CARDIOVASCULAR DRUGS:

STATINS AND LIPID LOWERING AGENTS

	INSTIs		NNRTIs		Pls	
	 BICTEGRAVIR (Biktarvy) DOLUTEGRAVIR (Tivicay, Triumeq, Juluca, Dovato) RALTEGRAVIR (Isentress) 	• ELVITEGRAVIR/ COBICISTAT (Stribild, Genvoya)	 DORAVIRINE (Pifeltro, Delstrigo) RILPIVIRINE (Edurant, Complera, Odefsey, Juluca) 	 EFAVIRENZ (Sustiva, Atripla) ETRAVIRINE (Intelence) NEVIRAPINE (Viramune) 	Boosted with ritonavir (Norvir) or cobicistat • ATAZANAVIR (Reyataz, Evotaz) • DARUNAVIR (Prezista, Prezcobix, Symtuza) • LOPINAVIR (Kaletra)	
STATINS						
• Atorvastatin (Lipitor)		Potential for ↑ statin		Potential for ↓ statin	Potential for ↑ statin. Use lowest statin dose possible (maximum 20 mg atorvastatin daily).	
• Rosuvastatin (Crestor)		Potential for ↑ statin			Potential for \uparrow statin. Use lowest statin dose possible (maximum 10 mg rosuvastatin daily).	
• Pitavastatin (Livalo)						
• Pravastatin (Pravachol)		Potential for ↑ statin			Potential for ↑ statin	
• Lovastatin (Mevacor), simvastatin (Zocor)		Potential for \uparrow statin and toxicity		Potential for ↓ statin	Potential for \uparrow statin and toxicity	

	INSTIs		NNRTIs		Pls
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FIBRATES					
 Fenofibrate, bezafibrate, gemfibrozil 					
CHOLESTEROL ABSO	ORPTION INHIBITOR				
• Ezetimibe (Ezetrol)					
GASTROINTESTINAL	LIPASE INHIBITOR				
• Orlistat (Xenical)	Potential for ↓ ARV absorption				
BILE ACID SEQUEST	RANTS				
• Cholestyramine (Olestyr), colestipol (Colestid)	Potential for ↓ ARV absorption				

Mechanism of Drug Interactions, Management and Monitoring

	Mechanism of Interaction	Main Interacting ARVs	Management	Monitoring
Hmg-Coa Reductase Inhibitors (Statins)				
Lovastatin, simvastatin	Inhibition of CYP3A4	Ritonavir and cobicistat- boosted protease inhibitors and elvitegravir	Contraindicated. Use alternate statin.	Statin toxicity: myalgia, rhabdomyolysis
Atorvastatin, rosuvastatin, pravastatin	Inhibition of CYP3A4, OATP1B1, BCRP	Ritonavir and cobicistat- boosted protease inhibitors and elvitegravir	Use lowest statin dose possible and titrate to effect	Statin toxicity: myalgia, rhabdomyolysis
Pitavastatin	Primarily cleared via UGT, OATP1B1	None	Most ARVs may be used	Statin toxicity: myalgia, rhabdomyolysis
Orlistat	Reduced absorption of ARVs by decreasing dietary fat absorption	All antiretrovirals, particularly lipophilic agents	Take ARVs at least 2 hours before/after orlistat	Antiretroviral efficacy
Bile acid sequestrants	Reduced absorption of ARVs	All antiretrovirals	Take ARVs either 1 hour before or 4-6 hours after bile acid sequestrants	Antiretroviral efficacy

Legend:

No dose adjustment required.

Use combination with caution. Adjustment in drug dose or frequency or additional/more frequent monitoring may be required.

May wish to consult with a pharmacist knowledgeable in HIV drug interactions.

Contraindicated/avoid combination.



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