

NRTI Mutations:	None																								
NNRTI Mutations:	None																								
RT Other Mutations:	K512R ^{100%} _{seen=1,021} • S519N ^{100%} _{seen=1,042} • Q524K ^{94%} _{seen=1,002} • K527E ^{99%} _{seen=1,965} • K530R ^{100%} _{seen=2,007} • A534S ^{100%} _{seen=2,054} • A554S ^{100%} _{seen=2,114} • I556V ^{100%} _{seen=2,114} • K558KR ^{91,127%, 91,27%} _{seen=3,039} • V559V ^{91,76%, 1,22%} _{seen=5,852}																								
<div><div>Nucleoside Reverse Transcriptase Inhibitors</div><div><table><tr><td>abacavir (ABC)</td><td>Susceptible</td></tr><tr><td>zidovudine (AZT)</td><td>Susceptible</td></tr><tr><td>stavudine (D4T)</td><td>Susceptible</td></tr><tr><td>didanosine (DDI)</td><td>Susceptible</td></tr><tr><td>emtricitabine (FTC)</td><td>Susceptible</td></tr><tr><td>lamivudine (3TC)</td><td>Susceptible</td></tr><tr><td>tenofovir (TDF)</td><td>Susceptible</td></tr></table></div></div> <div><div>Non-nucleoside Reverse Transcriptase Inhibitors</div><div><table><tr><td>doravirine (DOR)</td><td>Susceptible</td></tr><tr><td>efavirenz (EFV)</td><td>Susceptible</td></tr><tr><td>etravirine (ETR)</td><td>Susceptible</td></tr><tr><td>nevirapine (NVP)</td><td>Susceptible</td></tr><tr><td>rilpivirine (RPV)</td><td>Susceptible</td></tr></table></div></div>		abacavir (ABC)	Susceptible	zidovudine (AZT)	Susceptible	stavudine (D4T)	Susceptible	didanosine (DDI)	Susceptible	emtricitabine (FTC)	Susceptible	lamivudine (3TC)	Susceptible	tenofovir (TDF)	Susceptible	doravirine (DOR)	Susceptible	efavirenz (EFV)	Susceptible	etravirine (ETR)	Susceptible	nevirapine (NVP)	Susceptible	rilpivirine (RPV)	Susceptible
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No drug resistance mutations were found for NRTI.

No drug resistance mutations were found for NNRTI.

INSTI Major Mutations:	G118R ^{91,80%, 91,20%} _{seen=1,013} • N155H ^{12%} _{seen=1,439}										
INSTI Accessory Mutations:	None										
IN Other Mutations:	K14R ^{99%} _{seen=2,070} • V31I ^{99%} _{seen=3,074} • M50I ^{94%} _{seen=2,917} • I60M ^{96%} _{seen=1,423} • T112V ^{100%} _{seen=574} • I113V ^{100%} _{seen=574} • S119SR ^{91,80%, 91,20%} _{seen=1,005} • T124A ^{100%} _{seen=1,171} • T125A ^{99%} _{seen=1,175} • G134N ^{99%} _{seen=2,035} • K136Q ^{99%} _{seen=1,075} • V201I ^{99%} _{seen=3,042} • K211R ^{99%} _{seen=3,040} • S283G ^{100%} _{seen=4,725}										
<div><div>Integrase Strand Transfer Inhibitors</div><div><table><tr><td>bictegravir (BIC)</td><td>Intermediate Resistance</td></tr><tr><td>cabotegravir (CAB)</td><td>High-Level Resistance</td></tr><tr><td>dolutegravir (DTG)</td><td>High-Level Resistance</td></tr><tr><td>elvitegravir (EVG)</td><td>High-Level Resistance</td></tr><tr><td>raltegravir (RAL)</td><td>High-Level Resistance</td></tr></table></div></div>		bictegravir (BIC)	Intermediate Resistance	cabotegravir (CAB)	High-Level Resistance	dolutegravir (DTG)	High-Level Resistance	elvitegravir (EVG)	High-Level Resistance	raltegravir (RAL)	High-Level Resistance
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IN comments

Major

- G118R** is a nonpolymorphic mutation reported in a significant proportion of persons with VF and emergent HIVDR in persons receiving a DTG-containing regimen. It has occasionally been reported in persons receiving other INSTIs. It is associated with 5-10-fold reduced susceptibility to RAL, EVG, DTG and CAB, and 2-3 fold reduced susceptibility to BIC.
- N155H** is a common nonpolymorphic INSTI-resistance mutations. It has been reported in a high proportion of persons developing VF and HIVDR while receiving RAL, EVG, DTG, and CAB. Alone, it reduces RAL and EVG susceptibility about 10 and 30-fold, respectively. It has minimal effect on susceptibility to DTG, BIC, and CAB.

Other

- M50I** is a highly polymorphic mutation, which has a prevalence of 3% to 34% in INSTI-naïve persons depending on subtype. It has been selected in vitro by DTG and BIC in combination with R263K. It may contribute to reduced DTG and CAB susceptibility in combination with R263K.
- S119R** is a polymorphic mutation that is weakly selected by INSTIs usually in combination with several major INSTI-associated DRMs. Alone, it has little, if any effect, on INSTI susceptibility.

- There is evidence for high-level **DTG** resistance. If **DTG** is used, it should be administered twice daily.

Drug resistance mutation scores of INSTI:

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Rule	BIC ⚖	CAB ⚖	DTG ⚖	EVG ⚖	RAL ⚖
<u>G118R</u>	30	60	50	60	60
<u>N155H</u>	10	25	10	60	60
Total	40	85	60	120	120