

Pulmonary Anti-Hypertensives Treatment Selector

Charts produced November 2017. Full information available at www.hiv-druginteractions.org

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		ATV/r	DRV/r	LPV/r	EFV	ETV	NVP	RPV	MVC	DTG	RAL	ABC	FTC	3TC	TDF	ZDV	E/C/F/TAF	E/C/F/TDF
	Ambrisentan	1	1	1	\leftrightarrow													
Z d	Bosentan	↑ª	↑ª	↑ª	\	↓	↑p	↓	↓	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	↑ª	↑ª
	Macitentan	1	1	1	\	↓	↓	\leftrightarrow	1	1								
	Sildenafil	1	1	1	\	↓	↓	\leftrightarrow	1	1								
G	Tadalafil	1	1	1	\	↓	↓	\leftrightarrow	1	1								
5	Riociguat	1	1	1	1	1	1	\leftrightarrow	1	1								
	Epoprostenol	\leftrightarrow																
Δq	lloprost	\leftrightarrow																
	Treprostinil	1	1	1	1	\leftrightarrow												
P	Selexipag	↔c	↔c	↔c	\leftrightarrow	↔ ^c	↔°											

Colour Legend

	No clinically significant interaction expected.
	These drugs should not be coadministered.
	Potential interaction which may require a dosage adjustment or close monitoring.
	Potential interaction predicted to be of weak intensity. No a priori dosage adjustment is recommended.

Text Legend

- Potential increased exposure of the pulmonary antihypertensive
- Potential decreased exposure of HIV drug Potential decreased exposure of the pulmonary antihypertensive
- No significant effect

ERA Endothelin receptor antagonist PDE5 Phosphodiesterase 5 inhibitors sCG Soluble guanylate cyclase stimulators

Prostacyclin analogues

- When starting bosentan in individuals already on ritonavir or cobicistat containing regimens use a bosentan dose of 62.5 mg once daily or every other day. Discontinue bosentan at least 36 h prior to starting a ritonavir or cobicistat containing regimen and restart after at least 10 days at 62.5 mg once daily or every other day.
- Potential additive liver toxicity.
- Exposure of selexipag increased, but exposure of active metabolite unchanged.

RAL raltegravir