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Interactions with Experimental COVID-19 Therapies

Charts updated 9 April 2020

Please check www.covid19-druginteractions.org for updates.

Please note that if a drug is not listed it cannot automatically be assumed it is safe to coadminister. No recommendation to use experimental therapy for COVID-19 is made.

Drug interaction data for many agents are limited or absent; therefore, risk-benefit assessment for any individual patient rests with prescribers.

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Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



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Anaesthetics & Muscle Relaxants

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Alcuronium	\leftrightarrow	\leftrightarrow	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Bupivacaine	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	\leftrightarrow
Cisatracurium	\leftrightarrow								
Desflurane	\leftrightarrow								
Dexmedetomidine ♥	\leftrightarrow	↓ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Enflurane	\leftrightarrow								
Ephedrine	\leftrightarrow								
Etidocaine	1	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	↓	\leftrightarrow
Halothane	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	*	\leftrightarrow	\leftrightarrow	\leftrightarrow
Isoflurane	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ketamine	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	\leftrightarrow
Minaxolone	↑	↑	\leftrightarrow						
Nitrous oxide	\leftrightarrow	+	\leftrightarrow						
Propofol ♥	\leftrightarrow	↓ •	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Rocuronium	↑	↑	\leftrightarrow						
Sevoflurane ♥	\leftrightarrow	↔♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sufentanil	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	↓	\leftrightarrow
Suxamethonium (succinylcholine)	\leftrightarrow								
Tetracaine	\leftrightarrow								
Thiopental	\leftrightarrow								
Tizanidine ♥	\leftrightarrow	↓ •	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Vecuronium	\leftrightarrow								

Text Legend

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Analgesics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Alfentanil	1	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	\leftrightarrow
Aspirin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	+	\leftrightarrow	\leftrightarrow
Buprenorphine ♥	↑	↑~2% ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥		\	\leftrightarrow
Celecoxib	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow		\leftrightarrow	\leftrightarrow
Codeine	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow		\leftrightarrow		\leftrightarrow	\leftrightarrow
Dextropropoxyphene	↑	↑	\leftrightarrow	*		\leftrightarrow		↓	\leftrightarrow
Diamorphine (diacetylmorphine)	\leftrightarrow	\downarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow		\leftrightarrow	\leftrightarrow
Diclofenac	\leftrightarrow								
Dihydrocodeine	↑	^↓	\leftrightarrow			*		\leftrightarrow	\leftrightarrow
Fentanyl	↑	1	\leftrightarrow	+	\leftrightarrow	+	\leftrightarrow	\	\leftrightarrow
Hydrocodone ♥	↑↓	↑↓ v	\leftrightarrow	+	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Hydromorphone	\leftrightarrow	\	\leftrightarrow						
Ibuprofen	\leftrightarrow	\leftrightarrow	\leftrightarrow	+		\leftrightarrow		\leftrightarrow	\leftrightarrow
Mefenamic acid	\leftrightarrow	+	\leftrightarrow		+	+		\leftrightarrow	\leftrightarrow
Metamizole	↑↓	↑↓	₩		↓	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow
Methadone ♥	\leftrightarrow	↓ 53% ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Morphine	\leftrightarrow	+	\leftrightarrow						
Naproxen	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Nimesulide	\leftrightarrow								
Oxycodone	↑	1 160%	\leftrightarrow	\leftrightarrow		\leftrightarrow		\	\leftrightarrow
Paracetamol (Acetaminophen)	\leftrightarrow	\leftrightarrow	\leftrightarrow	1 14-16%		\leftrightarrow		\leftrightarrow	\leftrightarrow
Pethidine (Meperidine)	↑	\	\leftrightarrow						
Piroxicam	\leftrightarrow	\leftrightarrow	\leftrightarrow	+		\leftrightarrow		\leftrightarrow	\leftrightarrow
Remifentanil	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow		\leftrightarrow	\leftrightarrow
Tapentadol	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow		\leftrightarrow	\leftrightarrow
Tramadol ♥	1	↑ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow

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Notes:

Codeine and Tramadol + LPV/r

Potential decrease of the analgesic effect due to the reduced conversion to the active metabolite.

Diamorphine and Morphine + ATV

No effect on systemic exposure but inhibition of P-gp by atazanavir at the blood-brain barrier could potentiate the opiate effect in the CNS.

Diamorphine and Morphine + LPV/r

Ritonavir could reduce systemic exposure of diamorphine and morphine due to induction of glucuronidation. Ritonavir also inhibits P-gp at the blood-brain barrier and could potentiate the opiate effect in the CNS.

Hydrocodone + ATV or LPV/r

Hydrocodone concentrations are increased, but concentrations of the metabolite hydromorphone (which has also analgesic activity) are reduced.

Metamizole + CLQ, HCLQ, RBV, TCZ, IFN-B

Coadministration should be avoided due to the increased risk of haematological toxicity.

Paracetamol + FAVI

The daily dose of paracetamol in adults should be no more than 3000 mg/day (rather than 4000 mg/day).

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Antiarrhythmics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Amiodarone ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	+	\leftrightarrow
Bepridil ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	
Disopyramide ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Dofetilide♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Flecainide ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lidocaine (Lignocaine)	↑	↑	\leftrightarrow						
Mexiletine	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Propafenone	↑	↑	\leftrightarrow						
Quinidine ♥	↑	↑ ♥	\leftrightarrow	+	↔ ♥	↔♥	\leftrightarrow	\rightarrow	

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Note, please check product labels for any additional cardiac warnings.

Notes:

Amiodarone + LPV/r

The European product label for LPV/r contraindicates coadministration but the US product label suggests caution and concentration monitoring of amiodarone.

