

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:

I13V100%
from 23,612

 •

E35D98%
from 11,880

 •

M36I98%
from 11,875

 •

N37DE11-17%
from 11,875

 •

R41K98%
from 11,101

 •

R57RK11-98%
from 29,202

 •

I62IV11-12%
from 29,208

 •

H69K100%
from 23,764

 •

L89M98%
from 11,638

Protease Inhibitors	
atazanavir/r (ATV/r)	Susceptible
darunavir/r (DRV/r)	Susceptible
lopinavir/r (LPV/r)	Susceptible

No drug resistance mutations were found for PI.

NRTI Mutations:

M184MV11-100%
from 14,811

NNRTI Mutations:

K103KN11-100%
from 11,817

 •

N147S11-100%
from 11,817

 •

V108VI11-100%
from 11,817

 •

P225PH11-100%
from 11,817

RT Other Mutations:

E6D100%
from 14,800

 •

K20KR11-100%
from 14,800

 •

V35T98%
from 11,297

 •

V60I98%
from 11,928

 •

K122E98%
from 11,200

 •

I135T97%
from 11,838

 •

K173S97%
from 14,011

 •

Q174K97%
from 14,012

 •

D177E98%
from 14,384

 •

V179I97%
from 14,386

 •

G196E90%
from 11,858

 •

T200A98%
from 11,208

 •

Q207A98%
from 10,092

 •

R211S98%
from 9,482

 •

F214L98%
from 11,812

 •

V245E97%
from 14,330

 •

D250E98%
from 14,330

 •

T286A97%
from 14,870

 •

V292I97%
from 14,777

 •

I293V97%
from 14,778

 •

E297K97%
from 14,580

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

RT comments

NRTI

- 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.

NNRTI

- K103N is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.
- 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.
- P225H is a non-polymorphic EFV-selected mutation that usually occurs in combination with K103N. The combination of P225H and K103N synergistically reduces NVP, EFV and DOR susceptibility.

Other

- V179I is a polymorphic mutation that is frequently selected in persons receiving ETR and RPV. However, it has little, if any, direct effect on NNRTI susceptibility.

Drug resistance mutation scores of NRTI:

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Rule	ABC	AZT	FTC	3TC	TDF
M184MV	15	-10	60	60	-10

Drug resistance mutation scores of NNRTI:

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Rule	DOR	EFV	ETR	NVP	RPV
K103KN + P225PH	10	0	0	0	0
V108VI	10	10	0	15	0
P225PH	20	45	0	45	0
K103KN	0	60	0	60	0
Total	40	115	0	120	0