

Approval Form for Use of Casirivimab + Imdevimab for COVID-19 Pneumonia

Osler COVID Clinical Response Committee (CCRC) Recommendations:

- Given the short supply, Osler supports the use of casirivimab + imdevimab in patients with **moderate to critical illness** as outlined by the Ontario Science Advisory Table. For complete criteria details, please visit https://covid19-sciencetable.ca/
- In the moderate to critical illness setting a one time dose of 2,400 mg of casirivimab + imdevimab (to be given intravenously) is recommended
- Casirivimab/Imdevimab will be dispensed upon receipt of this completed approval form AND an accompanying order
- · Prescribers must dictate a note describing the informed consent discussion and that the patient meets prescribing criteria

CCRC recommends an informed discussion with patients who meet the following criteria for casirivimab + imdevimab therapy for COVID-19:

Inclusion Criteria (Note: All boxes must be checked off)	YES	NO
Moderately ¹ or critically ² ill		
Confirmed (current) COVID-19 infection		
Symptoms for 9 days or less (Date of Symptom onset)		
At risk of acute decompensation		

²Critically ill patients: Patients requiring ventilator and/or circulatory support, including high-flow nasal oxygen, non-invasive ventilation or extracorporeal membrane oxygenation (ECMO)

Exclusion Criteria (Note: All boxes must be checked off)	YES	NO
History of previous COVID-19 infection		
Fully vaccinated against COVID-19		
Severe hypersensitivity (e.g. anaphylaxis) to casirivimab, imdevimab or any component of the formulation		

COMMENTS:								
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REQUIRED APPROVALS:

	NAME	SIGNATURE	DATE
Ordering Prescriber*			

This form is to be completed in advance. Maintain a copy of the form along with the order in the chart. Scan the form and the order to Pharmacy.









¹Moderately ill patients: Patients newly requiring low-flow supplemental oxygen