

Drug resistance interpretation: PRHIVDB 9.5.1 (2023-11-05)

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:

L10V99%
seen:2,218

•

I13V99%
seen:2,332

•

I15V99%
seen:2,352

•

G16A99%
seen:2,334

•

E35D99%
seen:3,475

•

M36I99%
seen:3,475

•

N37D99%
seen:3,475

•

R41K99%
seen:3,543

•

K43R99%
seen:3,538

•

R57K99%
seen:3,434

•

D60DE99%
seen:3,434

•

G72V,R74V99%
seen:3,434

•

H69T99%
seen:3,314

•

L89M99%
seen:3,380

Protease Inhibitors

atazanavir/r (ATV/r)

Susceptible

darunavir/r (DRV/r)

Susceptible

fosamprenavir/r (FPV/r)

Susceptible

indinavir/r (IDV/r)

Susceptible

lopinavir/r (LPV/r)

Susceptible

nelfinavir (NFV)

Susceptible

saquinavir/r (SQV/r)

Susceptible

tipranavir/r (TPV/r)

Susceptible

PR comments

Other

- L10I/V are polymorphic, PI-selected accessory mutations that increase the replication of viruses with other PI-resistance mutations.

Mutation scoring: PRHIVDB 9.5.1 (2023-11-05)

No drug resistance mutations were found for PI.

Drug resistance interpretation: RTHIVDB 9.5.1 (2023-11-05)

NRTI Mutations:

M184V99%
seen:712

NNRTI Mutations:

K103KN99%
seen:1,864

•

E138G99%
seen:1,714

•

G190GA99%
seen:1,829

RT Other Mutations:

K11A99%
seen:2,232

•

K20R99%
seen:3,302

•

V21I99%
seen:3,302

•

T27S99%
seen:3,403

•

K32KN99%
seen:3,305

•

R32V,R34V99%
seen:3,305

•

V35T99%
seen:3,494

•

T39R99%
seen:3,334

•

K122E99%
seen:3,335

•

D123N99%
seen:3,335

•

I135T99%
seen:3,333

•

K166KR99%
seen:3,342

•

K173S99%
seen:3,352

•

Q174K99%
seen:3,352

•

D177E99%
seen:3,352

•

V179I99%
seen:4,713

•

I202V99%
seen:3,333

•

Q207A99%
seen:3,344

•

R211K99%
seen:3,333

•

F214FL99%
seen:3,333

•

R215V99%
seen:3,333

Nucleoside Reverse Transcriptase Inhibitors

abacavir (ABC)

Low-Level Resistance

zidovudine (AZT)

Susceptible

stavudine (D4T)

Susceptible

didanosine (DDI)

Potential Low-Level Resistance

emtricitabine (FTC)

High-Level Resistance

lamivudine (3TC)

High-Level Resistance

tenofovir (TDF)

Susceptible

Non-nucleoside Reverse Transcriptase Inhibitors

doravirine (DOR)

Susceptible

efavirenz (EFV)

High-Level Resistance

etravirine (ETR)

Low-Level Resistance

nevirapine (NVP)

High-Level Resistance

rilpivirine (RPV)

Intermediate Resistance

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.
- E138Q/G are non-polymorphic accessory mutations selected by ETR occasionally NVP and EFV. They cause low-level reductions in susceptibility to NVP, RPV, and ETR.
- 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

- 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.
- This virus is predicted to have intermediate-level reduced susceptibility to RPV. The use of the combination of CAB/ RPV should be considered to be contraindicated.

Mutation scoring: RTHIVDB 9.5.1 (2023-11-05)

Drug resistance mutation scores of NRTI:

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Rule	ABC	AZT	D4T	DDI	FTC	3TC	TDF
M184V	15	-10	-10	10	60	60	-10

Drug resistance mutation scores of NNRTI:

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Rule	DOR	EFV	ETR	NVP	RPV
K103KN	0	60	0	60	0
E138G	0	10	10	10	15
G190GA	0	45	10	60	15
Total	0	115	20	130	30