

Patients/HIV testing/cART and Dilemmas



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- A 24 y/o female , presents at 36 weeks of gestation to the ED with h/o uterine contractions
- She is admitted to labor and delivery for monitoring.
- Patient has had no prenatal care since her first trimester
- Laboratory values from 1st trimester
 - HIV: 4th generation Ag/Ab assay: Nonreactive
 - Syphilis serology: Negative
 - GC/Chlamydia: Negative



She was lost to follow up after her first trimester.

She continues to be sexually active, but has been in a monogamous relationship

HIV status of her partner is unknown

**What labs
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done**

1) None, as all her STI testing was negative in the 1st trimester

2) Repeat HIV test, syphilis serology, GC/Chlamydia

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MDHHS Perinatal Human Immunodeficiency Virus (HIV), Hepatitis B, Hepatitis C, and Syphilis Testing and Reporting Guidelines

Trimester	HIV Testing
1 st trimester	All women: (4th generation Ag/Ab assay)
3 rd Trimester	Btw 28- 32 weeks, regardless of perceived risk and/or previous negative test.
Women who are at high-risk for infection	Any time and as often regardless of previous negative results. Upon admission for delivery regardless of previous negative result. Women with S/S of acute HIV infection: plasma RNA test in conjunction with an HIV antibody test.
Women who present to L&D or ED with no available, documented test results or prenatal care	Test STAT with rapid or expedited point of care testing.

- All positive screening tests must be confirmed.

Labs and results:

HIV 4th generation Ag/Ab assay: **Reactive**

Syphilis serology: Negative

GC/Chlamydia: Negative



Clinical Decision:

1) Disregard the HIV test as false positive as she had a previous negative test

2) Start the mother on a 3 drug ART

3) Immediate C-section

4) Await result of HIV-1/HIV-2 differentiation assay and HIV RNA result



Clinical Decision:

