

Antiretroviral Treatment Options for Patients on Directly Acting Antivirals for Hepatitis C

	Simeprevir (Galexos®) 150 mg daily with food with sofosbuvir 400 mg daily	Daclatasvir (Daklinza®, DCV) 60 mg daily with sofosbuvir 400 mg daily	Ledipasvir/ Sofosbuvir (Harvoni®) 90/400 mg coformulation once daily	Velpatasvir/ Sofosbuvir (Epclusa®) 100/400 mg coformulation once daily	Holkira Pak®/ Viekira Pak® (US) (Abbvie 3D regimen: paritaprevir/ritonavir, ombitasvir 150/100/25 mg QD plus dasabuvir 250 mg BID)	Elbasvir/Grazoprevir (Zepatier®) 100 mg/50 mg coformulation once daily
PIs: atazanavir	Not recommended with ritonavir- or cobicistat boosted PIs (significant ↑ simeprevir AUC). ^{1,2}	↓ daclatasvir dose to 30 mg daily with atazanavir/ritonavir or atazanavir/ cobicistat. ^{3,4}	Potential for ↑ tenofovir concentrations when administered with concomitant booster. Monitor for toxicity. ⁵⁻⁷	OK with atazanavir/ritonavir. ⁸	OK with atazanavir 300 mg QD. ^{9,10}	Contraindicated with atazanavir ¹¹ : 10.58-fold ↑ grazoprevir AUC ¹² and 4.76-fold ↑ elbasvir exposures. ¹³
PIs: other		No dose modifications required with darunavir/ritonavir, darunavir/cobicistat or lopinavir/ritonavir. ⁴		OK with darunavir/ritonavir, lopinavir/ritonavir. ⁸	Darunavir: take without additional ritonavir; monitor HIV viral load due to decreased darunavir Ctough (Canadian monograph).	Contraindicated with darunavir, lopinavir, saquinavir, tipranavir ¹¹ : 7.5-12.86-fold ↑ grazoprevir AUC ¹² and 0.66-3.7-fold ↑ elbasvir exposures. ¹³
					US monograph: Not recommended due to potential for decreased darunavir Ctough. ¹⁰	
NNRTIs	Not recommended with efavirenz or nevirapine (71% ↓ simeprevir AUC). ^{1,2}	↑ daclatasvir dose to 90 mg once daily with efavirenz. ³	Efavirenz OK. ¹⁵	Do not use with efavirenz (50% ↓ velpatasvir AUC). ¹⁶	Contraindicated with efavirenz (increased risk of adverse events including LFT elevations). ^{10,17}	Contraindicated with efavirenz ¹¹ : (84% ↓ grazoprevir AUC ¹⁸ and 54% ↓ elbasvir AUC. ¹⁹
	Not recommended with etravirine. ²	No data. Coadministration not recommended with etravirine or nevirapine due to potential for ↓			Etravirine contraindicated due to risk of decreased 3D exposures. ¹⁰	Not recommended with etravirine ¹¹ due to potential for decreased elbasvir and grazoprevir concentrations.

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		daclatasvir. ²⁰				
	Rilpivirine OK. ²¹	Rilpivirine OK. ^{20, 22}	Rilpivirine OK. ¹⁵ Rilpivirine/FTC/tenofovir alafenamide: OK. ²³	Rilpivirine OK. ¹⁶	Not recommended with rilpivirine (116-273% ↑ rilpivirine exposures). ¹⁷	Rilpivirine OK. ²⁴
InSTIs		Dolutegravir OK. ^{20, 25}	Dolutegravir OK. Monitor for tenofovir-associated toxicities if using tenofovir-based backbone. ²⁶	Dolutegravir OK. ¹⁶	Dolutegravir OK. ²⁷	Dolutegravir OK. ²⁸
	Raltegravir OK. ²¹	Raltegravir OK. ²²	Raltegravir OK. ¹⁵	Raltegravir OK. ¹⁶	Raltegravir OK. ^{10, 17}	Raltegravir OK. ^{19, 29}
	Not recommended with cobicistat-boosted regimens. ¹	↓ daclatasvir dose to 30 mg daily with cobicistat ²⁰	Potential for ↑ tenofovir concentrations when administered with concomitant booster. Monitor for toxicity. ^{6, 7} NB: US monograph: combination not recommended. ⁵	Elvitegravir/cobicistat/FTC/tenofovir: 40% ↑ tenofovir AUC. Monitor for toxicity. ⁸		Not recommended with elvitegravir/cobicistat/tenofovir/emtricitabine due to potential for increased elbasvir and grazoprevir concentrations. ¹¹
			Elvitegravir/cobicistat/FTC/tenofovir alafenamide: OK. ²⁶	Elvitegravir/cobicistat/FTC/tenofovir alafenamide: OK. ⁸		
Maraviroc		Standard doses of both OK. ²⁰				
NRTIs	Tenofovir OK. ¹	Tenofovir OK. ³	Potential for ↑ tenofovir concentrations. Monitor for toxicity. ⁷	Potential for 40-81% ↑ tenofovir concentrations. Monitor for toxicity. ³⁰	Tenofovir OK. ^{9, 10}	Tenofovir OK. ^{19, 29}
			Tenofovir alafenamide OK. ²³	Tenofovir alafenamide OK. ³⁰		

Key:  = avoid combination  = caution/dose adjustment  = combination OK

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