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Antibacterials

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Azithromycin ♥	1	↔ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Bedaquiline ♥	↑	122% ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ciprofloxacin ♥	\leftrightarrow	↔♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Clarithromycin ♥	↑ ↑	^ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Clindamycin	1	1	\leftrightarrow						
Clofazimine ♥	\leftrightarrow	↔ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Delamanid ♥	1	^ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Erythromycin ♥	1	^ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Isoniazid	\leftrightarrow								
Levofloxacin ♥	\leftrightarrow	↔ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Linezolid	\leftrightarrow								
Metronidazole	\leftrightarrow								
Moxifloxacin ♥	1	↓ •	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ofloxacin ♥	\leftrightarrow	↔ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Pyrazinamide	\leftrightarrow								
Rifabutin	1	1	\downarrow	\leftrightarrow	1	↓ ↓	\leftrightarrow	\leftrightarrow	\leftrightarrow
Rifampicin	\downarrow	↓ 75%	\downarrow	\leftrightarrow	\downarrow	\downarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Rifapentine	\downarrow	Ŭ.	\downarrow	\leftrightarrow	\downarrow	\downarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sulfadiazine	\leftrightarrow	\downarrow	\leftrightarrow						
Telithromycin ♥	↑ ↑	↑↑ 🕶	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Tinidazole	1	1	\leftrightarrow						

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Notes:

No interactions are expected with the COVID-19 therapies listed and the following antibacterials:

amikacin, amoxicillin, ampicillin, capreomycin, cefalexin, cefazolin, cefixime, cefotaxime, ceftazidime, ceftriaxone, chloramphenicol, clavulanic acid, cloxacillin, cycloserine, dapsone, doxycycline, ertapenem, ethambutol, ethionamide, flucloxacillin, gentamicin, imipenem/cilastatin, kanamycin, meropenem, nitrofurantoin, para-aminosalicylic acid, penicillins, piperacillin, rifaximin, spectinomycin, streptomycin, tazobactam, tetracyclines, trimethoprim/sulfamethoxazole, vancomycin.

Clarithromycin + ATV or LPV/r

A dose reduction of clarithromycin may be required for patients with impaired renal function. Refer to product labels for details.

Delamanid + ATV or LPV/r

Coadministration is expected to increase concentrations of DM-6705, a delamanid metabolite which is associated with QT prolongation. Frequent ECG monitoring is recommended.

Isoniazid + RBV

Use of isoniazid should be carefully monitored with patients with current chronic liver disease. Severe and sometimes fatal hepatitis associated with isoniazid therapy may occur and may develop even after many months of treatment.

Linezolid + RBV

Myelosuppression has been reported with both linezolid and ribavirin. Close monitoring of blood counts is recommended.

Linezolid + TCZ or IFN-β

Caution is required due to potential additive haematological toxicity.

Metronidazole and Tinidazole + LPV/r

No interaction is expected with lopinavir tablets. Coadministration is not recommended with lopinavir oral solution as it contains alcohol.

Pyrazinamide + FAVI

No effect on pyrazinamide concentrations but coadministration increased blood uric acid concentrations. Monitor uric acid.

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Anti-coagulant, Anti-platelet and Fibrinolytic

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Acenocoumarol	\leftrightarrow	\	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	\leftrightarrow
Apixaban	↑	1	\leftrightarrow	\leftrightarrow	↑	1	\leftrightarrow	\	\leftrightarrow
Argatroban	\leftrightarrow								
Aspirin (anti-platelet)	\leftrightarrow								
Betrixaban ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Clopidogrel	\	\downarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	→	\leftrightarrow
Dabigatran	↑	↔ or ↓	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow
Dalteparin	\leftrightarrow								
Dipyridamole	↑	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Edoxaban	↑	1	\leftrightarrow	\leftrightarrow	1	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow
Eltrombopag	\leftrightarrow	↓ 17%	\leftrightarrow						
Enoxaparin	\leftrightarrow								
Fondaparinux	\leftrightarrow								
Heparin	\leftrightarrow								
Phenprocoumon	↑	^↓	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	→	\leftrightarrow
Prasugrel	\leftrightarrow	→	\leftrightarrow						
Rivaroxaban	↑	↑	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	→	\leftrightarrow
Streptokinase	\leftrightarrow								
Ticagrelor	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	\leftrightarrow
Warfarin	↑	\	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	+	\leftrightarrow

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Notes:

Apixaban + LPV/r

The US product label for apixaban suggests to use apixaban at a reduced dose (2.5 mg twice daily) if needed.

Betrixaban + ATV or LPV/r

The US product label for betrixaban recommends for patients receiving or starting a strong P-gp inhibitor to reduce betrixaban dose and use an initial dose of 80 mg followed by 40 mg once daily.

Clopidogrel + ATV or LPV/r

Decreased conversion to active metabolite leading to non-responsiveness to clopidogrel. Prasugrel should be preferred to clopidogrel with ATV or LPV/r.

Edoxaban + ATV or LPV/r

The European product label for edoxaban states to consider a dose reduction of edoxaban from 60 mg to 30 mg with strong P-gp inhibitors, however, the US product label recommends no dose modification.

Prasugrel + ATV or LPV/r

Concentrations of active metabolite are reduced but without a significant reduction in prasugrel activity.

Vitamin K antagonists + ATV or LPV/r

Monitor INR with vitamin K antagonists (e.g., acenocoumarol, phenprocoumon, warfarin).

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Anticonvulsants

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Carbamazepine	↑↓	↑↓	↓	\leftrightarrow	₩	↓	\leftrightarrow	\downarrow	\leftrightarrow
Clonazepam	↑	↑	\leftrightarrow						
Eslicarbazepine	₩	↓	₩	\leftrightarrow	₩	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ethosuximide	↑	↑	\leftrightarrow						
Gabapentin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lacosamide		\leftrightarrow			\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lamotrigine	+	↓ 50%	\leftrightarrow						
Levetiracetam		\leftrightarrow			\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Oxcarbazepine	↓	↓	↓	\leftrightarrow	₩	₩	\leftrightarrow	\leftrightarrow	\leftrightarrow
Perampanel	↑	↑	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Phenobarbital (Phenobarbitone)	↓	↓	₩	\leftrightarrow	₩	₩	\leftrightarrow	↓	\leftrightarrow
Phenytoin	↓	↓	₩	\leftrightarrow	₩	₩	\leftrightarrow	\	
Pregabalin	\leftrightarrow	*	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	
Primidone	↓	$\downarrow \downarrow$	₩	\leftrightarrow	₩	↓	\leftrightarrow	\	
Retigabine	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Rufinamide	₩	↓	₩	\leftrightarrow	1	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sultiame	↑	↑	\leftrightarrow						
Tiagabine	↑	↑	\leftrightarrow						
Topiramate	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Valproate (Divalproex)	\leftrightarrow	1 38%	\leftrightarrow						
Vigabatrin	+	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Zonisamide	+	*	\leftrightarrow	+	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow

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Notes:

Valproate + LPV/r

Case report of a 48% decrease in valproate concentration in previously stable patient who developed exacerbated mania on starting lopinavir/ritonavir; dose increase of valproate was required.

