

**Patient**

Name (Last, First): \_\_\_\_\_

DOB: \_\_\_\_\_ Age: \_\_\_\_\_ Primary Language: \_\_\_\_\_

**Provider**

Name (Last, First): \_\_\_\_\_

Phone: \_\_\_\_\_ Referral Help Line: (505) 291-2626

**Bamlanivimab and Casirivimab/Imdevimab Eligibility**

Bamlanivimab and Casirivimab/Imdevimab are unapproved monoclonal antibody therapies under FDA Emergency Use Authorization for the treatment of mild to moderate COVID-19 in adults and pediatric patients with positive results of direct SARS-CoV-2 viral testing who are 12 years of age and older; weighing at least 40 kg; and, who are at **high risk** for progressing to severe COVID-19 and/or hospitalization.

Monoclonal antibodies should be administered as soon as possible after positive SARS-CoV-2 (PCR or antigen testing) and **within 10 days of symptom onset**. Patients are not eligible for treatment if > 10 days from symptom onset. Not all Infusion Centers are able to facilitate pediatric infusions; please call your local site to confirm.

Per FDA requirements, patients will be provided an FDA *Fact Sheet for Patients, Parents and Caregivers* prior to infusion therapy.

**Patient Eligibility:** Patient **must meet at least one High Risk criterion** for treatment eligibility. Check all that apply.

- Body mass index (BMI)  $\geq$  35
- Chronic kidney disease
- Diabetes
- Immunosuppressive disease
- Currently receiving immunosuppressive treatment
- $\geq$  65 years of age
- $\geq$  55 years of age **AND** have at least one of the following. Check all that apply.
  - Cardiovascular disease
  - Hypertension
  - Chronic obstructive pulmonary disease/other chronic respiratory disease
- 12-17 years of age **AND** have at least one of the following. Check all that apply.
  - BMI  $\geq$  85th percentile for age and gender based on [CDC growth charts](#)
  - Sickle cell disease
  - Congenital or acquired heart disease
  - Neurodevelopmental disorders, for example cerebral palsy
  - A medical-related technological dependence, for example, tracheostomy, gastrostomy, or positive pressure ventilation (not related to COVID-19)
  - Asthma, reactive airway or other chronic respiratory disease that requires daily medication for control

**Required Patient Referral Information\***

Date of Symptom(s) Onset:		1 <sup>st</sup> COVID-19 Positive Test Result Date in past 90 days:	
Covid-19 Testing Location:		City:	State:

\*Include in the fax patient demographic information from the patient health record that includes a copy of their complete insurance data.

**Referring Provider Information**

Provider Name:		Provider Phone Number:	
Primary Practice Address:		City:	State:

**Therapy Location**

Please schedule patient for treatment at one of the following Presbyterian Infusion Centers:

Albuquerque      Santa Fe      Ruidoso      Clovis      Socorro      Tucumcari      Espanola

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**Provider Attestation**

- I attest the patient I am referring for Bamlanivimab (700 mg IV) or Casirivimab/Imdevimab (1200 mg/1200 mg IV) therapy:
- has a recent positive SARS-CoV-2 test result,
  - is under my medical care and will follow up with me for any necessary COVID-19- related medical issues,
  - has become symptomatic in the past 10 days (and can reasonably be scheduled for treatment within 10 days of symptom onset),
  - is  $\geq 12$  years of age,
  - weighs  $\geq 40$  kg,
  - has been informed (or parents/caregivers have been informed) of alternatives to receiving this medication,
  - has been informed (or parents/caregivers have been informed) that the medication is an unapproved product permitted for use under an Emergency Use Authorization only,
  - has been advised (or parents/caregivers have been advised) to report any suspected adverse drug events to this Infusion Center immediately for FDA reporting,
  - may receive either Bamlanivimab or Casirivimab/Imdevimab, depending on drug availability, as the indications and restrictions are similar for both therapies, and
  - does NOT meet ANY of the following exclusionary criteria:
    - o previously hospitalized due to COVID-19, OR
    - o requires oxygen therapy due to COVID-19, OR
    - o requires an increase in baseline oxygen flow rate due to COVID-19 in those on chronic oxygen therapy due to underlying non-COVID-19 related co-morbidity

**Provider Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

(Signature stamps not accepted, wet signature required)

**Referral Submission Instructions**Please fax all the of the following, in one fax, to the **Presbyterian Infusion Center Fax (505) 355-7158**.

- Fax coversheet from referring provider primary practice
- This Presbyterian Infusion Center Referral Form, fully complete, including wet signature of provider
- Patient Demographics (Patient Name, DOB, Address, Insurance, Primary Language)
- Copy of most recent clinical encounter (e.g. History and Physical, Progress Note, Telephone Note, Video Visit)
- Copy of positive COVID-19 test result.

Test result may be sent from provider's office OR patient may present hard copy, text, or email of positive test result for verification at time of appointment.

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*PLEASE NOTE: treatment will not be provided unless positive COVID-19 test result is confirmed.*

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**Referral Provider Comments**