PatientID: GU121699

Okitobba 06, 2023

## Color Code

HR: High-Level Resistance

PLR: Potential Low-Level F
IR: Intermediate Resistance PLR: Potential Low-Level Resistance

S: Susceptible

DRUG.CLASS	DRUG	RESISTANCE.PROFILE	DRMS.above.20.percent.prevalence
	BIC	LR	
	CAB	IR	
INSTI	DTG	$_{ m LR}$	N155NH;T66TA;H51HY
	EVG	HR	
	RAL	HR	

## Appendix

## Drug abbreviations in full

DRUG.CLASS	ABBREVIATION	DRUG.NAME
	ATV	Atazanavir
	DRV	Darunavir
	FPV	Fosamprenavir
PI	IDV	Indinavir
11	LPV	Lopinavir
	NFV	Nelfinavir
	SQV	Saquinavir
	TPV	Tipranavir
	ABC	Abacavir
	AZT	Azidothymidine
	DFT	Stavudine
NRTI	DDI	Didanosine
	FTC	Emtricitabine
	LMV	Lamivudine
	TDF	Tenofovir
	DOR	Doravirine
	EFV	Efavirenz
NNRTI	ETR	Etravirine
	NVP	Nevirapine
	RPV	Rilpivirine
	BIC	Bictegravir
	CAB	Cabotegravir
INSTI	DTG	Dolutegravir
	EVG	Elvitegravir
	RAL	Raltegravir

## Comments

DRUG.CLASS	COMMENTS	
PI		
NRTI		
NNRTI		
	H51Y is an uncommon nonpolymorphic accessory mutation selected in vitro by EVG,	
	DTG, and CAB. Alone, it has minimal if any effect on INSTI susceptibility.	
	N155H is a common nonpolymorphic INSTI-resistance mutations. It has been reported in	
	a high proportion of persons developing VF and HIVDR while receiving RAL, EVG, DTG,	
	and CAB. Alone, it reduces RAL and EVG susceptibility about 10 and 30-fold,	
INSTI	respectively. It has minimal effect on susceptibility to DTG, BIC, and CAB.	
	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.	