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Antidepressants

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Agomelatine	\leftrightarrow	+	\leftrightarrow						
Amitriptyline	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow
Bupropion	\leftrightarrow	↓ 57%	\leftrightarrow						
Citalopram ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Clomipramine	↑	1	\leftrightarrow		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow
Desipramine ♥	\leftrightarrow	↑ 5% ♥	\leftrightarrow		↑ ♥	↑ ♥		\leftrightarrow	\leftrightarrow
Doxepin	\leftrightarrow	↑	\leftrightarrow		\leftrightarrow			\leftrightarrow	*
Duloxetine	\leftrightarrow	↑↓	\leftrightarrow		↑	↑		\leftrightarrow	*
Escitalopram ♥	1	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥		+	*
Fluoxetine	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	+	\leftrightarrow
Fluvoxamine	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	+	\leftrightarrow
Imipramine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lithium ♥	\leftrightarrow	↔♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Maprotiline ♥	\leftrightarrow	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Mianserin ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Milnacipran	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Mirtazapine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Nefazodone	↑ ↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Nortriptyline ♥	\leftrightarrow	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑v	\leftrightarrow	\leftrightarrow	\leftrightarrow
Paroxetine	↑↓ ?	↑↓ ?	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow
Phenelzine	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow
Reboxetine	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow
Sertraline	↑	\	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow
St John's wort	↓	↓	₩	\leftrightarrow	↓	↓	+	\leftrightarrow	\leftrightarrow
Tranylcypromine	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow
Trazodone	1	↑	\leftrightarrow						
Trimipramine ♥	\leftrightarrow	↑v	\leftrightarrow	\leftrightarrow	↑ ♥	↑₩	\leftrightarrow	\leftrightarrow	\leftrightarrow
Venlafaxine ♥	1	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑₩	\leftrightarrow	\leftrightarrow	\leftrightarrow
Vortioxetine	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	↑	1		\leftrightarrow	\leftrightarrow

Text Legend

- ↑ Potential increased exposure of the comedication
- Potential decreased exposure of the comedication
- 1 Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- → No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

▼ These drugs have been identified by www.crediblemeds.org as having a known or possible QT or TdP risk. The risk may be concentration- or dose-related and/or additive if two or more such drugs are combined.

Note, please check product labels for any additional cardiac warnings.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



Charts updated 9 April 2020

Please check www.covid19-druginteractions.org for updates.

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Please note that if a drug is not listed it cannot automatically be assumed it is safe to coadminister. No recommendation to use experimental therapy for COVID-19 is made.

Drug interaction data for many agents are limited or absent; therefore, risk-benefit assessment for any individual patient rests with prescribers.

Anti-diabetics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Acarbose	\leftrightarrow								
Canagliflozin	\leftrightarrow	\	\leftrightarrow						
Dapagliflozin	\leftrightarrow								
Dulaglutide	₩	\leftrightarrow							
Empagliflozin	\leftrightarrow								
Exanatide	↓	\leftrightarrow							
Glibenclamide (Glyburide)	↑	↑	\leftrightarrow						
Gliclazide	\leftrightarrow	↓	\leftrightarrow						
Glimepiride	\leftrightarrow	→	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	*
Glipizide	\leftrightarrow	→	\leftrightarrow						
Insulin	\leftrightarrow								
Linagliptin	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	*
Liraglutide	₩	\leftrightarrow							
Metformin	\leftrightarrow								
Nateglinide	↑	^↓	\leftrightarrow						
Pioglitazone	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Repaglinide	↑	↑	\leftrightarrow	1 52%	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	
Rosiglitazone	\leftrightarrow	\rightarrow	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Saxagliptin	↑	↑	\leftrightarrow						
Sitagliptin	↑	↑	\leftrightarrow						
Tolbutamide	\leftrightarrow	↓	\leftrightarrow						
Vildagliptin	\leftrightarrow								

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Notes:

Canagliflozin +LPV/r

If coadministration is deemed necessary, increasing canagliflozin to 300 mg once daily may be considered if patients are currently tolerating canagliflozin 100 mg once daily, have an eGFR ≥60 mL/min/1.73m² or CrCl ≥60 mL/min, and require additional glycaemic control. Other glucose-lowering therapies should be considered for patients with an eGFR 45 mL/min/1.73m² to <60 mL/min/1.73m² or CrCl 45 mL/min to <60 mL/min taking canagliflozin 100 mg who are receiving concurrent therapy with a UGT enzyme inducer and who require additional glycaemic control.

Linagliptin + LPV/r

The increase in linagliptin exposure is not considered clinically significant as it is mainly eliminated unchanged and has a large safety window.

Saxagliptin + ATV or LPV/r:

The US product label for saxagliptin states the recommended dose of saxagliptin to be 2.5 mg once daily when coadministered with strong CYP3A4/5 inhibitors.

Sitagliptin + ATV or LPV/r

The increase in sitagliptin exposure is not considered clinically significant as it is mainly eliminated unchanged and has a large safety window.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-ß	Interferon beta

· ·
These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

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Antifungals

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Amphotericin B	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Anidulafungin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Caspofungin	↑	\leftrightarrow							
Fluconazole ♥	+	↔♥	\leftrightarrow	\leftrightarrow	îv	ſÌ♥	\leftrightarrow	+	\leftrightarrow
Flucytosine	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Griseofulvin	↓	↓	\leftrightarrow	\leftrightarrow	₩	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow
Isavuconazole	↑	1 96%	\leftrightarrow	\leftrightarrow	Î	1	\leftrightarrow	+	\leftrightarrow
Itraconazole	↑	1	\leftrightarrow	\leftrightarrow	Π	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ketoconazole	↑	1	\leftrightarrow	\leftrightarrow	Π	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Micafungin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Miconazole	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Nystatin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Posaconazole	1 270%	1	\leftrightarrow	\leftrightarrow	Î	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Terbinafine	↑	1	\leftrightarrow						
Voriconazole	$\downarrow \downarrow$	↑↓ ↑	\leftrightarrow	\leftrightarrow	ſ	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- → No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

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Note, please check product labels for any additional cardiac warnings...

Notes:

Griseofulvin + LPV/r

LPV/r oral solution contains alcohol. Consumption of alcohol in association with griseofulvin can result in a 'disulfram-like' type reaction. No such interaction is expected with LPV/r tablets.

Itraconazole or Ketoconazole + ATV or LPV/r

The daily dose of itraconazole or ketoconazole should not exceed 200 mg.

