

# PEP: Post-exposure Prophylaxis

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# What We Will Talk About

- Considerations When Thinking about Post-Exposure Prophylaxis (PEP)
  - Steps to Initiating PEP
  - Key Decision Points
  - HIV Testing
  - Antiretroviral Therapy
- HIV Testing: HIV Screen versus HIV RNA or DNA PCR
- PrEP

# **STEPS TO IMPLEMENTING PEP**

# TAKING THE HISTORY & MAKING DECISION TO USE PEP

*If it has been > 72 hours since exposure, do not use PEP*

- **Step 1:** Establish Risk of HIV Transmission Based on Exposure Type
- **Step 2:** Establish Risk of HIV Transmission Based on Source Person's HIV Infection Status
- **Step 3:** Decide approach to the use of PEP

# Risk of Transmission

Exposure Type	Risk of Transmission per 10,000 Exposure	Exposure Risk Category
<b>Cutaneous Exposures</b>		
Fluid on intact skin Bite without break in skin Mutual masturbation	No risk	No risk identified
Skin with compromised integrity		Low to intermediate
<b>Mucous Membranes Exposures</b>		
Kissing	No Risk	No risk identified
Oral sex Splash to eye or mouth	Risk of HIV transmission from oral sex is not known, although HIV rarely transmitted .	Low
<b>Vaginal sex without trauma</b>	<b>Unprotected receptive vaginal intercourse: 1 – 30 per 10,000</b>  <b>Unprotected insertive vaginal intercourse: 3-9 per 10,000</b>	<b>Intermediate</b>
<b>Receptive anal intercourse Traumatic sex with blood</b>	<b>Unprotected receptive anal intercourse: 50-320 per 10,000</b>	<b>High</b>
<b>Percutaneous Exposure</b>		
Bite with break in skin		Low

# Risk of Transmission

<b>Type of HIV Exposure</b>	<b>Risk of Transmission per 10,000 Exposure Events</b>
Blood transfusion	9,500
Perinatal exposure	1,300 to 4,500
Needle sharing (injection drug use)	67
Needle stick (health care professional)	32
Ingestion of human milk	0.1

# TAKING THE HISTORY & MAKING DECISION TO USE PEP

- Step 1: Establish Risk of HIV Transmission Based on Exposure Type
- **Step 2:** Establish Risk of HIV Transmission Based on **Source Person's HIV Infection Status**
- Step 3: Decide approach to the use of PEP

# Risk of HIV Transmission Base on Source Person's HIV Status

HIV Infection Status of Exposure Source	Risk of Transmission
Known not to have HIV	No risk
HIV status/risk status unknown/unknown source.	Unquantified
<b>HIV status unknown; does not have risk factors</b>	<b>Low</b>
<b>HIV status unknown: has 1 or more risk factors</b>	<b>Intermediate</b>
Known to have HIV infection	High

The following persons might be considered at high-risk for HIV: individuals with (1) history of multiple sexual partners; (2) STIs; (3) history of sex with MSM; (4) history of needle-sharing; (5) history of trading sex for goods or services. **Decisions for prophylaxis should not be made based on the likelihood that the perpetrator has HIV.**



# TAKING THE HISTORY & MAKING DECISION TO USE PEP

- Step 1: Establish Risk of HIV Transmission Based on Exposure Type
- Step 2: Establish Risk of HIV Transmission Based on Source Person's HIV Infection Status
- Step 3: Decide approach to the use of PEP

# One Approach to the Use of PEP

<b>Exposure Risk Category</b>	<b>HIV Infection Status of Source</b>	<b>Suggested Approach</b>
No risk identified	Any	No PEP
Any	Known to not have HIV	No PEP
<b>Low, intermediate, or high</b>	<b>Unknown</b>	<b>Consider PEP</b>
Low or intermediate risk	HIV infected	Consider PEP
High risk	HIV infected	Recommend PEP

# IMPLEMENTING PEP

*If it has been > 72 hours since exposure, do not use PEP*

- Step 4: Deciding the Medications to Include in PEP
- Step 5: Arranging Medications
- Step 6: Recommended Baseline Laboratory Testing for HIV and PEP

