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

PI Major Mutations: None

PI Accessory Mutations: None PR Other Mutations: K14R ... • E35D ... • M36I ... • R41K ... • H69K ... • L89M ...

Protease Inhibitors

atazanavir/r (ATV/r) Susceptible darunavir/r (DRV/r) Susceptible Susceptible fosamprenavir/r (FPV/r) indinavir/r (IDV/r) Susceptible lopinavir/r (LPV/r) Susceptible Susceptible nelfinavir (NFV) saquinavir/r (SQV/r) Susceptible Susceptible tipranavir/r (TPV/r)

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

No drug resistance mutations were found for PI.

Drug resistance interpretation: RT

None NRTI Mutations: NNRTI Mutations: None

RT Other Mutations:

Nucleoside Reverse Transcriptase Inhibitors

abacavir (ABC) Susceptible zidovudine (AZT) Susceptible stavudine (D4T) Susceptible Susceptible didanosine (DDI) emtricitabine (FTC) Susceptible lamivudine (3TC) Susceptible tenofovir (TDF) Susceptible Non-nucleoside Reverse Transcriptase Inhibitors

doravirine (DOR) Susceptible efavirenz (EFV) Susceptible Susceptible etravirine (ETR) Susceptible nevirapine (NVP) rilpivirine (RPV) Susceptible

RT comments

Other

Mutation scoring: RT

. 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.

No drug resistance mutations were found for NRTI.

No drug resistance mutations were found for NNRTI.

INSTI Major Mutations: None

INSTI Accessory Mutations: None

Drug resistance interpretation: IN

IN Other Mutations: K14R === * V31l === * V31l === * V31l === * V20ll === * T218S == * T228S == * T128S == * T228S == * T228S == * V20ll === * V20ll == *

Integrase Strand Transfer Inhibitors

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

IN comments

Mutation scoring: IN

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.

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