Voriconazole + ATV

The effect of atazanavir on voriconazole exposure is dependent on CYP2C19 metaboliser status. In the majority of patients decreases in both voriconazole and atazanavir exposures may be expected, leading to loss of therapeutic effect and possible development of resistance. The European SmPC for atazanavir recommends a patient's CYP2C19 genotype should be performed if feasible. In patients without a functional CYP2C19 allele, increased voriconazole exposures are expected.

Voriconazole + LPV/r

Coadministration may result in bidirectional interactions leading to increased concentrations of lopinavir/ritonavir and an increase or decrease in voriconazole. Administration of voriconazole with ritonavir (100 mg twice daily) decreased voriconazole AUC by 39%.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

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Potential interaction which may require a dose adjustment or close monitoring.
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No clinically significant interaction expected



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Anti-hypertensives - ACE inhibitors

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Benazepril	↑	\leftrightarrow							
Captopril				\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Cilazapril				\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Enalapril				\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Fosinopril		↑		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lisinopril				\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Perindopril	\leftrightarrow	+	\leftrightarrow						
Quinapril	\leftrightarrow	+	\leftrightarrow						
Ramipril	\leftrightarrow								
Trandolapril	\leftrightarrow								

Anti-hypertensives – Angiotensin antagonists

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Candesartan	\leftrightarrow								
Eprosartan	\leftrightarrow	+	+	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Irbesartan	\leftrightarrow	→		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	
Losartan	\leftrightarrow	\rightarrow		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	
Olmesartan	\leftrightarrow								
Telmisartan	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	
Valsartan	↑	↑	\leftrightarrow						

Anti-hypertensives – Diuretics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Amiloride	\leftrightarrow								
Bendroflumethiazide				\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	
Chlortalidone	\leftrightarrow								
Furosemide	\leftrightarrow								
Hydrochlorothiazide	\leftrightarrow	+	\leftrightarrow						
Indapamide	↑		\leftrightarrow						
Metolazone	\leftrightarrow								
Torasemide	\leftrightarrow	\	\leftrightarrow						
Xipamide	\leftrightarrow								

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required
No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

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Anti-hypertensives - Other agents

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Aliskiren	↑	↑	\leftrightarrow						
Clonidine	\leftrightarrow								
Digoxin	↑	↑	\leftrightarrow	\leftrightarrow	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Dopamine	\leftrightarrow								
Doxazosin	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Eplerenone	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Hydralazine		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Isosorbide dinitrate	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ivabradine	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Labetalol	↑	→	\leftrightarrow						
Lacidipine ♥	↑	↑ ♥	\leftrightarrow		↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lercanidipine	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Methyldopa		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Moxonidine		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Prazosin	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ranolazine	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sacubitril	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sodium nitroprusside	\leftrightarrow								
Spironolactone	\leftrightarrow								
Terazosin	↑	↑	\leftrightarrow						

Text Legend

- ↑ Potential increased exposure of the comedication
- ↓ Potential decreased exposure of the comedication
- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- \leftrightarrow No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

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Note, please check product labels for any additional cardiac warnings.

Notes:

Doxazosin + ATV or LPV/r

For patients already taking doxazosin, monitor blood pressure and reduce doxazosin dose as needed if hypotension occurs on starting ATV or LPV/r.

Isosorbide nitrate + ATV or LPV/r
Decreased active metabolite.

Sacubitril + ATV or LPV/r Increased active metabolite.

Terazosin + ATV or LPV/r

For patients already taking terazosin, monitor blood pressure and reduce terazosin dose as needed if hypotension occurs on starting ATV or LPV/r.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required
No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

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Anti-hypertensives – Pulmonary hypertension

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Ambrisentan	↑	↑	\leftrightarrow						
Bosentan	↑#	↑	₩	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Epoprostenol	\leftrightarrow								
lloprost		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow
Macitentan	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow
Riociguat		↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow
Selexipag	\leftrightarrow								
Sildenafil	↑	↑	\leftrightarrow						
Tadalafil	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow		
Treprostinil		\leftrightarrow	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow

Text Legend

- ↑ Potential increased exposure of the comedication
- ↓ Potential decreased exposure of the comedication
- Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- → No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

These drugs have been identified by www.crediblemeds.org as having a known or possible QT or TdP risk. The risk may be concentration- or dose-related and/or additive if two or more such drugs are combined.
Note, please check product labels for any additional cardiac warnings.

Notes:

Ambrisentan + ATV or LPV/r

Start ambrisentan at 5 mg and closely monitor the patient for tolerability.

Bosentan + LPV/r

When coadministered patients should be closely observed for bosentan toxicity, especially during the first week of co-administration. For patients on bosentan, the US product label for LPV/r suggests to discontinue bosentan at least 36 hours prior to initiation of LPV/r and after at least 10 days of LPV/r, to resume bosentan at 62.5 mg once daily or every other day based upon individual tolerability.

Riociauat + ATV or LPV/r

The European product label for riociguat does not recommend its use in presence of strong inhibitors of CYPs, P-gp and BCRP; the US product label recommends to start riociguat at a dose of 0.5 mg three times daily and to monitor for signs and symptoms of hypotension.

Tadalafil + ATV

The US product label for ATV suggests for patients receiving atazanavir for at least one week, to start tadalafil at 20 mg once daily and increase to 40 mg once daily based on individual tolerability. For patients on tadalafil, avoid the use of tadalafil when starting atazanavir. Stop tadalafil at least 24 hours before starting atazanavir. At least one week after starting atazanavir, resume tadalafil at 20 mg once daily and increase to 40 mg once daily based on individual tolerability.

Tadalafil + LPV/r

The European product label for LPV/r does not recommend tadalafil for the treatment of pulmonary arterial hypertension, but the US product label suggests for patients on tadalafil, to avoid use of tadalafil during the initiation of LPV/r and to stop tadalafil at least 24 hours prior to starting LPV/r. After at least one week following the initiation of LPV/r, resume tadalafil at 20 mg once daily. Increase to 40 mg once daily based upon individual tolerability.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



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Antipsychotics/Neuroleptics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Amisulpride	\leftrightarrow								
Aripiprazole ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Asenapine	↑	↓	\leftrightarrow						
Chlorpromazine ♥	\leftrightarrow	↑ ♥		\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Clozapine ♥	↑	↑ ♥		\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Fluphenazine	\leftrightarrow	↑		\leftrightarrow	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow
Haloperidol ♥	↑	↑ ♥		\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
lloperidone ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levomepromazine ♥	\leftrightarrow	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Olanzapine	\leftrightarrow	\	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Paliperidone ♥	↑	↑	\leftrightarrow						
Perazine	↑	1	\leftrightarrow						
Periciazine	↑	1	\leftrightarrow						
Perphenazine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Pimozide ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Pipotiazine	\leftrightarrow	↑	\leftrightarrow						
Quetiapine	↑	↑	\leftrightarrow						
Risperidone	↑	1	\leftrightarrow	\leftrightarrow	1	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sulpiride ♥	\leftrightarrow	↔ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Thioridazine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Tiapride ♥	\leftrightarrow	↔ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ziprasidone	↑	↑	\leftrightarrow						
Zotepine ♥	1	↑♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Zuclopenthixol ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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- Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- → No significant effect

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Note, please check product labels for any additional cardiac warnings.

