

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

| | Simeprevir (Galexos®) 150 mg daily with food | Sofosbuvir (Sovaldi®) 400 mg daily | Sofosbuvir/Ledipasvir (Harvoni®) 400 mg/90 mg coformulation once daily | Daclatasvir (Daklinza®, DCV, BMS-790052) 60 mg daily (licensed in Japan & EU) | Holkira Pak®/Viekira Pak® (US) (Abbvie 3D regimen: paritaprevir/ritonavir, ombitasvir 150/100/25 mg QD plus dasabuvir 250 mg BID) | Grazoprevir (MK-5172) 100 mg/50 mg coformulation once daily (investigational) | Elbasvir (MK-8742) | | |
|-----------------|--|--|--|--|---|---|--|---|---|
| PIs: atazanavir | Not recommended with ritonavir- or cobicistat boosted PIs (significant ↑ simeprevir AUC). ^{1,2} | OK ³ | Potential for ↑ tenofovir concentrations. Monitor for toxicity. ^{4,5} | ↓ daclatasvir dose to 30 mg daily with atazanavir/ritonavir or atazanavir/cobicistat. ^{6,7} | OK with atazanavir 300 mg QD. ^{8,9} | Not recommended with atazanavir/ritonavir (10.58-fold ↑ MK-5172 AUC). ¹⁰ | Atazanavir concentrations OK, but significant ↑ MK-8742 exposures (4.76-fold). ¹¹ | | |
| PIs: other | | | | No dose modifications required with darunavir/ritonavir, darunavir/cobicistat or lopinavir/ritonavir. ⁷ | Darunavir: take without additional ritonavir; monitor HIV viral load due to decreased darunavir Ctrough (Canadian monograph). US monograph: Not recommended due to potential for decreased darunavir Ctrough. ⁹ | | | Not recommended with darunavir/ritonavir, lopinavir/ritonavir (7.5-12.86-fold ↑ MK-5172 AUC). ¹⁰ | Darunavir and lopinavir concentrations OK, but significant ↑ MK-8742 exposures (0.66-3.7-fold). ¹¹ |
| | | | | | Not recommended with lopinavir/ritonavir due to higher GI side effects and ↑ paritaprevir exposures. ¹² | | | | |
| NNRTIs | Not recommended with efavirenz or nevirapine (71% ↓ simeprevir AUC). ^{1,2} | Efavirenz OK ³ | Efavirenz OK. ¹³ | ↑ daclatasvir dose to 90 mg once daily with efavirenz. ⁶ | Contraindicated with efavirenz (increased risk of adverse events including LFT elevations). ^{9,14} | Avoid efavirenz (84% ↓ MK-5172 AUC). ¹⁵ | Avoid efavirenz (54% ↓ MK-8742 AUC). ¹⁶ | | |
| | Not recommended with etravirine. ² | | | | | | | No data. Coadministration not recommended with | No data. |

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|-----------|--|--|---|---|---|-------------------------------|-------------------------------|
| | | | | etravirine or nevirapine due to potential for ↓ daclatasvir. ¹⁷ | | | |
| | Rilpivirine OK. ¹⁸ | Rilpivirine OK ³ | Rilpivirine OK. ¹³ | Rilpivirine OK. ¹⁷ | Not recommended with rilpivirine (116-273% ↑ rilpivirine exposures). ¹⁴ | | |
| InSTIs | | | | Dolutegravir OK. ¹⁷ | | | |
| | Raltegravir OK. ¹⁸ | Raltegravir OK ³ | Raltegravir OK. ¹³ | Raltegravir OK. ¹⁴ | Raltegravir OK. ⁹ | Raltegravir OK. ¹⁹ | Raltegravir OK. ¹⁶ |
| | Not recommended with cobicistat-boosted regimens. ¹ | | Potential for ↑ tenofovir concentrations. Monitor for toxicity. ⁵ NB: US monograph: combination not recommended. ⁴ | ↓ daclatasvir dose to 30 mg daily with cobicistat ¹⁷ | | | |
| Maraviroc | | | | Standard doses of both OK. ¹⁷ | | | |
| NRTIs | Tenofovir OK. ¹ | Tenofovir OK ³ | Tenofovir OK. ¹³ | Tenofovir OK. ⁶ | Tenofovir OK. ^{8,9} | Tenofovir OK. ¹⁹ | Tenofovir OK. ¹⁶ |

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

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