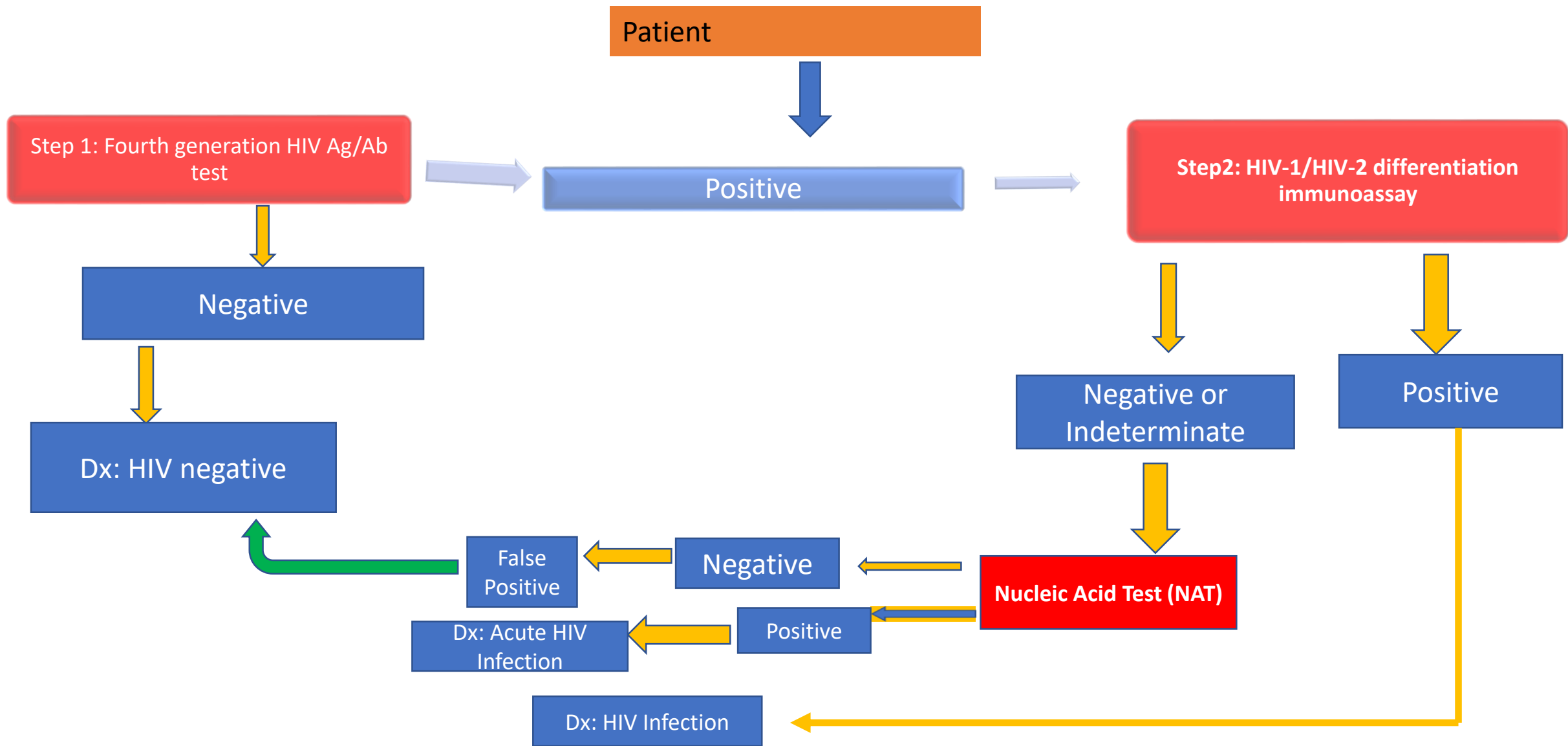
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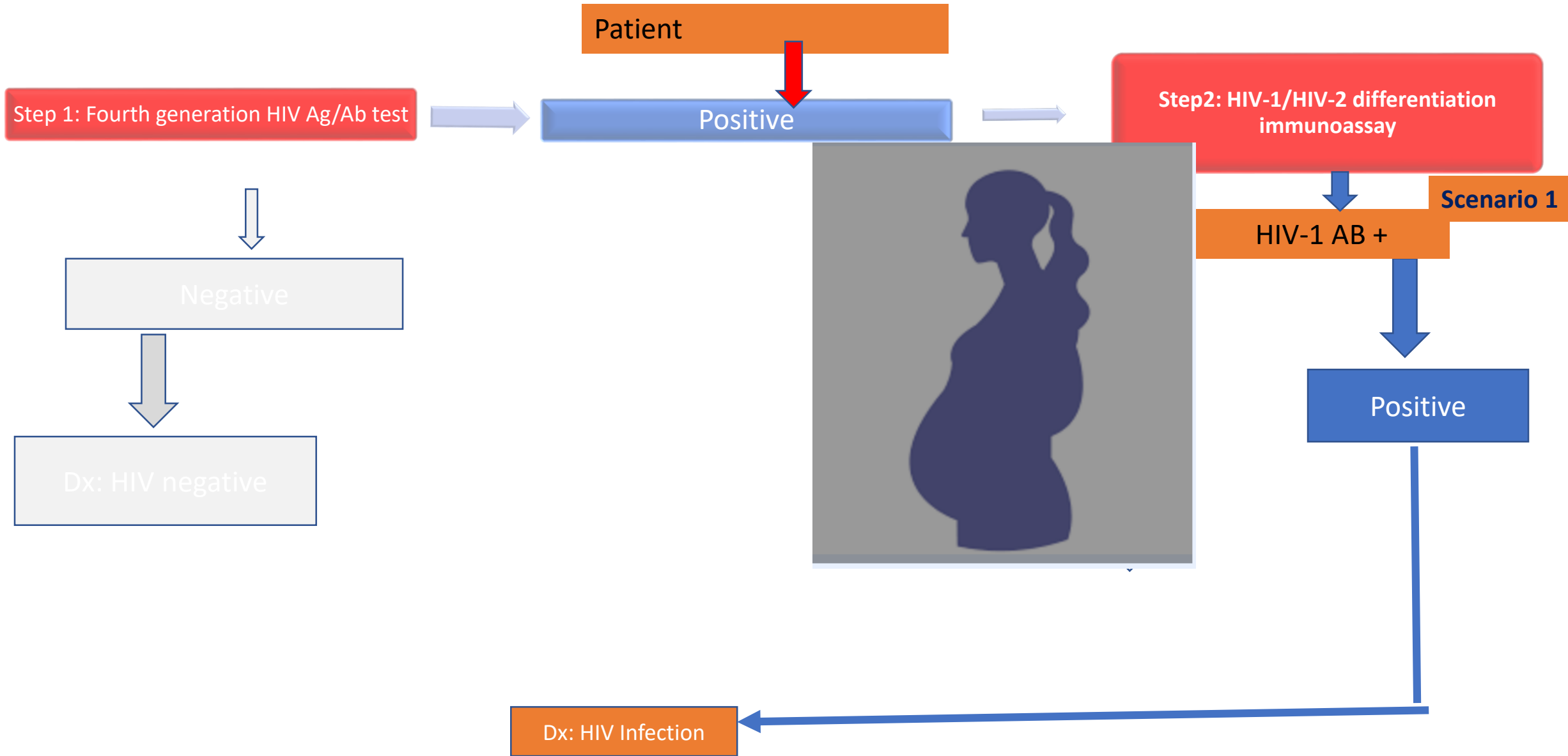
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CDC Recommendations for HIV Testing



