#### Drug resistance interpretation: PR HIVDB 9.5.1 (2023-11-05)

PI Major Mutations: None

PI Accessory Mutations: None

PR Other Mutations: 

## 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 saguinavir/r (SQV/r) Susceptible tipranavir/r (TPV/r) Susceptible

# Mutation scoring: PR

Drug resistance interpretation: RT

HIVDB 9.5.1 (2023-11-05)

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No drug resistance mutations were found for PI.

NRTI Mutations: None NNRTI Mutations: K103N ---

RT Other Mutations: 

# Nucleoside Reverse Transcriptase Inhibitors

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

## Non-nucleoside Reverse Transcriptase Inhibitors

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

## RT comments

# NNRTI

Other

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

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)

No drug resistance mutations were found for NRTI.

Drug resistance mutation scores of NNRT1:				Download CSV	
Rule	DOR ÷	EFV ≑	ETR ≑	NVP ≑	RPV :
K103N	0	60	0	60	0

# Drug resistance interpretation: IN

HIVDB 9.5.1 (2023-11-05)

INSTI Major Mutations: None None INSTI Accessory Mutations:

IN Other Mutations:

K14R - V32I - S39N - L45V - M50I - I72V - T112V - T112V - T124A - T125A - V126VF - G134D - I35V - D167E - V201I - K211R - N22K - L234I - S283G - C125A - C125A

# Integrase Strand Transfer Inhibitors

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

### IN comments

# Other

M50I is a highly polymorphic mutation, which has a prevalence of 3% to 34% in INSTI-naïve persons depending on subtype. It has been selected in vitro by DTG and BIC in combination with R263K. It may contribute to reduced DTG and CAB susceptibility in combination with R263K.

No drug resistance mutations were found for INSTI.

Mutation scoring: IN HIVDB 9.5.1 (2023-11-05)