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

PI Major Mutations: None PI Accessory Mutations: None

PR Other Mutations: 616E ... • E35D ... • M36V ... • R41K ... • R57K ... • L63C ... • H69K ... • L89M ...

Protease Inhibitors

 atazanavir/r (ATV/r)
 Susceptible

 darunavir/r (DRV/r)
 Susceptible

 lopinavir/r (LPV/r)
 Susceptible

Mutation scoring; PR

No drug resistance mutations were found for PI.

Drug resistance interpretation: RT

NRTI Mutations: Nor

NNRTI Mutations: K103N White V106M White P225H White

RT Other Mutations: 15V sm • V35T sm • V601 sm • V122E sm • D123N sm • V179L sm • V179L sm • V245E sm • V245E

Nucleoside Reverse Transcriptase Inhibitors Non-nucleoside Reverse Transcriptase Inhibitors doravirine (DOR) abacavir (ABC) Susceptible High-Level Resistance zidovudine (AZT) Susceptible efavirenz (EFV) High-Level Resistance emtricitabine (FTC) Susceptible etravirine (ETR) Susceptible lamivudine (3TC) Susceptible nevirapine (NVP) High-Level Resistance tenofovir (TDF) Susceptible 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.
- V106M is a non-polymorphic mutation that confers high-level resistance to NVP and EFV. It is selected in vitro and in vivo by DOR and preliminary data suggests it reduces DOR susceptibility about 3-fold.
- P225H is a non-polymorphic EFV-selected mutation that usually occurs in combination with K103N. The combination of P225H and K103N synergistically reduces NVP, EFV and DOR susceptibility.

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

Total

No drug resistance mutations were found for NRTL

60

Drug resistance mutation scores of NNRTI:

-						
Rule	DOR ÷	EFV ≑	ETR ÷	NVP ≑	RPV ÷	
K103N + P225H	10	0	0	0	0	
V106M	30	60	0	60	0	

45

165

0

45 60

165

0

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