# Medications Recommended for PEP

Patient Category	PEP Recommendation
Persons $\geq$ 40 kg Can swallow pills	Dolutegravir 1 50 mg tablet daily AND Truvada 1 tablet PO daily:
Persons < 40 kg Can chew tablets	Zidovudine AND Lamivudine (3TC) AND Raltegravir chewable tablets
Children 2 weeks to 2 years of age  Persons > 2 yrs who cannot swallow pills or chew tablets	AZT AND 3TC AND Kaletra Oral Solution

# IMPLEMENTING PEP

- Step 4: Deciding the Medications to Include in PEP
- **Step 5: Arranging Medications**
- Step 6: Recommended Baseline Laboratory Testing for HIV and PEP

# Arranging Medications

- PEP needs to be started early...preferably within 2 hours, but always before 72 hours
- Should not interrupt PEP. Clients need the meds when they leave your facility if at all possible
- Consider giving a start packet
- Talk with pharmacy you are sending the script to— make sure they have the meds or can get them quickly
- If nausea a concern can use zofran
- Call to client next day to see how they are doing, taking meds correctly, have questions, and actually have all the meds
- Calls to remind youth of follow up testing

# IMPLEMENTING PEP

- Step 4: Deciding the Medications to Include in PEP
- Step 5: Arranging Medications
- Step 6: Recommended Baseline Laboratory Testing for HIV and PEP

# Baseline Testing

- CBC with auto differential
- If Kaletra or AZT ordered – add ALT to baseline testing,
- If Truvada ordered--add creatinine to baseline testing.
- Check Hepatitis C antibody.
- Check a Hepatitis B panel (Hep B surface antibody, surface antigen, core antibody)
  - If exposed patient not fully Hepatitis B immunized: patient should complete the vaccination series-give the next dose immediately
  - If Hepatitis B surface antibody present-no further action
  - If Hepatitis B surface antibody negative and patient completed vaccine series administer booster vaccine and re-evaluate serologic status in 1 month to determine if full revaccination is necessary.
  - If Hepatitis B surface antibody negative and the source is known to be Hepatitis B Surface Antigen positive—give the exposed patient Hepatitis B immune globulin (per FDA 0.06 ml/kg as a onetime dose)