Notes:

Clozapine + RBV, CLQ or HCLQ

The risk of haematological toxicity may be potentially increased as clozapine, ribavirin, chloroquine and hydroxychloroquine can cause myelosuppression. Closely monitor haematological parameters.

Clozapine + TCZ or IFN-β

Caution is required due to potential additive haematological toxicity.

Quetiapine + ATV or LPV/r

Coadministration contraindicated in the European product label for quetiapine, however, US product label recommends quetiapine should be reduced to one sixth of the original dose if coadministered with a potent CYP3A4 inhibitor.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

	•
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	Potential interaction which may require a dose adjustment or close monitoring.
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Antivirals – Covid-19 therapies

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Atazanavir		×	\leftrightarrow	\leftrightarrow	↑	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow
Lopinavir/ritonavir ♥	×		\leftrightarrow	\leftrightarrow	↑ ♥	↑ v	\leftrightarrow	\leftrightarrow	\leftrightarrow
Remdesivir		\leftrightarrow			\leftrightarrow	\leftrightarrow	\leftrightarrow		*
Favipiravir	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	*
Chloroquine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow		×	\leftrightarrow	\leftrightarrow	\leftrightarrow
Hydroxychloroquine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	×		\leftrightarrow	\leftrightarrow	\leftrightarrow
Ribavirin	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow
Tocilizumab	\leftrightarrow		\leftrightarrow						
Interferon beta-1a		\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	

Text Legend

- ↑ Potential increased exposure of the comedication
- ↓ Potential decreased exposure of the comedication.
- Potential increased exposure of COVID drug
- Potential decreased exposure of COVID drug
- → No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

♥ These drugs have been identified by www.crediblemeds.org as having a known or possible QT or TdP risk. The risk may be concentration- or dose-related and/or additive if two or more such drugs are combined. Note, please check product labels for any additional cardiac warnings.

Notes: ATV + LPV/r

These drugs are not intended to be combined for the treatment of COVID-19.

Chloroquine and hydroxychloroquine should not be coadministered.

CLQ or HCLQ + LPV/r

LPV/r may increase concentrations of chloroquine or hydroxychloroquine, but to a moderate extent. Since LPV/r and chloroquine or hydroxychloroguine can cause QT prolongation, ECG monitoring is recommended when coadministering these agents.

CLQ or HCLQ + RBV, TCZ or IFN-β

Use with caution due to potential additive toxicity.

The risk of haematological toxicity may be potentially increased as ribavirin and tocilizumab can cause myelosuppression. Closely monitor haematological parameters.

Use with caution due to increased risk of haematological toxicity.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir		Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
1		IFN-β	Interferon beta

I	These drugs should not be coadministered
I	Potential interaction which may require a dose adjustment or close monitoring.
I	Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
ĺ	No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

Charts updated 9 April 2020

Please check www.covid19-druginteractions.org for updates.

Please note that if a drug is not listed it cannot automatically be assumed it is safe to coadminister. No recommendation to use experimental therapy for COVID-19 is made.

Drug interaction data for many agents are limited or absent; therefore, risk-benefit assessment for any individual patient rests with prescribers.

Antivirals - HCV DDAs

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Elbasvir/Grazoprevir	↑ *	↑ *	\leftrightarrow						
Glecaprevir/Pibrentasvir	^ *	↑ *	\leftrightarrow						
Ledipasvir/Sofosbuvir		\leftrightarrow							
Ombitasvir/Paritaprevir/r	1	↑ *	\leftrightarrow	\leftrightarrow	↑	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ombitasvir/Paritaprevir/r + Dasabuvir	↑	↑ *	\leftrightarrow	↑	↑	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sofosbuvir		\leftrightarrow							
Sofosbuvir/Velpatasvir		+	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Sofosbuvir/Velpatasvir/Voxilaprevir	↑ *	↑ *	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

- Potential increased exposure of the comedication
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- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- \leftrightarrow No significant effect

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Note, please check product labels for any additional cardiac warnings.

Notes:

* Increased risk of ALT elevations due to an expected or observed significant increase in DAA concentrations.

All DAAs + CLQ and HCLQ

The product labels for chloroquine and hydroxychloroquine recommend caution in impaired hepatic function.

Ledipasvir/sofosbuvir + LPV/r

Case reports of possible interaction between ledipasvir and LPV with patients having drug-induced liver injury manifesting as significant bilirubin rise within two weeks of starting ledipasvir/sofosbuvir while on LPV-containing HIV regimens.

Ombitasvir/Paritaprevir/r + ATV

Paritaprevir AUC increased by 187%. Exposures of paritaprevir greater than this have been evaluated in phase 2 studies and were not expected to have a clinically meaningful impact on safety.

Ombitasvir/Paritaprevir/r + Dasabuvir + ATV

Paritaprevir AUC increased by 94%. Exposures of paritaprevir greater than this have been evaluated in phase 2 studies and were not expected to have a clinically meaningful impact on safety. However, the combination carries an increased risk for hyperbilirubinaemia (including ocular icterus), particularly when ribavirin is also prescribed.

Ombitasvir/Paritaprevir/r + Dasabuvir + FAVI

Coadministration may increase dasabuvir concentrations. However, due to dasabuvir's large therapeutic index, a clinically relevant effect is not anticipated.

Ombitasvir/Paritaprevir/r ± Dasabuvir + CLQ

Inhibition of CYP3A4 by ritonavir may decrease exposure to chloroquine active metabolites, but this may not affect overall activity. No a priori dose alteration for chloroquine is recommended.

Ombitasvir/Paritaprevir/r ± Dasabuvir + HCLQ

Inhibition of CYP3A4 by ritonavir may decrease exposure to hydroxychloroquine, although to a moderate extent due to the multiple elimination pathways. No dose alteration is required.

