

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:

L10I14%
from 29,287

 •

I13V18%
from 40,012

 •

K14R10%
from 40,008

 •

G16E10%
from 40,608

 •

L19I13%
from 42,087

 •

K20R12%
from 42,107

 •

E35D14%
from 47,082

 •

M36I18%
from 47,073

 •

R41K10%
from 47,082

 •

R57K10%
from 42,002

 •

L63T10%
from 39,011

 •

H69Q10%
from 38,688

 •

L89M10%
from 25,109

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

PR comments

Other

- **L10I/V** are polymorphic, PI-selected accessory mutations that increase the replication of viruses with other PI-resistance mutations.
- **K20R** is a highly polymorphic PI-selected accessory mutation that increases replication fitness in viruses with PI-resistance mutations.

No drug resistance mutations were found for PI.

NRTI Mutations:

L74V12%
from 12,609

 •

M184V11%
from 21,803

NNRTI Mutations:

L100I10%
from 8,500

 •

K103N10%
from 8,780

RT Other Mutations:

E6D14%
from 18,104

 •

V35T18%
from 20,702

 •

V50I18%
from 18,703

 •

K102R10%
from 8,807

 •

K122E10%
from 10,005

 •

D123N10%
from 10,005

 •

I135T10%
from 14,208

 •

I142V10%
from 15,708

 •

K173N5-12%
from 20,008

 •

Q174KL4-10%
from 11,008

 •

D177E10%
from 20,003

 •

I178L10%
from 11,805

 •

V179I10%
from 11,812

 •

Q207A10%
from 18,100

 •

R211S10%
from 19,008

 •

F214L14%
from 17,000

 •

V243E13%
from 20,300

 •

E248D10%
from 10,003

 •

D250E18%
from 13,800

 •

A272P10%
from 10,000

 •

T286TA1-80%
from 6,101

 •

V292VI1-80%
from 10,000

 •

I293V14%
from 5,000

 •

P294PT1-80%
from 5,000

 •

K311KR0-70%
from 5,100

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

RT comments

NRTI

- **L74V** causes intermediate ABC resistance. L74I causes low-level ABC resistance.
- **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

- **L100I** is a non-polymorphic mutation that usually occurs in combination with K103N. In this setting it confers high-level resistance to NVP, EFV, and RPV and intermediate resistance to ETR and DOR.
- **K103N** is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.

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:

Download CSV

▼

Rule	ABC ⚡	AZT ⚡	FTC ⚡	3TC ⚡	TDF ⚡
<u>L74V</u>	30	0	0	0	0
<u>L74V + M184V</u>	15	0	0	0	0
<u>M184V</u>	15	-10	60	60	-10
Total	60	-10	60	60	-10

Drug resistance mutation scores of NNRTI:

Download CSV

▼

Rule	DOR ⚡	EFV ⚡	ETR ⚡	NVP ⚡	RPV ⚡
<u>L100I</u>	15	60	30	60	60
<u>L100I + K103N</u>	15	0	0	0	0
<u>K103N</u>	0	60	0	60	0
Total	30	120	30	120	60