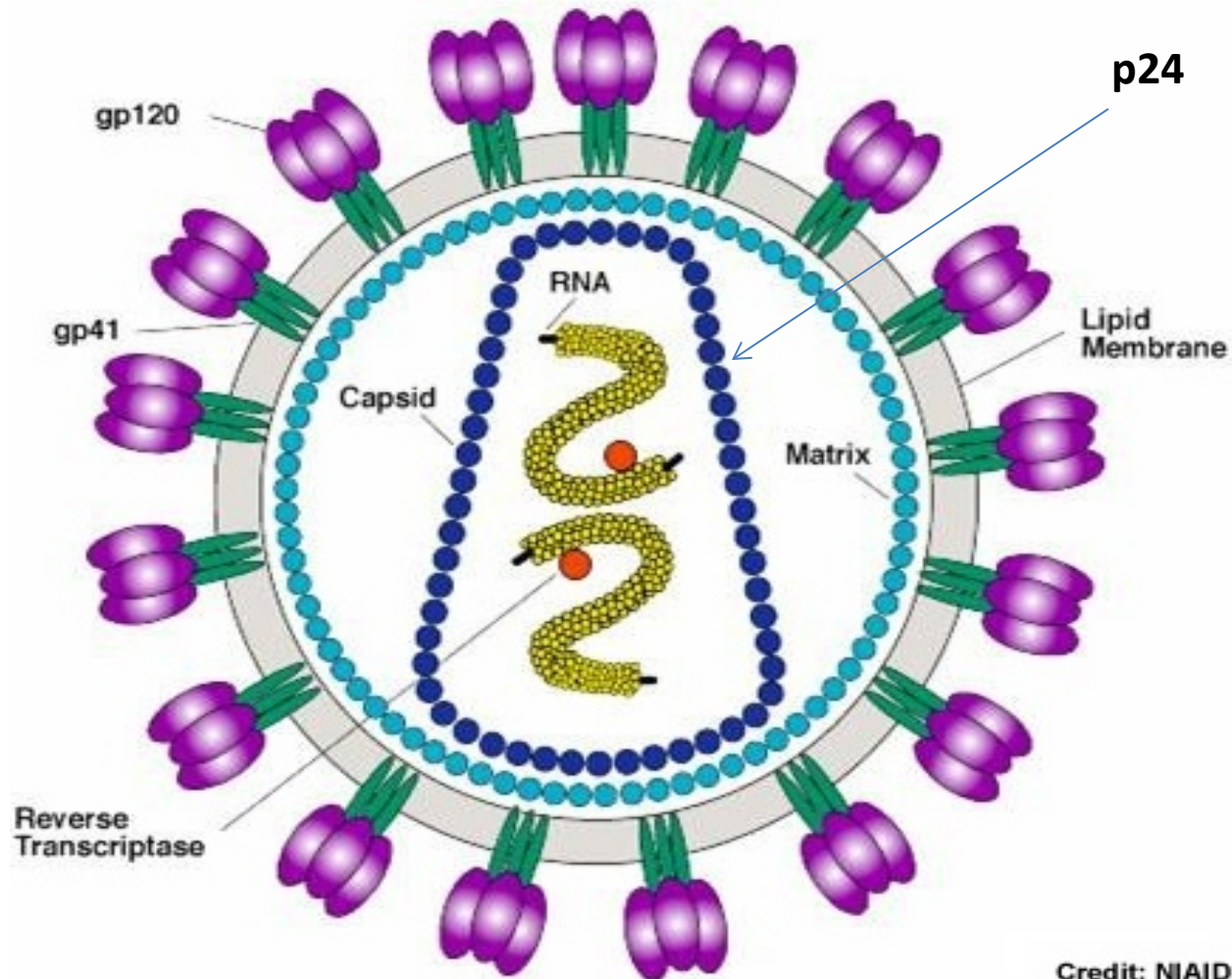
# Baseline HIV Diagnostic Testing

- **HIV antibody or antigen/antibody test**
  - Be sure to obtain verbal consent and document
  - If < 14 need parental consent
  - If 14 and older, youth must consent to testing
- If the patient had multiple sexual exposures in the recent past, they may already have early HIV infection. In addition to the HIV antibody or Ag/Ab test please obtain an **HIV RNA PCR** at baseline

