

## Lopinavir ritonavir+interferon alfa vs Ribavirin +Interferon alfa for COVID-19

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**Question:** Should Lopinavir/ritonavir+interferon alfa compared to Ribavirin +Interferon alfa for COVID-19

**Setting:** inpatient

Certainty assessment							№ of patients		Effect		Certainty
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Lopinavir/ritonavir+interferon alfa	Ribavirin +Interferon alfa	Relative (95% CI)	Absolute (95% CI)	
<b>All-cause mortality</b>											
1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	No death reported				⊕⊕○○ LOW
<b>SARS-CoV-2 clearance</b>											
1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	22/36 (61.1%)	17/33 (51.5%)	<b>RR 1.19</b> (0.78 to 1.81)	<b>98 more per 1.000</b> (from 113 fewer to 417 more)	⊕⊕○○ LOW
<b>Progression of COVID-19 disease</b>											
1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	2/36 (5.6%)	1/33 (3.0%)	<b>RR 1.83</b> (0.17 to 19.29)	<b>25 more per 1.000</b> (from 25 fewer to 554 more)	⊕⊕○○ LOW
<b>Number of patients with any adverse event</b>											
1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	26/36 (72.2%)	23/33 (69.7%)	<b>RR 1.04</b> (0.77 to 1.40)	<b>28 more per 1.000</b> (from 160 fewer to 279 more)	⊕⊕○○ LOW
<b>Number of patients with serious adverse events</b>											
1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	No serious adverse event reported				⊕⊕○○ LOW

Certainty assessment							№ of patients		Effect		Certainty
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Lopinavir/ritonavir+interferon alfa	Ribavirin +Interferon alfa	Relative (95% CI)	Absolute (95% CI)	

#### Time to SARS-CoV 2 clearance

1 <sup>1</sup>	randomised trials	serious <sup>a</sup>	not serious	not serious	serious <sup>b</sup>	none	The study reports that the negative conversion time (SARS-CoV-2 clearance) from baseline to follow-up did not differ between the two HR groups: 1.50 (95% CI 0.89,2.53)			
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#### Explanations

a. Downgraded of one level for high risk of performance bias

b. Downgraded of one level for small sample size

#### References

1. Huang YQ, Tang SQ, Xu XL, et al. No Statistically Apparent Difference in Antiviral Effectiveness Observed Among Ribavirin Plus Interferon-Alpha, Lopinavir/Ritonavir Plus Interferon-Alpha, and Ribavirin Plus Lopinavir/Ritonavir Plus Interferon-Alpha in Patients With Mild to Moderate Coronavirus Disease 2019: Results of a Randomized, Open-Labeled Prospective Study. *Front Pharmacol.* 2020;11:1071. Published 2020 Jul 14. doi:10.3389/fphar.2020.01071