

PI Major Mutations:None

PI Accessory Mutations:**L33F**

93%  
cov=37,290

PR Other Mutations:**I13V**

99%  
cov=29,322

 • **K14R**

94%  
cov=29,347

 • **G16E**

95%  
cov=30,660

 • **M36I**

99%  
cov=37,611

 • **N37ND**

N: 73%, D: 26%  
cov=37,909

 • **P39S**

93%  
cov=37,650

 • **R57K**

96%  
cov=35,557

 • **D60E**

93%  
cov=32,626

 • **E65D**

89%  
cov=28,610

 • **H69K**

96%  
cov=28,512

 • **L89M**

98%  
cov=38,476

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

PR comments

Accessory

- L33F** is a relatively non-polymorphic accessory mutation selected by each of the PIs. In combination with other PI-resistance mutations, it is associated with reduced susceptibility to LPV, ATV, and DRV.

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 ⚡
<u>L33F</u>	5	5	10	5	5	10	5	10

NRTI Mutations:**D67N**

92%  
cov=12,556

 • **K70R**

95%  
cov=12,965

 • **M184V**

89%  
cov=6,417

 • **T215FI**

I: 63%, F: 27%  
cov=8,149

 • **K219E**

92%  
cov=8,493

NNRTI Mutations:**L100LI**

I: 81%, L: 17%  
cov=13,089

 • **K103N**

99%  
cov=13,942

RT Other Mutations:**E6N**

90%  
cov=16,029

 • **V35T**

94%  
cov=15,051

 • **T39TL**

T: 53%, L: 29%  
cov=14,950

 • **E40ED**

D: 53%, E: 44%  
cov=14,955

 • **K49R**

93%  
cov=13,864

 • **V60I**

99%  
cov=13,708

 • **K102KR**

K: 63%, R: 36%  
cov=13,172

 • **K122E**

96%  
cov=10,444

 • **D123N**

91%  
cov=10,434

 • **I135T**

95%  
cov=8,681

 • **K166T**

89%  
cov=6,690

 • **K173S**

86%  
cov=6,215

 • **Q174K**

79%  
cov=6,217

 • **D177E**

92%  
cov=6,386

 • **V179I**

77%  
cov=6,343

 • **T200A**

96%  
cov=7,309

 • **I202V**

92%  
cov=7,441

 • **E203EK**

E: 76%, K: 18%  
cov=7,432

 • **Q207D**

85%  
cov=7,565

 • **L210F**

85%  
cov=7,440

 • **R211K**

92%  
cov=7,437

 • **D218E**

91%  
cov=6,497

 • **L228R**

88%  
cov=9,068

 • **V245E**

95%  
cov=15,713

 • **E248D**

95%  
cov=16,036

 • **K249KR**

K: 51%, R: 49%  
cov=16,037

 • **D250E**

96%  
cov=16,040

 • **L283I**

96%  
cov=25,387

 • **I293V**

99%  
cov=26,766

 • **P294T**

92%  
cov=26,780

 • **L295M**

91%  
cov=26,777

 • **V314VA**

V: 75%, A: 23%  
cov=29,002

 • **G335D**

97%  
cov=257

 • **R356K**

89%  
cov=227

 • **M357K**

94%  
cov=227

 • **G359S**

94%  
cov=215

 • **K366R**

92%  
cov=181

 • **T369A**

93%  
cov=179

 • **E370A**

92%  
cov=176

 • **A371V**

93%  
cov=176

 • **I375IV**

V: 86%, I: 12%  
cov=168

 • **A376V**

93%  
cov=265

 • **T377M**

94%  
cov=165

Nucleoside Reverse Transcriptase Inhibitors		Non-nucleoside Reverse Transcriptase Inhibitors	
<b>abacavir (ABC)</b>	High-Level Resistance	<b>doravirine (DOR)</b>	Intermediate Resistance
<b>zidovudine (AZT)</b>	High-Level Resistance	<b>efavirenz (EFV)</b>	High-Level Resistance
<b>stavudine (D4T)</b>	High-Level Resistance	<b>etravirine (ETR)</b>	Intermediate Resistance
<b>didanosine (DDI)</b>	High-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>	Intermediate 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.
- **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.
- **T215Y/F** are TAMs that causes intermediate/high-level resistance to AZT and potentially low-level resistance to ABC and TDF.
- T215Y/F are TAMs that causes intermediate/high-level resistance to AZT and potentially low-level resistance to ABC and TDF. **T215S/C/D/E/I/V/N/A/L** do not reduce NRTI susceptibility but arise from viruses that once contained T215Y/F. The presence of one of these revertant mutations suggests that the patient may have once been infected with a virus containing T215Y/F.
- **K219E/Q/N/R** are accessory TAMS that usually occur in combination with multiple other TAMs.

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.

Mutation scoring: RT

HIVDB 9.5.1 (2023-11-05)

Drug resistance mutation scores of NRTI:

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Rule	ABC ↕	AZT ↕	D4T ↕	DDI ↕	FTC ↕	3TC ↕	TDF ↕
<u>D67N</u>	5	15	15	5	0	0	5
<u>D67N + K70R + M184V + K219E</u>	10	0	0	0	0	0	0
<u>D67N + K70R + K219E</u>	10	15	10	10	10	10	10
<u>D67N + T215FI + K219E</u>	5	5	5	5	0	0	5
<u>K70R</u>	5	30	15	10	0	0	5
<u>M184V</u>	15	-10	-10	10	60	60	-10
<u>T215FI</u>	10	60	40	15	0	0	10
<u>K219E</u>	5	10	10	5	0	0	5
<u>K70R + T215FI</u>	0	0	5	5	0	0	0
Total	65	125	90	65	70	70	30

Drug resistance mutation scores of NNRTI:

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Rule	DOR ↕	EFV ↕	ETR ↕	NVP ↕	RPV ↕
<u>L100LI</u>	15	60	30	60	60
<u>L100LI + K103N</u>	15	0	0	0	0
<u>K103N</u>	0	60	0	60	0
Total	30	120	30	120	60