# **HIV TESTING**

# HIV: The Virus

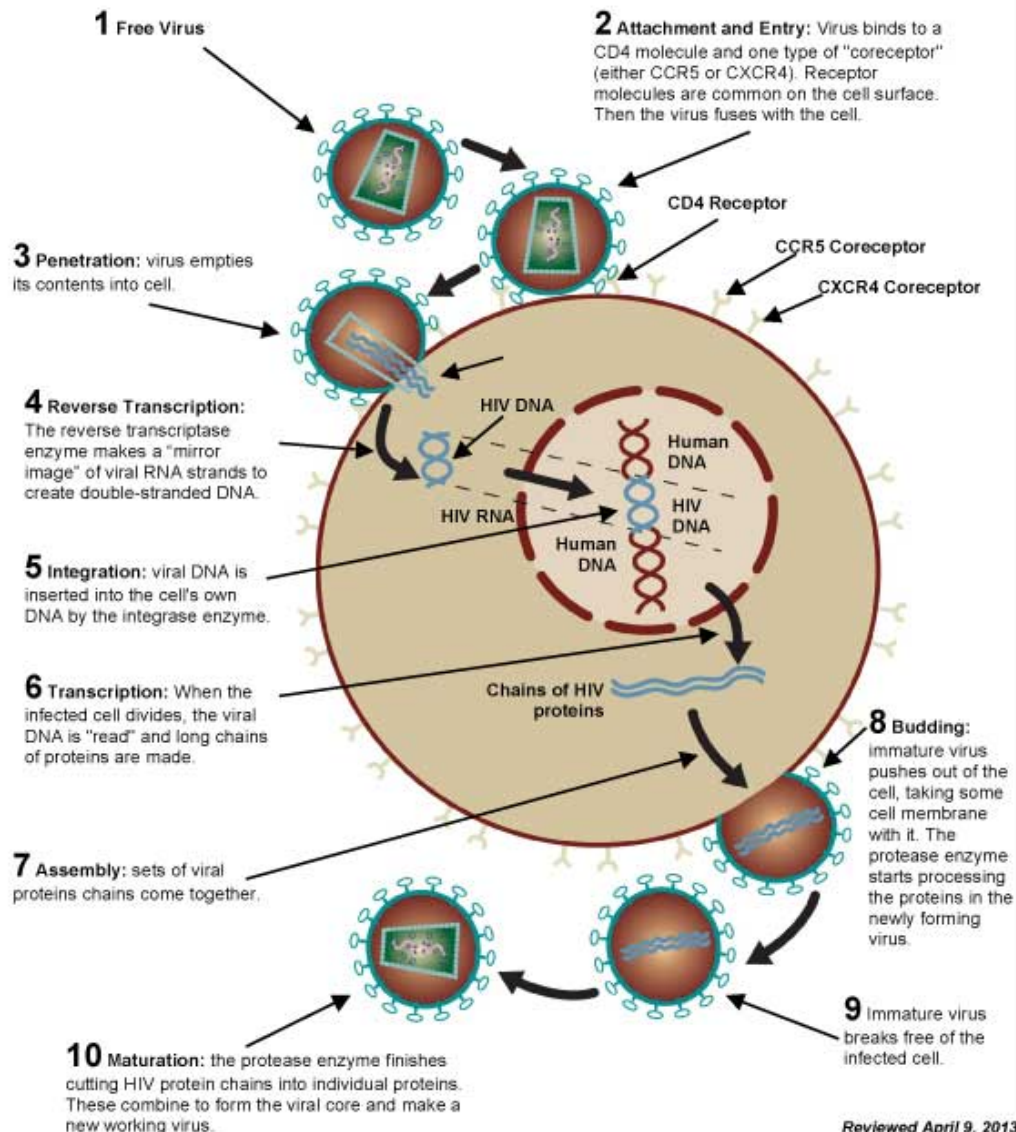
## This is the Culprit



Credit: NIAID



# HIV LIFE CYCLE



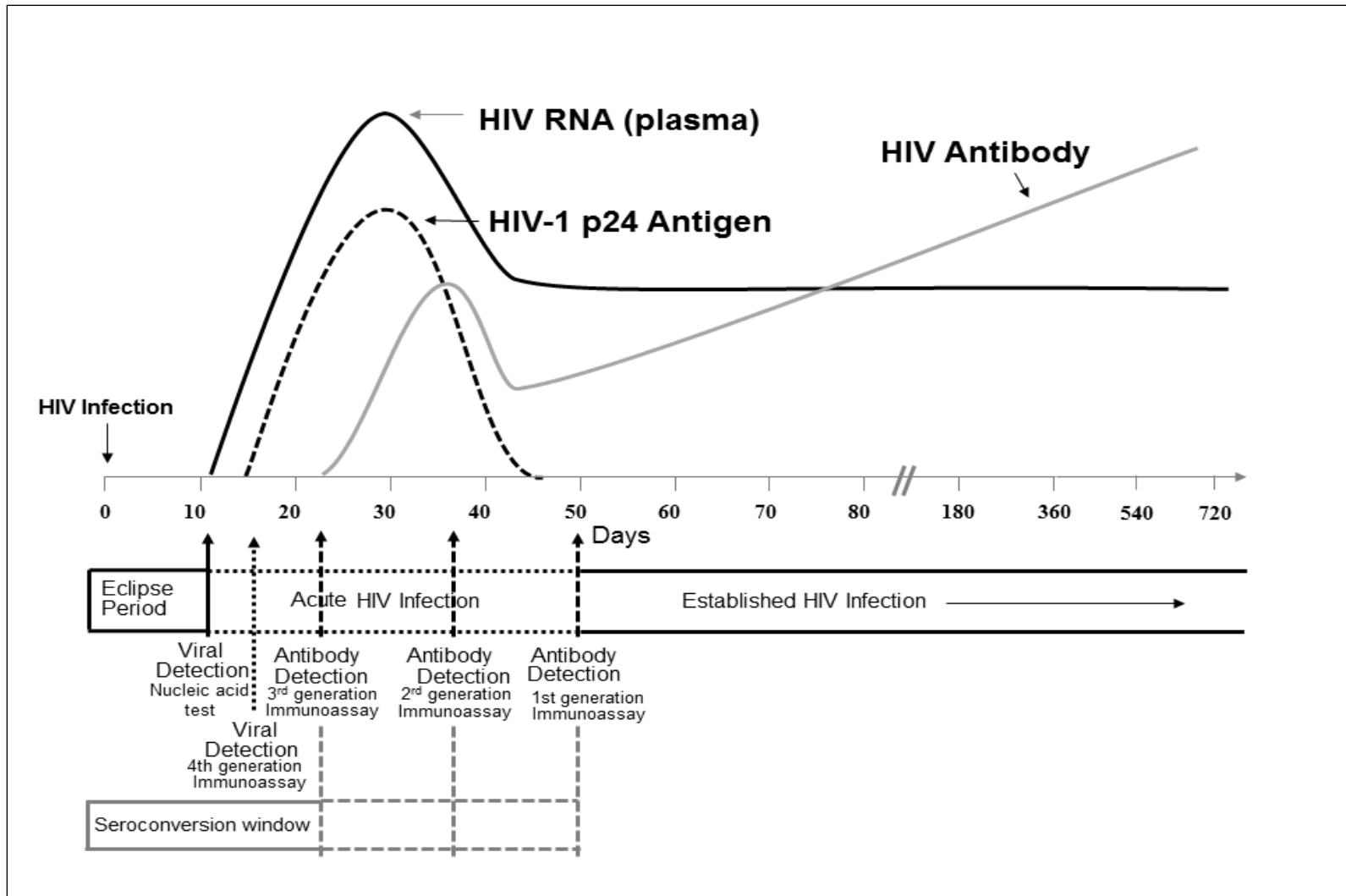
Reviewed April 9, 2013

# **SEQUENCE OF MARKER APPEARANCE**

# Sequence of HIV Testing

Day after HIV Acquisition	HIV Test	Then what happens
Day 0	HIV Infection	
Day 10	<b>HIV RNA or DNA PCR</b>	Peaks about day 30 after infection. Plateaus around day 40.
Day 15	4th generation ( <b>HIV antibody/antigen test</b> ) Detects antigen (p24)	The antigen marker, p24 peaks about 30 days after infection, rapidly decreases & may become undetectable about 45 days after infection
Day 22	3rd generation ( <b>HIV antibody screen</b> ) Detects IgM & IgG	The test marker, IgM, peaks at about 35 days then decreases until about 38 days when IgG antibodies are detectable using a 2nd generation test
Day 38	2nd generation <b>HIV antibody screen &amp; antibody differentiation tests (monospot)</b> Detects IgG	The test marker, IgG continues to increase until they are detectable by first generation tests and the western blot by day 45-50

# Appearance of Laboratory Markers for HIV-1 Infection



# 17 Year Old with Newly Diagnosed HIV: Testing Sequence

- Baseline Test: HIV DNA PCR was positive (Friday)
- After baseline testing returned we performed the following tests on the SAME blood sample
  - antigen/antibody screen negative
