PatientID: HIVDR-765-23

Sebuttemba 27, 2023

Color Code

HR: High-Level Resistance

LR: Low-Level Resistance

IR: Intermediate Resistance

S: Susceptible

DRUG.CLASS	DRUG	RESISTANCE.PROFILE	DRMS.above.20.percent.prevalence
PI	ATV	\mathbf{S}	
	DRV	\mathbf{S}	
	FPV	\mathbf{S}	
	IDV	\mathbf{S}	
	LPV	\mathbf{S}	
	NFV	\mathbf{S}	
	SQV	\mathbf{S}	
	TPV	${f S}$	
NRTI	ABC	IR	
	AZT	IR	
	D4T	IR	
	DDI	IR	M184MV;F77LF;M41L;K219R;V75M
	FTC	$_{ m HR}$	
	LMV	$_{ m HR}$	
	TDF	${f S}$	
NNRTI	DOR	$_{ m HR}$	
	EFV	$_{ m HR}$	
	ETR	LR	A98G;V108I;H221Y;P225H;K103N
	NVP	$_{ m HR}$	
	RPV	IR	
	BIC	S	
INSTI	CAB	${f S}$	
	DTG	${f S}$	
	EVG	${f S}$	
	RAL	${f S}$	

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			
	F77L usually occurs in combination with the multi-NRTI resistance mutation Q151M. When it occurs alone, its clinical significance is uncertain.		
	K219E/Q/N/R are accessory TAMS that usually occur in combination with multiple other		
	TAMs.		
	M184V/I cause high-level in vitro resistance to 3TC and FTC and low/intermediate		
	resistance to ABC (3-fold reduced susceptibility). M184V/I are not contraindications to		
	continued treatment with 3TC or FTC because they increase susceptibility to AZT and		
	TDF and are associated with clinically significant reductions in HIV-1 replication.		
	M41L is a TAM that usually occurs with T215Y. In combination, M41L plus T215Y confer		
NRTI	intermediate / high-level resistance to AZT and d4T and contribute to reduced ddI, ABC		
	and TDF susceptibility.		
	V75T/M/A/S are nonpolymorphic accessory NRTI-selected mutations. They appear to		
	have minimal phenotypic effects on AZT, ABC, and TDF.		
	A98G is a non-polymorphic accessory mutation associated with low-level reduced susceptibility to each of the NNRTIs.		
	H221Y is a non-polymorphic accessory mutation selected primarily by NVP, RPV, and		
	DOR. It frequently occurs in combination with Y181C.		

NNRTI	K103N is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM. 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. V108I is a relatively non-polymorphic accessory mutation selected in vitro and/or in vivo with each of the NNRTIs. It appears to contribute to reduced susceptibility to most NNRTIs only in combination with other NNRTI-resistance mutations.
INSTI	