

NRTI Mutations:None

NNRTI Mutations:None

RT Other Mutations:K530R 100%
seen=177 • A534S 100%
seen=264 • **W533*** 100%
seen=129 • **G543R** 100%
seen=148 • A554N 100%
seen=402 • K558R 100%
seen=1,043

Nucleoside Reverse Transcriptase Inhibitors		Non-nucleoside Reverse Transcriptase Inhibitors	
abacavir (ABC)	Susceptible	doravirine (DOR)	Susceptible
zidovudine (AZT)	Susceptible	efavirenz (EFV)	Susceptible
stavudine (D4T)	Susceptible	etravirine (ETR)	Susceptible
didanosine (DDI)	Susceptible	nevirapine (NVP)	Susceptible
emtricitabine (FTC)	Susceptible	rilpivirine (RPV)	Susceptible
lamivudine (3TC)	Susceptible		
tenofovir (TDF)	Susceptible		

No drug resistance mutations were found for NRTI.

No drug resistance mutations were found for NNRTI.

INSTI Major Mutations:**R263K** 100%
seen=2,290

INSTI Accessory Mutations:**G163R** 100%
seen=2,752

IN Other Mutations:S17N 100%
seen=2,508 • K71KE 10.100%
seen=1,114 • I72V 100%
seen=2,630 • **L74I** 100%
seen=2,680 • T112I 100%
seen=1,179 • T124A 100%
seen=2,613 • T125A 100%
seen=2,613 • **W132*** 100%
seen=2,605 • G134N 100%
seen=2,230 • K136Q 100%
seen=2,230 • V201I 100%
seen=4,947 • **R224Q** 100%
seen=4,778 • L234V 100%
seen=4,629 • V260I 100%
seen=2,134 • R269K 100%
seen=2,170 • S283G 100%
seen=1,368

Integrase Strand Transfer Inhibitors	
bictegravir (BIC)	Intermediate Resistance
cabotegravir (CAB)	Intermediate Resistance
dolutegravir (DTG)	Intermediate Resistance
elvitegravir (EVG)	Intermediate Resistance
raltegravir (RAL)	Intermediate Resistance

IN comments

Major

- R263K** is a nonpolymorphic mutation selected in vitro by EVG, DTG, BIC, and CAB. It occurs in a high proportion of persons who develop VF and emergent HIVDR while receiving DTG. Alone, it reduces DTG, BIC, and CAB susceptibility about 2-fold.

Accessory

- G163R/K** are nonpolymorphic in all subtypes except subtype F. They are accessory resistance mutations as they usually occur in combination with other INSTI-resistance mutations particularly N155H.

Other

- L74I** is a highly polymorphic mutation with a prevalence of 3% to 30% depending on subtype. It is the consensus amino acid in subtype A viruses belonging to the A6 clade. It does not appear to be selected by any of the INSTIs or to reduce their susceptibility.

- This virus is predicted to have intermediate-level reduced susceptibility to **CAB**. The use of the combination of **CAB**/RPV should be considered to be contraindicated.
- There is evidence for intermediate **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>R263K</u>	30	30	30	30	25
<u>G163R</u>	0	0	0	15	15
Total	30	30	30	45	40