

# Bronchodilators (for COPD) Treatment Selector

Charts revised December 2023. Full information available at [www.hiv-druginteractions.org](http://www.hiv-druginteractions.org)

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	ATV/c	ATV/r	DRV/c	DRV/r	LPV/r	DOR	EFV	ETV	NVP	RPV oral	FTR	LEN	MVC	BIC/F/TAF	CAB oral	CAB/RPV	DTG	EVG/c/F/TAF	EVG/c/F/TDF	RAL	FTC/TAF	FTC/TDF
<b>Long acting muscarinic antagonists</b>																						
Acidinium bromide	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
Glycopyrronium bromide	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
Tiotropium bromide	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
Umeclidinium bromide	↑	↑	↑	↑	↑	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↑	↑	↔	↔	↔
<b>Short acting muscarinic antagonist</b>																						
Ipratropium bromide	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
<b>Long acting β2 agonists</b>																						
Formoterol	↔♥	↔♥	↔	↔	↔♥	↔	↔♥	↔	↔	↔♥	↔♥	↔	↔	↔	↔	↔♥	↔	↔	↔	↔	↔	↔
Indacaterol	↑a	↑a	↑a	↑a	↑a	↔	↓	↓	↓	↔	↔	↑	↔	↔	↔	↔	↔	↔	↑a	↑a	↔	↔
Olodaterol	↑	↑	↑	↑	↑	↔	↔	↔	↔	↔	↔	↑	↔	↔	↔	↔	↔	↔	↑	↑	↔	↔
Salmeterol	↑	↑	↑	↑	↑	↔	↓	↓	↓	↔	↔	↔	↔	↔	↔	↔	↔	↔	↑	↑	↔	↔
Vilanterol	↑	↑	↑	↑	↑	↔	↓	↓	↓	↔	↔	↑	↔	↔	↔	↔	↔	↔	↑	↑	↔	↔
<b>Short acting β2 agonists</b>																						
Salbutamol (albuterol)	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
Terbutaline	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
<b>Methylxanthines</b>																						
Aminophylline	↔	↓	↔	↓	↓	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
Theophylline	↔	↓	↔	↓	↓	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔
<b>Phosphodiesterase 4 inhibitors</b>																						
Roflumilast	↑	↑	↑	↑	↑	↔	↓	↓	↓	↔	↔	↔	↔	↔	↔	↔	↔	↔	↑	↑	↔	↔
<b>Inhaled corticosteroids</b>																						
Beclometasone	↑b	↑b	↔c	↓11% c	↑b	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↑b	↑b	↔	↔
Budesonide	↑d	↑d	↑d	↑d	↑d	↔	↓	↓	↓	↔	↔	↑d	↔	↔	↔	↔	↔	↔	↑d	↑d	↔	↔
Ciclesonide	↑e	↑e	↑e	↑e	↑e	↔	↔	↔	↔	↔	↔	↑e	↔	↔	↔	↔	↔	↔	↑e	↑e	↔	↔
Fluticasone	↑d	↑d	↑d	↑d	↑d	↔	↓	↓	↓	↔	↔	↑d	↔	↔	↔	↔	↔	↔	↑d	↑d	↔	↔
Mometasone	↑d	↑d	↑d	↑d	↑d	↔	↓	↓	↓	↔	↔	↑d	↔	↔	↔	↔	↔	↔	↑d	↑d	↔	↔

**Interactions with CAB/RPV long acting injections**  
Pharmacokinetic interactions shown are mostly with RPV. QT interactions shown are with RPV.

**Interactions with Lenacapavir**  
Residual LEN may affect exposure of sensitive CYP3A4 substrates initiated within 9 months after stopping subcutaneous LEN.

**Interactions with Ibalizumab**  
None

**Interactions with Abacavir (ABC), Lamivudine (3TC), Tenofovir-DF (TDF) or Zidovudine (ZDV)**  
ABC: No clinically relevant interactions expected.  
3TC: No clinically relevant interactions expected.  
TDF: No clinically relevant interactions expected.  
ZDV: No clinically relevant interactions expected.

**Colour Legend**

- No clinically significant interaction expected.
- These drugs should not be coadministered.
- Potential interaction which may require a dose 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 bronchodilator
  - ↓ Potential decreased exposure of the bronchodilator
  - ↔ No significant effect
  - ♥ One or both drugs may cause QT and/or PR prolongation. ECG monitoring is advised if coadministered with atazanavir or lopinavir. Efavirenz has a potential risk of QT prolongation relating specifically to homozygous carriers of CYP2B6\*6/\*6. Rilpivirine and fostemsavir were shown to prolong the QT interval at supratherapeutic doses. Caution is advised with rilpivirine. ECG monitoring is advised with fostemsavir and drugs with a known QT prolongation risk.
- Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

**Text Legend**

- ↑ Potential increased exposure of the bronchodilator
- ↓ Potential decreased exposure of the bronchodilator
- ↔ No significant effect

**Notes**

- a Exposure can be increased by up to 2-fold with ritonavir (and may be similar with cobicistat), however, this increase does not raise any concerns based on indacaterol's safety data.
- b Coadministration of ritonavir (100 mg twice daily) increased the AUC of the active metabolite (beclometasone-17-monopropionate) by 108% but no significant effect on adrenal function was seen. Caution is still warranted, use the lowest possible corticosteroid dose and monitor for corticosteroid side effects.
- c DRV/r decreased the AUC of active metabolite (beclometasone-17-monopropionate) by 11%, but no significant effect on adrenal function was seen.
- d Risk of elevated corticosteroid levels, Cushing's syndrome and adrenal suppression. This risk is present for oral and injected administration, and also for topical, inhaled or eye drop formulations.
- e No dose adjustment required but monitor closely, especially for signs of Cushing's syndrome when using a high dose or prolonged administration.