

Table 238.2. Interactions of etravirine with other co-administered drugs.

Drug	Interaction
Nucleoside and nucleotide reverse transcriptase inhibitors	
Didanosine (ddl)	etravirine =; ddl =
Tenofovir (TDF)	etravirine ↓; TDF =
All other NRTIs	etravirine =; NRTI =
Protease inhibitors	
Atazanavir (ATV) (unboosted)	etravirine ↑↑; ATV ↓
Atazanavir (DRV) (cobicistat boosted)	etravirine ↓; cobicistat ↓; ATV ↓
Atazanavir (ritonavir boosted)	etravirine ↑; ATV ↓
Cobicistat (used to boost protease inhibitors plasma levels)	cobicistat ↓↓
Darunavir (DRV) (cobicistat boosted)	etravirine ↓; cobicistat ↓; DRV ↓
Darunavir (DRV) (ritonavir boosted)	etravirine ↓; DRV =
Indinavir (IDV) (unboosted)	etravirine ↑↑; IDV ↓↓
Lopinavir (LPV) (ritonavir boosted)	etravirine ↑; LPV ↓
Nelfinavir (NFV)	etravirine ↑; NFV ↑
Ritonavir (high dose as sole protease inhibitor)	etravirine ↓↓
Saquinavir (SQV) (unboosted)	etravirine =; SQV ↓↓
Saquinavir (SQV) (ritonavir boosted)	etravirine ↓; SQV =
Tipranavir (TPV) (ritonavir boosted)	etravirine ↓↓; TPV =
Nonnucleoside reverse transcriptase inhibitors	
Efavirenz (efavirenz)	etravirine ↓↓
Nevirapine (nevirapine)	etravirine ↓↓
Rilpivirine	etravirine =; rilpivirine ↓
Fusion inhibitor	
Enfuvirtide (ENF)	etravirine =; ENF =
CCR5 antagonist	
Maraviroc (without ritonavir-boosted protease inhibitor)	etravirine =; maraviroc ↓↓
Maraviroc (in combination with ritonavir-boosted protease inhibitor)	etravirine =; maraviroc ↑
Integrase inhibitors	
Dolutegravir	etravirine =; dolutegravir ↓
Eltegravir (unboosted)	etravirine =; eltegravir =
Raltegravir	etravirine =; raltegravir ↓
Nucleoside and nucleotide analogs for hepatitis B and C	
Tenofovir	etravirine =; nucleotide analog =
Adefovir, entecavir, lamivudine, ribavirin, telbivudine	

No significant interaction; etravirine AUC increased 11%
Changes clinically insignificant; etravirine AUC decreased 19%; TDF AUC increased 15%

No interactions expected (not studied)

Co-administration is not recommended owing to low ATV plasma level (down 20%); etravirine AUC increased 50%

Co-administration is not recommended because cobicistat plasma concentrations and, consequently, atazanavir concentrations are expected to decrease significantly

Changes not clinically significant; etravirine AUC increased 30%; ATV level down 14%

Co-administration is not recommended owing to low cobicistat plasma levels

Co-administration is not recommended because cobicistat plasma concentrations and consequently darunavir concentrations are expected to decrease significantly

Changes probably not clinically significant but etravirine AUC decreased 37%, DRV AUC up only 6%

Co-administration is not recommended because IDV AUC decreased 50%; and etravirine AUC increased 50%

Changes not clinically significant; etravirine AUC increased 17%; LPV AUC decreased 20%

Interaction expected (not studied)

Co-administration is not recommended owing to low etravirine plasma levels

Co-administration is not recommended because SQV AUC decreased 50%

Changes not clinically significant; SQV AUC decreased 13%

Co-administration is not recommended because etravirine AUC decreased 76%; TPV AUC increased 18%

Co-administration is not recommended owing to low etravirine plasma levels as well as other reasons

Co-administration is not recommended owing to low etravirine plasma levels as well as other reasons

Co-administration is not recommended owing to low rilpivirine plasma levels as well as other reasons

No interaction

Maraviroc should be given in a higher dose of 600 mg twice daily

Maraviroc should be given in a lower dose of 150 mg twice daily

C_{max} , AUC, C_{trough} decreased 52–88%

No significant interaction

Changes not clinically significant

See entries for nucleoside and nucleotide reverse transcriptase inhibitors

Interaction unexpected (not studied)