

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:I13V 98%
n=1,832 • G16E 98%
n=1,263 • R41K 99%
n=1,387 • L63A 99%
n=1,388

Protease Inhibitors	
atazanavir/r (ATV/r)	Susceptible
darunavir/r (DRV/r)	Susceptible
fosamprenavir/r (FPV/r)	Susceptible
indinavir/r (IDV/r)	Susceptible
lopinavir/r (LPV/r)	Susceptible
nelfinavir (NFV)	Susceptible
saquinavir/r (SQV/r)	Susceptible
tipranavir/r (TPV/r)	Susceptible

No drug resistance mutations were found for PI.

NRTI Mutations:[V75M](#) 99%
n=1,372 • [M184V](#) 100%
n=1,382 • [T215C](#) 100%
n=1,328

NNRTI Mutations:[V108I](#) 100%
n=1,677 • [H221Y](#) 99%
n=1,282

RT Other Mutations:[P4H](#) 100%
n=1,875 • [L34V](#) 100%
n=1,231 • [V35T](#) 100%
n=1,231 • [T39I](#) 100%
n=1,268 • [V60L](#) 100%
n=1,626 • [K102R](#) 100%
n=1,632 • [V118V](#) 9.69%
n=1,630 • [D121Y](#) 100%
n=1,561 • [K122E](#) 100%
n=1,351 • [D123E](#) 100%
n=1,328 • [I135T](#) 100%
n=1,738 • [D177E](#) 100%
n=1,681 • [I202V](#) 100%
n=1,628 • [Q207E](#) 99%
n=1,302 • [F214L](#) 100%
n=1,228 • [V241I](#) 99%
n=1,632 • [V245I](#) 94%
n=1,282 • [D250E](#) 100%
n=1,06

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

RT comments

NRTI

- [V75T](#)/[M](#)/[A](#)/[S](#) are nonpolymorphic accessory NRTI-selected mutations. They appear to have minimal phenotypic effects on AZT, ABC, and TDF.
- [M184V](#)/[I](#) cause high-level in vitro resistance to 3TC and FTC and low/intermediate resistance to ABC (3-fold reduced susceptibility). [M184V](#)/[I](#) are not contraindications to continued treatment with 3TC or FTC because they increase susceptibility to AZT and TDF and are associated with clinically significant reductions in HIV-1 replication.
- T215Y/[F](#) are TAMs that causes intermediate/high-level resistance to AZT and potentially low-level resistance to ABC and TDF. [T215S](#)/[C](#)/[D](#)/[E](#)/[H](#)/[V](#)/[N](#)/[A](#)/[L](#) do not reduce NRTI susceptibility but arise from viruses that once contained T215Y/[F](#). The presence of one of these revertant mutations suggests that the patient may have once been infected with a virus containing T215Y/[F](#).

NNRTI

- [V108I](#) is a relatively non-polymorphic accessory mutation selected in vitro and/or in vivo with each of the NNRTIs. It appears to contribute to reduced susceptibility to most NNRTIs only in combination with other NNRTI-resistance mutations.
- [H221Y](#) is a non-polymorphic accessory mutation selected primarily by NVP, RPV, and DOR. It frequently occurs in combination with Y181C.

Other

- [V118I](#) is a polymorphic accessory NRTI-resistance mutation that often occurs in combination with multiple TAMs.

- This virus is predicted to have low-level reduced susceptibility to [RPV](#). The use of the combination of CAB/[RPV](#) should be considered to be relatively contraindicated.

Drug resistance mutation scores of NRTI:

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Rule	ABC ⇅	AZT ⇅	D4T ⇅	DDI ⇅	FTC ⇅	3TC ⇅	TDF ⇅
M184V	15	-10	-10	10	60	60	-10
V75M	0	10	30	15	0	0	0
T215C	0	10	20	10	0	0	0
Total	15	10	40	35	60	60	-10

Drug resistance mutation scores of NNRTI:

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Rule	DOR ⇅	EFV ⇅	ETR ⇅	NVP ⇅	RPV ⇅
V108I	10	10	0	15	0
H221Y	10	10	10	15	15
Total	20	20	10	30	15