  - viral load > 10,000,000
- By Monday the antigen/antibody screen was reactive



**GOAL OF PEP**

# **GOAL of ARV in PEP**

- Reduce viral load to prevent HIV infection

# FOLLOW-UP CARE

- Step 8: Educate the Patient/Family about Follow-up Care
- Step 9: Follow-up Testing

# Follow up Testing

- Follow-up Lab testing:
  - **HIV antibody or Ag/Ab test** at 4 to 6 weeks and 4 months after exposure
  - For patients with **multiple sexual exposures** in the recent past, perform **an HIV DNA PCR** with the 4-6 week HIV antibody or HIV Ab/Ag test.
  - Recheck patient's Hepatitis C antibody status at 4 month post exposure.
  - Hepatitis B follow-up testing depends on baseline results.

# PEP works

- A case-control study demonstrating an **81% reduction** in the odds of HIV transmission among health care workers with percutaneous exposure to HIV who received zidovudine (ZDV) prophylaxis.

# PrEP

- We owe it to youth who participate in high risk activity to discuss PrEP
- Truvada-1 pill daily works effectively to prevent HIV infection
- PrEP= Pre post exposure prophylaxis
- For youth at high risk for HIV infection

# References

- <http://www.cdc.gov/hiv/pdf/programresources/cdc-hiv-npep-guidelines.pdf>
- WI HIV Primary Care Support Network  
“Guidelines for HIV Prophylaxis after Sexual Assault in Children and Youth”
- Please call/email if you ever have questions about PEP or PrEP