

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Casirivimab and imdevimab

INITIATION – Treatment of profoundly immunocompromised patients

Re-assessment required after 2 weeks

Prerequisites (tick boxes where appropriate)

- Patient has confirmed (or probable) COVID-19
- and The patient is in the community (treated as an outpatient) with mild to moderate disease severity*
- and Patient is profoundly immunocompromised** and is at risk of not having mounted an adequate response to vaccination against COVID-19 or is unvaccinated
- and Patient's symptoms started within the last 10 days
- and Patient is not receiving high flow oxygen or assisted/mechanical ventilation
- and Casirivimab and imdevimab is to be administered at a maximum dose of no greater than 2,400 mg

Note: * Mild to moderate disease severity as described on the [Ministry of Health Website](#)
** Examples include B-cell depletive illnesses or patients receiving treatment that is B-Cell depleting.

INITIATION – mild to moderate COVID-19-hospitalised patients

Re-assessment required after 2 weeks

Prerequisites (tick boxes where appropriate)

- Prescribed by, or recommended by any relevant practitioner, or in accordance with a protocol or guideline that has been endorsed by the Te Whatu Ora Hospital.
- and Patient has confirmed (or probable) COVID-19
- and Patient is an in-patient in hospital with mild to moderate disease severity*
- and Patient's symptoms started within the last 10 days
- and Patient is not receiving high flow oxygen or assisted/mechanical ventilation
- and Age > 50
- or BMI > 30
- or Patient is Māori or Pacific ethnicity
- or Patient is at increased risk of severe illness from COVID-19, excluding pregnancy, as described on the Ministry of Health website (see Notes)
- and Patient is unvaccinated
- or Patient is seronegative where serology testing is readily available or strongly suspected to be seronegative where serology testing is not available
- and Casirivimab and imdevimab is to be administered at a maximum dose of no greater than 2,400 mg

Note: * Mild to moderate disease severity as described on the [Ministry of Health Website](#)
** (<https://www.health.govt.nz/our-work/diseases-and-conditions/covid-19-novel-coronavirus/covid-19-information-specific-audiences/covid-19-advice-higher-risk-people>)

I confirm that the above details are correct:

Signed: Date: