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

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INSTI Major Mutations: T66TA • N155NH

INSTI Accessory Mutations:

IN Other Mutations: L1011 • T1121 • V201VI • K211Q • I217V • K240KR • K266KR

Integrase Strand Transfer Inhibitors

bictegravir (BIC) Low-Level Resistance cabotegravir (CAB) Intermediate Resistance dolutegravir (DTG) Low-Level Resistance elvitegravir (EVG) High-Level Resistance raltegravir (RAL) High-Level Resistance

IN comments

- T66A/I are non-polymorphic mutations selected in persons receiving EVG, RAL, and DTG usually in combination with other INSTI-resistance mutations. They cause moderate reductions in EVG susceptibility but do not appear to reduce susceptibility to other INSTIs.
- . N155H is a common nonpolymorphic INSTI-resistance mutations. It has been reported in a high proportion of persons developing VF and CAB. Alone, it reduces RAL and EVG susceptibility about 10 and 30-fold, respectively. It has minimal effect on susceptibility to DTG, BIC, and CAB.

Accessory

- HS1Y is an uncommon nonpolymorphic accessory mutation selected in vitro by EVG, DTG, and CAB. Alone, it has minimal if any effect on INSTI susceptibility.
- . This virus is predicted to have intermediate-level reduced susceptibility to CAB. The use of the combination of CAB/RPV should be considered to be contraindicated.
- . There is evidence for low-level DTG resistance. If DTG is used, it should be administered twice daily.

Mutation scoring: IN

Drug resistance mutation scores of INSTI:

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BIC ≑	CAB ≑	DTG ÷	EVG ‡	RAL :
10	15	10	15	15
10	25	10	60	60
0	0	0	60	15
20	40	20	135	90
	10 10 0	10 15 10 25 0 0	10 15 10 10 25 10 0 0 0	10 15 10 15 10 25 10 60 0 0 0 60