Sofosbuvir/Velpatasvir + FAVI

Coadministration may increase velpatasvir concentrations. However, due to velpatasvir's large therapeutic index, a clinically relevant effect is not anticipated.

Sofosbuvir/Velpatasvir/Voxilaprevir + FAVI

Coadministration may increase velpatasvir concentrations. However, due to velpatasvir's large therapeutic index, a clinically relevant effect is not anticipated.

Key to abbreviations

	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-ß	Interferon beta

	•
	These drugs should not be coadministered
	Potential interaction which may require a dose adjustment or close monitoring.
	Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
	No clinically significant interaction expected



Charts updated 9 April 2020 Page 17 of 29

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Antivirals - Others

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Oseltamivir	+	\leftrightarrow	\leftrightarrow	1 4%	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

- ↑ Potential increased exposure of the comedication
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- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- \leftrightarrow No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

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Note, please check product labels for any additional cardiac warnings.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
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Interactions with Experimental COVID-19 Therapies

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Anxiolytics/Hypnotics/Sedatives

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Alprazolam	↑	1	\leftrightarrow						
Bromazepam	↑	1	\leftrightarrow						
Buspirone	↑	1	\leftrightarrow						
Chlordiazepoxide	↑	1	\leftrightarrow						
Clobazam	↑	1	\leftrightarrow						
Clorazepate	↑	1	\leftrightarrow						
Diazepam	↑	1	\leftrightarrow						
Estazolam	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Flunitrazepam	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Flurazepam	↑	1	\leftrightarrow						
Hydroxyzine	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	*
Lorazepam	\leftrightarrow								
Lormetazepam	\leftrightarrow								
Midazolam (oral)	↑	↑	\leftrightarrow						
Midazolam (parenteral)	↑	↑	\leftrightarrow						
Oxazepam	\leftrightarrow								
Temazepam	\leftrightarrow								
Triazolam	↑	1	\leftrightarrow						
Zaleplon	↑	1	\leftrightarrow						
Zolpidem	1	1	\leftrightarrow						
Zopiclone	↑	↑	\leftrightarrow						

Text Legend

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- ↑ Potential increased exposure of COVID drug
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Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
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		IFN-β	Interferon beta

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No clinically significant interaction expected



Charts updated 9 April 2020 Page 19 of 29

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Beta Blockers

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Atenolol	\leftrightarrow								
Bisoprolol	↑	↑	\leftrightarrow						
Carvedilol	↑	^↓	\leftrightarrow						
Metoprolol	\leftrightarrow	↑	\leftrightarrow						
Nebivolol	\leftrightarrow	↑	\leftrightarrow						
Oxprenolol	↑	\	\leftrightarrow						
Pindolol	\leftrightarrow	↑	\leftrightarrow						
Propranolol	\leftrightarrow	↑	\leftrightarrow						
Timolol	\leftrightarrow	↑	\leftrightarrow						

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

· ·
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Interactions with Experimental COVID-19 Therapies

Charts updated 9 April 2020

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Bronchodilators

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Aclidinium bromide	\leftrightarrow								
Aminophylline	\leftrightarrow	\	\leftrightarrow	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow	↓	↑
Formoterol	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Glycopyrronium bromide	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Indacaterol	1	↑	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ipratropium bromide	\leftrightarrow								
Montelukast	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Olodaterol	↑	↑	\leftrightarrow						
Roflumilast	↑	↑	\leftrightarrow						
Salbutamol	\leftrightarrow								
Salmeterol	↑	1	\leftrightarrow						
Theophylline	\leftrightarrow	\	\leftrightarrow	1 17-27%	\leftrightarrow	\leftrightarrow	\leftrightarrow	\	1
Tiotropium bromide	\leftrightarrow								
Umeclidinium bromide	↑	↑	\leftrightarrow	\leftrightarrow	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow
Vilanterol	↑	↑	\leftrightarrow						

Text Legend

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- ↓ Potential decreased exposure of the comedication
- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- ↔ No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

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Note, please check product labels for any additional cardiac warnings.

Notes:

Aminophylline + TCZ

Aminophylline is a complex of theophylline and ethylenediamine and is given for its theophylline activity. Coadministration may decrease theophylline concentrations.

Aminophylline or theophylline + IFN-β

Coadministration may increase theophylline concentrations but this is unlikely to be clinically significant. (Aminophylline is a complex of theophylline and ethylenediamine and is given for its theophylline activity.)

Indacaterol + ATV or LPV/r

Exposure can be increased by up to 2-fold with ritonavir (and may be similar with atazanavir), however, this increase does not raise any concerns based on indacaterol's safety data.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
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No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

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Calcium Channel Blockers

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Amlodipine	↑	↑	\leftrightarrow						
Diltiazem	1 25%	↑	\leftrightarrow						
Felodipine	↑	1	\leftrightarrow						
Nicardipine ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Nifedipine	↑	1	\leftrightarrow						
Nisoldipine	↑	1	\leftrightarrow						
Nitrendipine	↑	1	\leftrightarrow						
Verapamil	↑	↑	\leftrightarrow	\leftrightarrow	1	Î	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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- Potential decreased exposure of the comedication
- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- \leftrightarrow No significant effect

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Note, please check product labels for any additional cardiac warnings.

Notes:

Amlodipine + LPV/r

If coadministration is indicated, consider a dose reduction for amlodipine of 50%.

Diltiazem + ΔT\/

If coadministration is indicated, an initial dose reduction of diltiazem by 50% is recommended, with subsequent titration as needed and ECG monitoring.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

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Contraceptives

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Desogestrel (COC)	1	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Desogestrel (POP)	↑	1	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Drospirenone (COC)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ethinylestradiol	1 48%	↓ 42%	\leftrightarrow	1 43%	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Etonogestrel (implant)	↑	↑ 52%	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Etonogestrel (vaginal ring)	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Gestodene (COC)	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levonorgestrel (COC)	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levonorgestrel (emergency con.)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levonorgestrel (implant)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levonorgestrel (IUD)	\leftrightarrow								
Levonorgestrel (POP)	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	*
Medroxyprogesterone (depot inj)	\leftrightarrow	↑70%	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+
Norelgestromin (patch)	↑	↑83%	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+
Norethisterone (COC)	110%	↓ 17%	\leftrightarrow	1 47%	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norethisterone (IM depot)	\leftrightarrow	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norethisterone(POP)	↑	↑	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norgestimate (COC)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norgestrel (COC)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Ulipristal	↑	↑	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Notes:

COC - Combined oral contraceptive; POP - Progestogen only pill; IUD - Intra-uterine device

Contraceptives + RBV

Extreme care must be taken to avoid pregnancy in female patients and in female partners of male patients taking ribavirin. The European product labels for ribavirin state that effective contraception must be used during ribavirin treatment and for 4 months after treatment has been concluded in female patients and for 7 months in female partners of male patients. The US product labels for ribavirin state that effective contraception must be used during ribavirin treatment and for 6 months after treatment has been concluded in female patients and female partners of male patients.

