

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:[L10G](#) • [V11\\*](#) • T12V • I13S • K14N • [I15D](#) • G16R • Q18M • K20Q • L33V • M36I • R41K • L63P • I64V • I72V

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

PR comments

Other

- L10F is a common non-polymorphic, PI-selected accessory mutation associated with reduced in vitro susceptibility to LPV and DRV. L10I/V are polymorphic, PI-selected accessory mutations that increase the replication of viruses with other PI-resistance mutations. L10R/Y are rare, non-polymorphic PI-selected mutations. Their effects on PI susceptibility have not been well studied. **L10G** is a highly unusual mutation at this position.
- L33I/V** are minimally polymorphic mutations that do not appear to be selected by PIs or to reduce their susceptibility.

No drug resistance mutations were found for PI.

NRTI Mutations:[K70Q](#) • [M184I](#)

NNRTI Mutations:[K101E](#) • [K103N](#) • [G190A](#)

RT Other Mutations:V35I • T39E • V60I • V90I • S105T • D121Y • K122E • I135K • Q174R • D177E • I178V • T200I • Q207E • R211K • [K219X](#) • [P225X](#) • P226S • L228R • P236S • [L246T](#) • P247A • E248R • D250E • [N255M](#) • [D256I](#) • [I257Y](#) • Q258R • [K259V](#) • L260V • [V261E](#) • [G262N](#) • [L264W](#)

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

RT comments

NRTI

- K70E/Q/N/T/S/G** cause low-leve resistance to 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.

NNRTI

- K101E** is a non-polymorphic accessory mutation that confers intermediate resistance to NVP and RPV and low-level reductions in susceptibility to EFV, ETR, and DOR when it occurs with other NNRTI-resistance mutations.
- K103N** is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.
- G190A** is a non-polymorphic mutation that causes high-level resistance to NVP and intermediate resistance to EFV. It does not significantly reduce susceptibility to RPV, ETR, or DOR.

Other

- V90I** is a polymorphic accessory mutation weakly selected by each of the NNRTIs. It is associated with minimal, if any, detectable reduction in NNRTI susceptibility.
- P236L is a rare mutation selected commonly by DLV, which appears to have little if any effect on current NNRTIs. **P236S** is a highly unusual mutation at this position.

Drug resistance mutation scores of NRTI:

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Rule	ABC	AZT	D4T	DDI	FTC	3TC	TDF
<u>K70Q</u>	15	0	15	15	10	10	15
<u>M184I</u>	15	-10	-10	10	60	60	-10
<u>K70Q + M184I</u>	0	0	10	0	0	0	10
Total	30	-10	15	25	70	70	15

Drug resistance mutation scores of NNRTI:

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Rule	DOR	EFV	ETR	NVP	RPV
<u>K101E</u>	15	15	15	30	45
<u>K101E + G190A</u>	5	0	5	0	0
<u>K103N</u>	0	60	0	60	0
<u>G190A</u>	0	45	10	60	15
<u>K101E + M184I</u>	0	0	0	0	15
Total	20	120	30	150	75