

Drug resistance interpretation: PR

HVDB 9.5.1 (2023-11-05)

PI Major Mutations:None

PI Accessory Mutations:None

PR Other Mutations:L10LV100%
seen(8,229) • K14KR91%
seen(5,282) • L19I90%
seen(5,285) • L33LV100%
seen(7,283) • M36MI100%
seen(5,259) • R41K100%
seen(5,286) • I72IV100%
seen(5,382)

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.
- L33I/V are minimally polymorphic mutations that do not appear to be selected by PIs or to reduce their susceptibility.

Mutation scoring: PR

HVDB 9.5.1 (2023-11-05)

No drug resistance mutations were found for PI.

Drug resistance interpretation: RT

HVDB 9.5.1 (2023-11-05)

NRTI Mutations:None

NNRTI Mutations:[K103N](#)100%
seen(1,803)

RT Other Mutations:V10A100%
seen(2,788) • K11KT100%
seen(2,762) • K14R100%
seen(2,762) • V35T100%
seen(2,549) • T39A99%
seen(2,587) • E40D100%
seen(2,539) • K49R100%
seen(2,282) • V60I100%
seen(2,587) • K102Q100%
seen(1,754) • D121Y100%
seen(1,592) • K122E100%
seen(1,592) • T16SI100%
seen(2,544) • D17TE100%
seen(2,553) • I178IM100%
seen(2,535) • T200A100%
seen(2,581) • Q20TE100%
seen(2,786) • V245N100%
seen(151) • E248D100%
seen(729) • D250E100%
seen(586) • A554N99%
seen(48) • V559I100%
seen(1,25)

Nucleoside Reverse Transcriptase Inhibitors

abacavir (ABC)	Susceptible
zidovudine (AZT)	Susceptible
stavudine (D4T)	Susceptible
didanosine (DDI)	Susceptible
emtricitabine (FTC)	Susceptible
lamivudine (3TC)	Susceptible
tenofovir (TDF)	Susceptible

Non-nucleoside Reverse Transcriptase Inhibitors

doravirine (DOR)	Susceptible
efavirenz (EFV)	High-Level Resistance
etravirine (ETR)	Susceptible
nevirapine (NVP)	High-Level Resistance
rilpivirine (RPV)	Susceptible

RT comments

NNRTI

- K103N is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.

Mutation scoring: RT

HVDB 9.5.1 (2023-11-05)

No drug resistance mutations were found for NRTI.

Drug resistance mutation scores of NNRTI:

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Rule	DOR	EFV	ETR	NVP	RPV
K103N	0	60	0	60	0

Drug resistance interpretation: IN

HVDB 9.5.1 (2023-11-05)

INSTI Major Mutations:None

INSTI Accessory Mutations:None

IN Other Mutations:K14R100%
seen(257) • L45V100%
seen(528) • I72V100%
seen(585) • T112V100%
seen(247) • I113V100%
seen(287) • S119T100%
seen(286) • H171Q100%
seen(275) • V201I100%
seen(382) • L234I100%
seen(354) • A265V97%
seen(479) • S283G100%
seen(435)

Integrase Strand Transfer Inhibitors

bictegravir (BIC)	Susceptible
cabotegravir (CAB)	Susceptible
dolutegravir (DTG)	Susceptible
elvitegravir (EVG)	Susceptible
raltegravir (RAL)	Susceptible

Mutation scoring: IN

HVDB 9.5.1 (2023-11-05)

No drug resistance mutations were found for INSTI.