Ethinylestradiol and/or progestins + ATV, LPV/r, FAVI

Concentrations of ethinylestradiol and progestins may be affected but no action is needed due to the short treatment duration of the COVID-19 therapy.

Levonorgestrel (emergency contraception) and Ulipristal + ATV or LPV/r

Any increase in exposure of levonorgestrel or ulipristal is unlikely to be clinically significant when used as a single dose.

Key to abbreviations

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FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

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ı	No clinically significant interaction expected



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Gastrointestinal Agents

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Alosetron	\leftrightarrow	\	\leftrightarrow						
Antacids	↓	\leftrightarrow	\leftrightarrow	\leftrightarrow	₩	₩	\leftrightarrow	\leftrightarrow	\leftrightarrow
Bisacodyl	\leftrightarrow								
Cimetidine	↓	\leftrightarrow							
Cisapride ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Esomeprazole	↓	\leftrightarrow							
Famotidine	↓ 41%	\leftrightarrow	*						
Lactulose	\leftrightarrow	*							
Lansoprazole	↓	+	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	
Loperamide	↑	1	\leftrightarrow						
Mesalazine	\leftrightarrow	+	\leftrightarrow						
Omeprazole	↓	\leftrightarrow							
Pantoprazole	↓	\leftrightarrow							
Prucalopride	\leftrightarrow	\leftrightarrow	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	
Rabeprazole	↓	\leftrightarrow							
Ranitidine	↓	\leftrightarrow							
Senna	\leftrightarrow								

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Notes:

Antacids + ATV

Antacids can reduce absorption of atazanavir. Atazanavir should be taken at least 2 h before or 1 h after antacids.

Antacids + CLQ

Antacids can reduce absorption of chloroquine. Antacids should be taken at least 2 h before or 2 h after chloroquine.

Antacids +HCLQ

Antacids can reduce absorption of hydroxychloroquine. Antacids should be taken at least 4 h before or 4 h after hydroxychloroquine.

Cimetidine, famotidine, ranitidine + ATV

Unboosted atazanavir is not recommended with H2RAs as they can reduce absorption of atazanavir. If coadministration is necessary, atazanavir 400 mg once daily with food should be administered at least 2 hours before and at least 10 hours after a dose of the H2RA.

Esomeprazole, lansoprazole, omeprazole, pantoprazole, rabeprazole + ATV

When possible, discontinue proton pump inhibitor treatment for the duration of atazanavir treatment.

Loperamide + ATV or LPV/r

Caution is advised with high doses of loperamide used for reducing stoma output, particularly as patients may be at increased risk of cardiac events due to electrolytes disturbances.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
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	•
	These drugs should not be coadministered
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Charts updated 9 April 2020 Page 24 of 29

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Gastrointestinal Agents – Anti-emetics

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Aprepitant	↑	↑	\leftrightarrow						
Dolasetron ♥	↑	↔♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Domperidone ♥	↑	^ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Dronabinol	↑	↑	\leftrightarrow						
Granisetron ♥	↑	↑ ♥	\leftrightarrow	\leftrightarrow	↔♥	↔♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Metoclopramide	\leftrightarrow								
Ondansetron ♥	↑	^ ♥	\leftrightarrow	\leftrightarrow	↔ ♥	↔ ♥	\leftrightarrow	\leftrightarrow	\leftrightarrow
Prochlorperazine	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	↑	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

- ↑ Potential increased exposure of the comedication
- Potential decreased exposure of the comedication
- ↑ Potential increased exposure of COVID drug
- ↓ Potential decreased exposure of COVID drug
- \leftrightarrow No significant effect

Numbers refer to increase or decrease in AUC as observed in drug-drug interaction studies.

▼ These drugs have been identified by www.crediblemeds.org as having a known or possible QT or TdP risk. The risk may be concentration- or dose-related and/or additive if two or more such drugs are combined.

Note, please check product labels for any additional cardiac warnings.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

· ·
These drugs should not be coadministered
Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



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Interactions with Experimental COVID-19 Therapies

Charts updated 9 April 2020

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Hormone Replacement Therapy

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Drospirenone (HRT)	↑	↑	\leftrightarrow						
Dydrogesterone (HRT)	↑	↑	\leftrightarrow	↑	\leftrightarrow	+	\leftrightarrow	\leftrightarrow	\leftrightarrow
Estradiol	↑	+	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Levonorgestrel (HRT)	↑	↑	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Medroxyprogesterone (oral)	↑	↑	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norethisterone (HRT)	↑	↑	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Norgestrel (HRT)	↑	↑	\leftrightarrow	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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- ↓ Potential decreased exposure of COVID drug
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Note, please check product labels for any additional cardiac warnings.

Notes:

Estradiol and + ATV, LPV/r or FAVI

Concentrations of estradiol may alter but no action is needed due to the short treatment duration of the COVID-19 therapy.

Progestins + ATV, LPV/r or FAVI

Concentrations of progestins may increase but no action is needed due to the short treatment duration of the COVID-19 therapy.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

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No clinically significant interaction expected



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Immunosuppressants

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Adalimumab	\leftrightarrow								
Anti-thymocyte globulin	\leftrightarrow		+	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Azathioprine	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	↑	\leftrightarrow	\leftrightarrow
Basiliximab	\leftrightarrow								
Belatacept	\leftrightarrow								
Ciclosporin	1	↑	\leftrightarrow	\leftrightarrow	↑	1	\leftrightarrow	↓	\leftrightarrow
Mycophenolate	\leftrightarrow	^↓	\leftrightarrow						
Pirfenidone	\leftrightarrow	\	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	1
Sirolimus	↑	↑	\leftrightarrow	\leftrightarrow	↑	1	\leftrightarrow	\	\leftrightarrow
Tacrolimus ♥	1	↑♥	\leftrightarrow	\leftrightarrow	↑ ♥	↑v	\leftrightarrow	\	\leftrightarrow

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Note, please check product labels for any additional cardiac warnings.

