

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:T12TP 1:47% (n=126) • I13V 98% (n=26,308) • I15V 90% (n=27,629) • K20I 97% (n=28,229) • E35D 96% (n=30,885) • M36I 98% (n=30,883) • R41K 98% (n=30,714) • I62V 92% (n=26,882) • H69K 96% (n=25,835) • T74S 92% (n=22,038) • L89M 98% (n=33,800)

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

- PR comments
- Other
- K20I is the consensus amino acid in subtype G and CRF02_AG. In subtypes B and C, K20I is a PI-selected mutation of uncertain effects on currently used PIs.
 - T74S is a PI-selected accessory mutation that is polymorphic in most non-B subtypes.

No drug resistance mutations were found for PI.

NRTI Mutations:

K70R 94% (n=9,224) • L74I 94% (n=8,763) • M184V 97% (n=11,805) • K219E 90% (n=11,202)

NNRTI Mutations:

K103N 97% (n=7,862) • M230L 90% (n=22,264)

RT Other Mutations:T7A 90% (n=14,582) • K20R 92% (n=14,721) • V35T 97% (n=13,382) • T39G 98% (n=12,621) • V118I 98% (n=8,261) • K122E 97% (n=8,830) • I135T 97% (n=10,977) • I142T 98% (n=11,869) • K173S 98% (n=12,621) • Q174K 97% (n=12,820) • V179I 98% (n=12,288) • T200A 97% (n=8,387) • I202V 98% (n=10,325) • Q207K 94% (n=9,308) • R211S 90% (n=10,321) • F214L 90% (n=9,802) • P225R 94% (n=11,548) • L228H 90% (n=11,982) • V243Q 98% (n=14,888) • E248D 98% (n=14,582) • A272P 97% (n=14,582) • K281R 92% (n=11,861) • T286A 97% (n=12,263) • E291D 98% (n=11,317) • V292I 98% (n=11,118) • I293V 97% (n=11,170) • E312D 93% (n=9,374) • V317A 92% (n=9,263)

Nucleoside Reverse Transcriptase Inhibitors		Non-nucleoside Reverse Transcriptase Inhibitors	
abacavir (ABC)	Intermediate Resistance	doravirine (DOR)	High-Level Resistance
zidovudine (AZT)	Intermediate Resistance	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
- K70R is a TAM that confers intermediate resistance to AZT and contributes to reduced ABC and TDF susceptibility in combination with other TAMs.
 - 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.
 - K219E/Q/N/R are accessory TAMs that usually occur in combination with multiple other TAMs.
- NNRTI
- K103N is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.
 - M230L is an uncommon non-polymorphic mutation selected in persons receiving EFV, NVP, and RPV. It causes intermediate to high-level resistance to each of the NNRTIs.
- Other
- V118I is a polymorphic accessory NRTI-resistance mutation that often occurs in combination with multiple TAMs.
 - 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.
 - 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. P225R is a highly unusual mutation at this position.

Drug resistance mutation scores of NRTI:

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Rule	ABC ⚡	AZT ⚡	FTC ⚡	3TC ⚡	TDF ⚡
K70R	5	30	0	0	5
L74I	15	0	0	0	5
M184V	15	-10	60	60	-10
K219E	5	10	0	0	5
Total	40	30	60	60	5

Drug resistance mutation scores of NNRTI:

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Rule	DOR ⚡	EFV ⚡	ETR ⚡	NVP ⚡	RPV ⚡
M230L	60	45	30	60	60
K103N	0	60	0	60	0
Total	60	105	30	120	60