

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:I13V 98% cov=29,142 • I15V 90% cov=30,440 • E35D 93% cov=33,802 • M36I 98% cov=33,593 • R41K 98% cov=33,440 • I62V 91% cov=28,080 • H69K 94% cov=23,990 • T74S 90% cov=24,471 • L89M 98% cov=33,983

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

PR comments

- Other
- 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:D67N 91% cov=14,578 • K70R 90% cov=14,408 • K219Q 94% cov=17,071

NNRTI Mutations:K103N 97% cov=14,255 • E138A 90% cov=16,914 • P225H 94% cov=16,517

RT Other Mutations:E6G 92% cov=16,727 • V8VI I: 75%, V: 23% cov=17,657 • K20R 94% cov=16,408 • V35T 98% cov=15,654 • V60I 97% cov=15,282 • K101Q 90% cov=14,155 • K122E 99% cov=15,178 • D123S 90% cov=14,442 • I135T 97% cov=16,726 • K173A 98% cov=21,800 • Q174K 97% cov=21,804 • T200A 98% cov=20,685 • Q207A 98% cov=17,054 • R211S 96% cov=18,117 • F214L 94% cov=17,258 • V245T 91% cov=11,331 • D250E 94% cov=10,918 • T286A 93% cov=7,172 • E291D 88% cov=6,517 • V292VI I: 87%, V: 12% cov=6,518 • I293V 99% cov=6,308 • P294T 89% cov=6,505 • E300EV V: 53%, E: 46% cov=6,262 • E312ED D: 87%, E: 10% cov=5,840

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

RT comments

- NRTI
- D67N is a non-polymorphic TAM associated with low-level resistance to AZT.
  - K70R is a TAM that confers intermediate resistance to AZT and contributes to reduced ABC and TDF susceptibility in combination with other TAMs.
  - 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.
  - E138A is a common polymorphic accessory mutation weakly selected in persons receiving ETR and RPV. It reduces ETR and RPV susceptibility ~2-fold. Its effect on ETR- and RPV-containing regimens is likely to be minimal.
  - 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
- K101Q is a relatively non-polymorphic mutation that is weakly selected in persons receiving NVP and EFV. It is of uncertain phenotypic and clinical significance.
  - 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 ⚡	FTC ⚡	3TC ⚡	TDF ⚡
<u>D67N</u>	5	15	0	0	5
<u>D67N + K70R + K219Q</u>	10	15	10	10	10
<u>K70R</u>	5	30	0	0	5
<u>K219Q</u>	5	10	0	0	5
Total	25	70	10	10	25

Drug resistance mutation scores of NNRTI:

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Rule	DOR ⚡	EFV ⚡	ETR ⚡	NVP ⚡	RPV ⚡
<u>K103N + P225H</u>	10	0	0	0	0
<u>P225H</u>	20	45	0	45	0
<u>K103N</u>	0	60	0	60	0
<u>E138A</u>	0	0	10	0	15
Total	30	105	10	105	15