Result of HIV-1/HIV-2 differentiation Immunoassay: Scenario 1



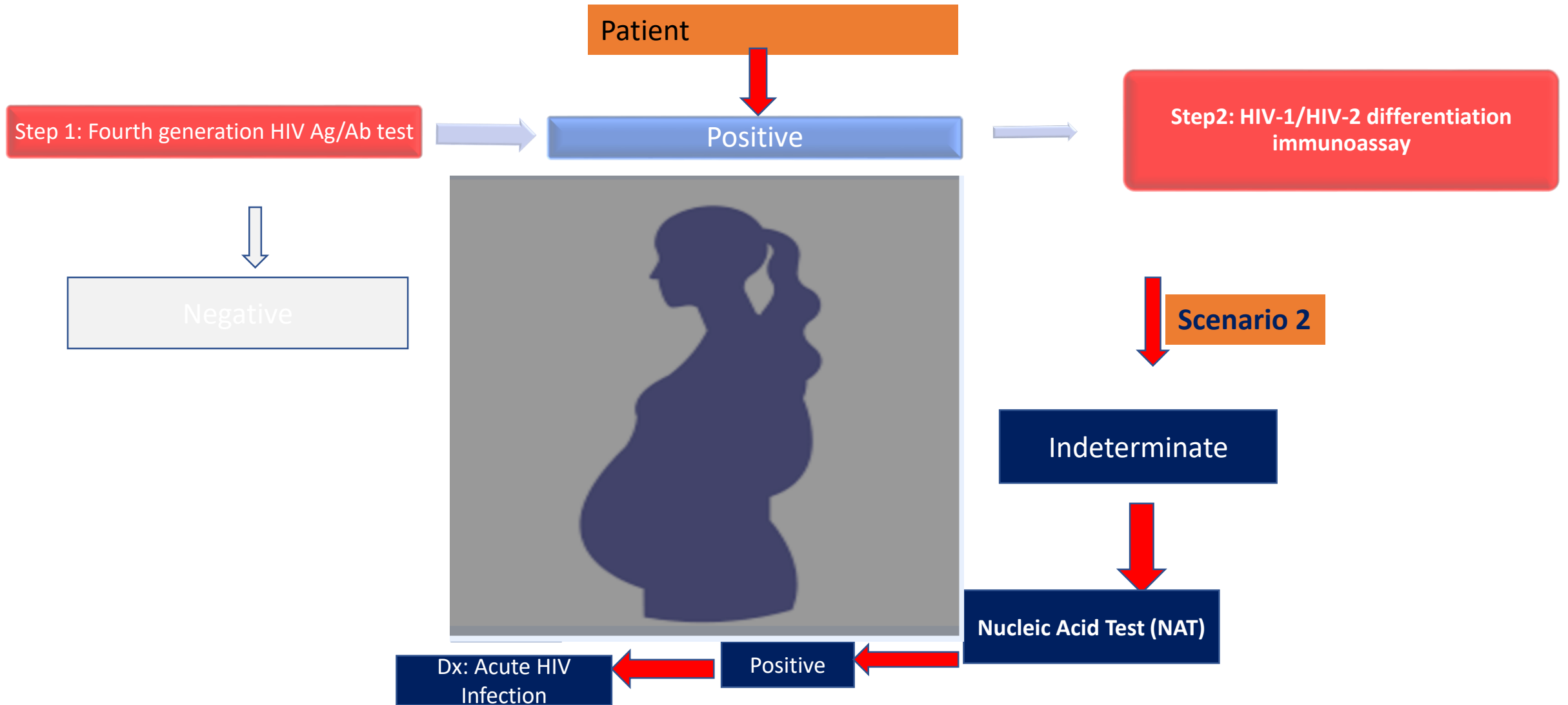


Mode of Delivery and ART for Mother; HIV+ at 36 weeks of gestation

Scheduled cesarean delivery at
38 weeks gestation

HIV RNA: Either >1000 copies/ml
or unknown: Intrapartum IV ZDV
administered to the mother

