

Bronchodilators (for COPD)

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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 |
|---|-------|-------|-------|--------|-------|-----|-----|-----|-----|----------|-----|-----|-----|-----------|----------|---------|-----|-------------|-------------|-----|---------|---------|
| Long acting muscarinic antagonists | | | | | | | | | | | | | | | | | | | | | | |
| Acidinium bromide | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Glycopyrronium bromide | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Tiotropium bromide | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Umeclidinium bromide | ↑ | ↑ | ↑ | ↑ | ↑ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↑ | ↔ | ↔ | ↔ |
| Short acting muscarinic antagonist | | | | | | | | | | | | | | | | | | | | | | |
| Ipratropium bromide | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Long acting β2 agonists | | | | | | | | | | | | | | | | | | | | | | |
| Formoterol | ↔♥ | ↔♥ | ↔ | ↔ | ↔♥ | ↔ | ↔♥ | ↔ | ↔ | ↔♥ | ↔♥ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔♥ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Indacaterol | ↑a | ↑a | ↑a | ↑a | ↑a | ↔ | ↓ | ↓ | ↓ | ↔ | ↔ | ↑ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑a | ↑a | ↔ | ↔ |
| Olodaterol | ↑ | ↑ | ↑ | ↑ | ↑ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↑ | ↔ | ↔ |
| Salmeterol | ↑ | ↑ | ↑ | ↑ | ↑ | ↔ | ↓ | ↓ | ↓ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↑ | ↔ | ↔ |
| Vilanterol | ↑ | ↑ | ↑ | ↑ | ↑ | ↔ | ↓ | ↓ | ↓ | ↔ | ↔ | ↑ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↑ | ↔ | ↔ |
| Short acting β2 agonists | | | | | | | | | | | | | | | | | | | | | | |
| Salbutamol (albuterol) | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Terbutaline | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Methylxanthines | | | | | | | | | | | | | | | | | | | | | | |
| Aminophylline | ↔ | ↓ | ↔ | ↓ | ↓ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Theophylline | ↔ | ↓ | ↔ | ↓ | ↓ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ |
| Phosphodiesterase 4 inhibitors | | | | | | | | | | | | | | | | | | | | | | |
| Roflumilast | ↑ | ↑ | ↑ | ↑ | ↑ | ↔ | ↓ | ↓ | ↓ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↔ | ↑ | ↑ | ↔ | ↔ |
| Inhaled corticosteroids | | | | | | | | | | | | | | | | | | | | | | |
| 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

- Green: No clinically significant interaction expected.
- Red: These drugs should not be coadministered.
- Orange: Potential interaction which may require a dose adjustment or close monitoring.
- Yellow: 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.
- ♥ Efavirenz has a potential risk of QT prolongation relating specifically to homozygous carriers of CYP2B6*6/*6. 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.