PatientID: HDR44

Okitobba 06, 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	S	
	DRV	\mathbf{S}	
	FPV	\mathbf{S}	
	IDV	${f S}$	
	LPV	${f S}$	
	NFV	\mathbf{S}	
	SQV	\mathbf{S}	
	TPV	\mathbf{S}	
NRTI	ABC	$^{ m HR}$	
	AZT	${f S}$	
	D4T	$_{ m HR}$	
	DDI	$_{ m HR}$	K65R;M184V;K219N;V75M
	FTC	$_{ m HR}$	
	LMV	$_{ m HR}$	
	TDF	IR	
NNRTI	DOR	PLR	
	EFV	$_{ m HR}$	
	ETR	$_{ m LR}$	V106I;K103N;G190A
	NVP	$_{ m HR}$	
	RPV	LR	

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	
	K219E/Q/N/R are accessory TAMS that usually occur in combination with multiple other
	TAMs.
	K65R confers intermediate reductions in susceptibility to TDF, ABC, and 3TC/FTC. It
	increases AZT susceptibility. In NRTI-experienced, INSTI-naive patients with K65R,
	TDF+3TC+DTG is usually highly effective and more effective than AZT/3TC/DTG.
	However, in patients receiving TDF+3TC+DTG, there is a risk of emergent DTG
	resistance that does not arise in NRTI-naive patients receiving TDF+3TC+DTG.
NRTI	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.
	V75T/M/A/S are nonpolymorphic accessory NRTI-selected mutations. They appear to
	have minimal phenotypic effects on AZT, ABC, and TDF.
	G190A is a non-polymorphic mutation that causes high-level resistance to NVP and
	intermediate resistance to EFV. It does not significantly reduce susceptibility to RPV,
	ETR, or DOR.

NNRTI	K103N is a non-polymorphic mutation that confers high-level reductions in NVP and EFV susceptibility. It is the most commonly transmitted DRM.
	V106I occurs in 1% to 2% of viruses from untreated persons. It contributes to reduced
	NNRTI susceptibility only in combination with other NNRTI-resistance mutations. It is
	commonly selected in persons receiving DOR in combination with mutations at position
	227.
INSTI	