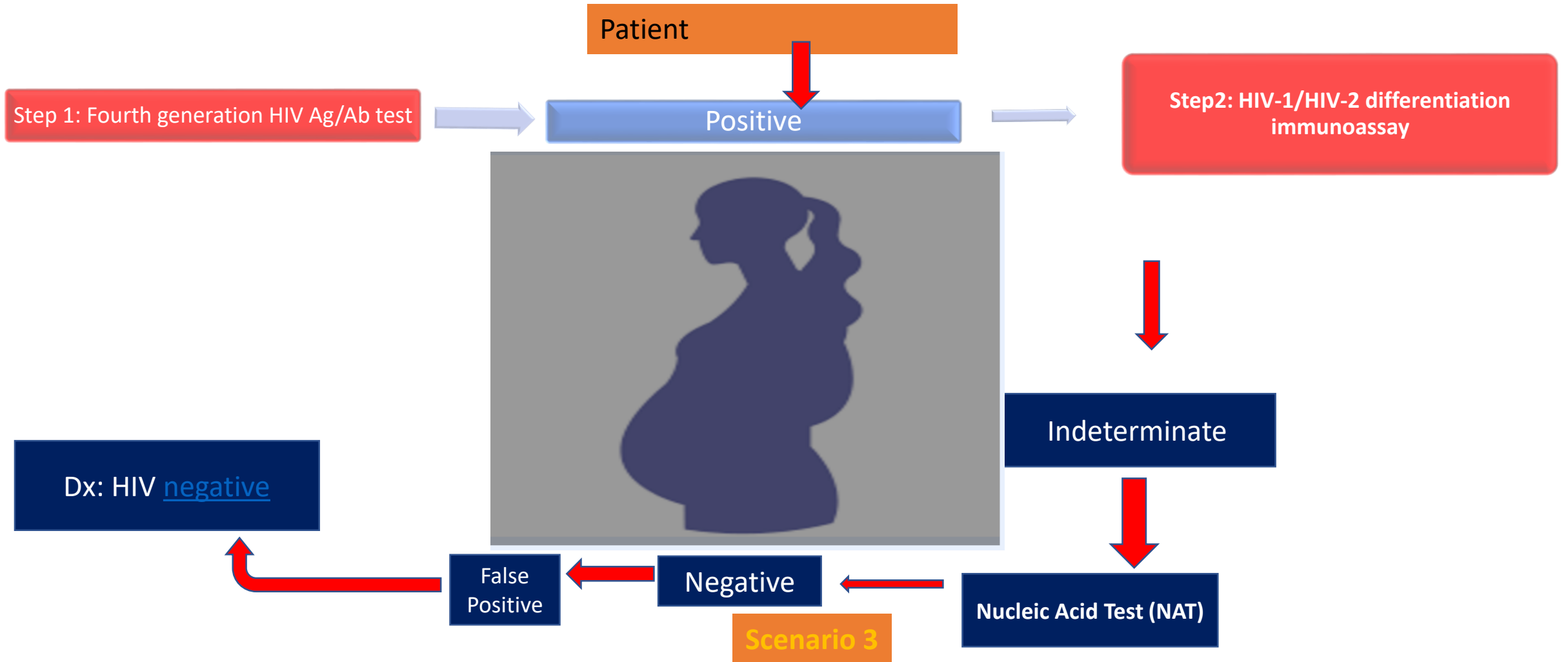
Result of HIV-1/HIV-2 differentiation Immunoassay: Scenario 2



Risk of perinatal transmission after maternal acute HIV infection

- Acute HIV infection during pregnancy is associated with a high risk of vertical transmission of HIV.
- Cesarean delivery is necessary when there is insufficient time to fully suppress a patient's viral load

