- existing conduction disease or who are on concomitant drugs that may prolong the PR or QRS intervals.
		Indirect hyperbilirubinemia (clinical jaundice)	ATV/r	This phenomenon is clinically benign but potentially stigmatizing. Substitute with LPV/r only if adherence is compromised.
		History of nephrolithiasis		Substitute with LPV/r or DRV/r. If boosted PIs are contraindicated, and NNRTIs have failed in first-line ART, consider salvage therapy.
	DRV/r-based regimen	Hepatotoxicity Severe skin and hypersensitivity reactions	DRV/r	Substitute with ATV/r or LPV/r. When it is used in third-line ART, limited options are available. For hypersensitivity reactions, substitute with another therapeutic class.
	ETV-based regimen	Severe skin and hypersensitivity reactions	ETV	Substitute with another therapeutic class (integrase inhibitors or boosted PIs).

Kisiizi provides access to the full consolidated guidelines via Stre@mline including the 2018 updates on ART for People Living with HIV.

Go to "Hospital Resources" then select "HIV" Category and the list of available resources will appear and can be accessed 24 hours a day. These documents are stored on the Kisiizi server so are not internet dependent.



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