

PI Major Mutations:None

PI Accessory Mutations:[K20T](#)

PR Other Mutations:T12M • I13S • K14V • [I15D](#) • [G16S](#) • Q18K • E35D • M36I • R41K • K43R • R57K • [Y59\\*](#) • [D60V](#) • H69K • [N83X](#) • L89M

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>	Low-Level Resistance
<b>saquinavir/r (SQV/r)</b>	Susceptible
<b>tipranavir/r (TPV/r)</b>	Susceptible

PR comments

Accessory

- K20T** is a non-polymorphic accessory PI-selected mutation associated with reduced susceptibility to ATV and LPV.

Drug resistance mutation scores of PI:

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Rule	ATV/r ⚡	DRV/r ⚡	FPV/r ⚡	IDV/r ⚡	LPV/r ⚡	NFV ⚡	SQV/r ⚡	TPV/r ⚡
<a href="#">K20T</a>	5	0	5	5	0	15	5	0

NRTI Mutations:[M184V](#)

NNRTI Mutations:[K103N](#)

RT Other Mutations:[P1Q](#) • V35T • P55S • V60I • K122E • D123N • I135T • K173A • Q174K • D177E • I178V • [T200X](#) • Q207K • R211K • F214L • [K219X](#) • K238Q • [Q242X](#) • V245Q • P247L • D250N • S251T • [T253D](#) • [V254C](#) • [N255H](#) • I257L • K259N • [L264S](#) • [N265T](#) • [W266R](#) • [A267V](#) • I274L • [K275X](#) • [L283S](#)

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

Other

- K238T/N** are uncommon non-polymorphic mutations selected in persons receiving NVP and EFV usually in combination with K103N. Alone, **K238T/N** appear to have minimal effects on NNRTI susceptibility. **K238Q** 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>M184V</u>	15	-10	-10	10	60	60	-10

Drug resistance mutation scores of NNRTI:

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Rule	DOR	EFV	ETR	NVP	RPV
<u>K103N</u>	0	60	0	60	0