Result of HIV-1/HIV-2 differentiation Immunoassay: Scenario 3



Clinical Decision

- False Positive : Fourth generation HIV Ag/Ab test
- No further HIV intervention needed

In Summary

Patient presented at 36 weeks of gestation, repeat 4th generation HIV Ag/AB test was positive. 3 possible scenarios:

HIV 4 th generation Ag/Ab test	HIV-1/HIV-2 Immunoassay	HIV Viral Load	Final Diagnoses	Intervention
Positive	HIV-1 Ab positive	Not done	HIV Infection +	C-section at 38 weeks, IV AZT
Positive	Indeterminate	Positive	Acute HIV Infection	C-section
Positive	Indeterminate	Undetected	False Positive 4 th . Generation test	No further intervention

Drug Drug Interactions

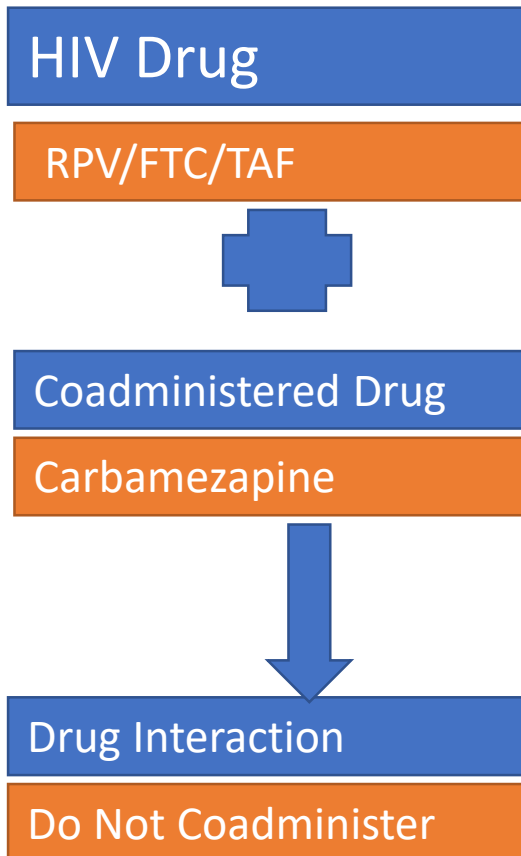
- A 42 y/o male, HIV+ for past 20 years. Initial cART: (Efavirenz / FTC/TDF). Current cART: Odefsey (RPV / FTC/TAF) for the past 4 years. Labs:
 - CD4+: > 500 cells/mm³
 - HIV VL: Undetectable
- Presented to the clinic with h/o multiple falls secondary to weakness in his legs
- Work up revealed a diagnosis of Polymyositis
- He was started on Prednisone, with slow recovery of function
- He subsequently developed severe calf muscle fasciculations and was started on Carbamazepine

Clinical Decision

1. There is potential of drug interaction between Carbamazepine and components of Odefsey (TAF+FTC+rilpivirine)
2. There is no booster in Odfesey, aka ritonavir or Cobicistat, drug interaction is unlikely

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Coadministration contraindicated:

Carbamezapine may cause significant decreases in the plasma concentrations of ART by

- 1) Inducing CYP3A : Decrease the plasma concentrations of rilpivirine
- 2) Induction of P-glycoprotein: Decrease the absorption of TAF resulting in decreased plasma concentration of TAF.

STRs and Carbamazepine

STR	Coadministered Drug	Drug Interaction
Atripla: Efavirenz / Emtricitabine/TDF	Carmazepinie	Potential interaction
Dovato: Dolutegravir/ Lamivudine	Carmazepinie	Potential interaction
Triumeq: Dolutegravir/Abacavir/ Lamivudine	Carmazepinie	Potential interaction
Biktarvy: Bictegravir/ Emtricitabine/Tenofovir alafenamide	Carmazepinie	Do no coadminister
Genvoya: Elvitegravir/Cobi/ Emtricitabine/TAF	Carmazepinie	Do not coadminister
Symtuza: Darunavir/Cobi/ Emtricitabine/TAF	Carmazepinie	Do not coadminister
Truvada+Doravirine	Carmazepinie	Do not coadminister

Clinical Decision

- Switched Carbamezapine to Levetiracetam
- No known drug interactions with antiretrovirals
- Patient was continued on Odefsey