

# Bronchodilators (for COPD) Treatment Selector

Charts reviewed June 2022. 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	MVC	BIC/F/TAF	CAB oral	CAB/RPV	DTG	EVG/c/F/TAF	EVG/c/F/TDF	RAL	FTC/TAF	FTC/TDF	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	↔	↔	↔
Ciclesonide	↑e	↑e	↑e	↑e	↑e	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↔	↑e	↑e	↔	↔	↔
Fluticasone	↑d	↑d	↑d	↑d	↑d	↔	↓	↓	↓	↔	↔	↔	↔	↔	↔	↔	↔	↑d	↑d	↔	↔	↔
Mometasone	↑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 Abacavir (ABC), Lamivudine (3TC) or Zidovudine (ZDV)**  
ABC: No clinically relevant interactions expected.  
3TC: No clinically relevant interactions expected.  
ZDV: No clinically relevant interactions expected.

**Interactions with Ibalizumab**  
None

**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. 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.