

NRTI Mutations:	None																								
NNRTI Mutations:	None																								
RT Other Mutations:	K512R <small>100% pos=5,279</small> • L517I <small>99% pos=5,750</small> • S519N <small>99% pos=5,720</small> • Q520S <small>97% pos=5,720</small> • Q524K <small>100% pos=5,975</small> • K527E <small>100% pos=7,295</small> • A534S <small>100% pos=7,900</small> • A554S <small>100% pos=7,812</small>																								
<div><div>Nucleoside Reverse Transcriptase Inhibitors</div><div><table><tr><td>abacavir (ABC)</td><td>Susceptible</td></tr><tr><td>zidovudine (AZT)</td><td>Susceptible</td></tr><tr><td>stavudine (D4T)</td><td>Susceptible</td></tr><tr><td>didanosine (DDI)</td><td>Susceptible</td></tr><tr><td>emtricitabine (FTC)</td><td>Susceptible</td></tr><tr><td>lamivudine (3TC)</td><td>Susceptible</td></tr><tr><td>tenofovir (TDF)</td><td>Susceptible</td></tr></table></div></div> <div><div>Non-nucleoside Reverse Transcriptase Inhibitors</div><div><table><tr><td>doravirine (DOR)</td><td>Susceptible</td></tr><tr><td>efavirenz (EFV)</td><td>Susceptible</td></tr><tr><td>etravirine (ETR)</td><td>Susceptible</td></tr><tr><td>nevirapine (NVP)</td><td>Susceptible</td></tr><tr><td>rilpivirine (RPV)</td><td>Susceptible</td></tr></table></div></div>		abacavir (ABC)	Susceptible	zidovudine (AZT)	Susceptible	stavudine (D4T)	Susceptible	didanosine (DDI)	Susceptible	emtricitabine (FTC)	Susceptible	lamivudine (3TC)	Susceptible	tenofovir (TDF)	Susceptible	doravirine (DOR)	Susceptible	efavirenz (EFV)	Susceptible	etravirine (ETR)	Susceptible	nevirapine (NVP)	Susceptible	rilpivirine (RPV)	Susceptible
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No drug resistance mutations were found for NRTI.

No drug resistance mutations were found for NNRTI.

INSTI Major Mutations:	None										
INSTI Accessory Mutations:	None										
IN Other Mutations:	D3DG <small>0-67% pos=0,001</small> • L14R <small>99% pos=5,760</small> • M50I <small>100% pos=5,500</small> • K71R <small>99% pos=2,360</small> • I72V <small>100% pos=2,362</small> • L101I <small>100% pos=1,905</small> • T112V <small>100% pos=2,000</small> • I113V <small>99% pos=2,000</small> • T124A <small>99% pos=3,175</small> • T125A <small>99% pos=3,276</small> • G134D <small>99% pos=3,960</small> • K136Q <small>99% pos=3,958</small> • D167E <small>100% pos=3,811</small>										
<div><div>Integrase Strand Transfer Inhibitors</div><div><table><tr><td>bictegravir (BIC)</td><td>Susceptible</td></tr><tr><td>cabotegravir (CAB)</td><td>Susceptible</td></tr><tr><td>dolutegravir (DTG)</td><td>Susceptible</td></tr><tr><td>elvitegravir (EVG)</td><td>Susceptible</td></tr><tr><td>raltegravir (RAL)</td><td>Susceptible</td></tr></table></div></div>		bictegravir (BIC)	Susceptible	cabotegravir (CAB)	Susceptible	dolutegravir (DTG)	Susceptible	elvitegravir (EVG)	Susceptible	raltegravir (RAL)	Susceptible
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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.