Notes:

Adalimumab and azathioprine + CLQ or HCLQ

The risk of haematological toxicity may be potentially increased as adalimumab, azathioprine, chloroquine and hydroxychloroquine can cause myelosuppression. Closely monitor haematological parameters.

Adalimumab + RBV

The risk of haematological toxicity may be potentially increased as adalimumab and ribavirin can cause myelosuppression. Closely monitor haematological parameters.

Adalimumab and basiliximab + TCZ

Avoid coadministration due to the enhanced immunosuppressive effect.

Adalimumab + IFN-β

Caution is required due to potential additive haematological toxicity.

Azathioprine + RBV

Ribavirin may interfere with azathioprine metabolism possibly leading to an accumulation of 6-methylthioinosine monophosphate, which has been associated with myelotoxicity.

Azathioprine + TCZ or IFN-β

Caution is required due to potential additive haematological toxicity.

Pirfenidone + IFN-β

Any increase is pirfenidone is unlikely to be clinically relevant, except in the presence of hepatic impairment as moderate hepatic impairment also increases pirfenidone exposure (by 60%). No a priori dosage adjustment is recommended in patients with hepatic impairment but monitor for increased toxicity.

Key to abbreviations

ATV	Atazanavir	CLQ	Chloroquine
LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir		Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
1		IFN-β	Interferon beta

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Potential interaction which may require a dose adjustment or close monitoring.
Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected



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Inotropes & Vasopressors

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Adrenaline (Epinephrine)	+	\leftrightarrow							
Dobutamine		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+
Noradrenaline	+	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Vasopressin		\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow

Text Legend

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Note, please check product labels for any additional cardiac warnings.

Notes:

Remdesivir

Pressor requirement to maintain blood pressure is a key exclusion criteria to eligibility for remdesivir use.

See https://rdvcu.gilead.com/ for further details.

Key to abbreviations

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LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir	RBV	Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
		IFN-β	Interferon beta

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Potential interaction which may require a dose adjustment or close monitoring.
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No clinically significant interaction expected



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Lipid Lowering Agents

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Atorvastatin	1	1 490%	\leftrightarrow						
Bezafibrate	\leftrightarrow	+							
Clofibrate	\leftrightarrow								
Evolocumab	\leftrightarrow	*							
Ezetimibe	↑	\leftrightarrow							
Fenofibrate	\leftrightarrow								
Fish oils	\leftrightarrow								
Fluvastatin	↑	\leftrightarrow							
Gemfibrozil	\leftrightarrow	↓ 41%	\leftrightarrow						
Lovastatin	↑	1	\leftrightarrow						
Pitavastatin	1 31%	↓ 20%	\leftrightarrow						
Pravastatin	↑	↑33%	\leftrightarrow						
Rosuvastatin	↑	1 08%	\leftrightarrow						
Simvastatin	↑	1	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	+

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Note, please check product labels for any additional cardiac warnings.

Notes:

Atorvastatin + ATV

Coadministration is not recommended. If the use of atorvastatin is considered necessary, use the lowest possible dose of atorvastatin with careful safety monitoring. The daily atorvastatin dose should not exceed 10 mg.

Atorvastatin + LPV/r

Do not exceed a daily dose of 20 mg with careful safety monitoring.

Evolocumab + TCZ

Avoid coadministration due to the enhanced immunosuppressive effect.

Rosuvastatin + ATV or LPV/r

Do not exceed rosuvastatin 10 mg/day.

Key to abbreviations

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LPV/r	Lopinavir/ritonavir	HCLQ	Hydroxychloroquine
RDV	Remdesivir		Ribavirin
FAVI	Favipiravir	TCZ	Tocilizumab
1		IFN-β	Interferon beta

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Steroids

	ATV	LPV/r ♥	RDV	FAVI	CLQ ♥	HCLQ ♥	RBV	TCZ	IFN-β
Beclometasone	\leftrightarrow	1	\leftrightarrow						
Betamethasone	^* ↓	^* ↓	₩	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow	\leftrightarrow
Budesonide	↑ *	↑ *	\leftrightarrow						
Ciclesonide	1	1	\leftrightarrow						
Clobetasol	↑ *	↑ *	\leftrightarrow						
Dexamethasone	^* ↓	1 * ↓	₩	\leftrightarrow	\leftrightarrow		\leftrightarrow	\leftrightarrow	\leftrightarrow
Fludrocortisone	↑ *	↑ *	\leftrightarrow						
Flunisolide	1	1	\leftrightarrow						
Fluocinolone	↑ *	↑ *	\leftrightarrow						
Fluticasone	^ *	↑ *	\leftrightarrow						
Hydrocortisone (oral)	↑ *	↑ *	\leftrightarrow						
Hydrocortisone (topical)	\leftrightarrow								
Megestrol acetate	\leftrightarrow								
Methylprednisolone	↑ *	↑ *	\leftrightarrow						
Mometasone	↑ *	↑ *	\leftrightarrow						
Nandrolone	\leftrightarrow								
Oxandrolone	\leftrightarrow								
Prednisolone	↑ *	↑ *	\leftrightarrow						
Prednisone	↑ *	↑ *	\leftrightarrow						
Stanazolol	↑	↑	\leftrightarrow						
Testosterone	↑	1	\leftrightarrow						
Triamcinolone	↑ *	↑ *	\leftrightarrow						

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Note, please check product labels for any additional cardiac warnings.

Notes:

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 drops corticosteroids

Beclometasone + LPV/r

Ritonavir (100 mg twice daily) increased the AUC of the active metabolite 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.

Betamethasone or Dexamethasone + ATV, LPV/r or RDV

Betamethasone and dexamethasone are moderate inducers of CYP3A4 and could decrease exposure and efficacy of ATV, LPV/r or RDV particularly when administered orally or intravenously at high doses or for a long duration.

Ciclesonide + ATV or LPV/r

No dose adjustment required but monitor closely, especially for Cushing's syndrome, when using a high dose or prolonged administration.

Flunisolide + ATV or LPV/r

Use the lowest possible flunisolide dose with monitoring for corticosteroid side effects.

Prednisolone or Prednisone + LPV/r

Based on DDI study with LPV/r, exposure of prednisolone (obtained also after conversion from prednisone) is increased modestly (+30%). A 30% dose reduction of the corticosteroid might be considered during concomitant treatment.

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Potential interaction likely to be of weak intensity. Additional action/monitoring or dosage adjustment unlikely to be required.
No clinically significant interaction expected