

 Patient Name:
 DOB:

 Address:
 \_\_\_\_\_\_Phone Number\_\_\_\_\_

PLEASE ATTACH: 1. Patient Demographic Info <u>AND</u> 2. Copy of Positive Covid-19 Result

## Bamlanivimab Infusion Order

Diagnosis : Covid-19 Positive (ICD-10 Code) U07.1		Date symptom onset
Diagnosis :	(ICD-10 Code)	Date positive results

Hold infusion and notify provider for:

- Unable to sign consent
- Patient with first positive result for SARS-CoV-2 virus and onset of symptoms NOT within 10 day
- Weight less than 40 kg
- Age less than 18 years
- Requires oxygen therapy or Sat <93%, or HR >130, or RR >30
- Has already received prior dose of Bamlanivimab.

If infusion-related reaction occurs, stop infusion and follow Hypersensitivity Reaction Management Policy/Protocol as clinically indicated.

Administer: Bamlanivimab 700mg in 250 mL 0.9% sodium chloride over 60 minutes. Then flush line with 20 mL 0.9% NaCl. Use in-line or add-on 0.2/0.22 micron polyethersulfone (PES) filter and polyvinyl chloride (PVC) or polyethylene (PE)-lined PVC

Ondansetron 4mg PO (oral disintegrating tab) PRN nausea; may repeat x1 after 30 minutes

(Check all that apply)-must have one

\_\_\_\_Greater than or equal to 65 years of age

Body Mass Index (BMI) greater than or equal to 35

\_\_\_\_Chronic Kidney Disease (Stage IV or greater)

\_\_\_\_Diabetes with A1C ≥ 8 or random blood sugar > 300 mg/dL

Immunosuppressive Condition (solid organ transplant, ESRD or ESLD, advanced HIV, active chemotherapy, chronic high dose steroids (>30mg prednisone for >30 days), use of biologic agents for treatment of underlying diseases (i.e. TNF alpha inhibitor for RA or Crohn's)

## OR

Are greater than or equal to 55 years of age with one or more of the following:

Cardiovascular disease other than hypertension

\_\_\_\_Currently receiving treatment with medication for hypertension

\_\_\_\_Chronic Obstructive Pulmonary Disease, Interstitial Lung Disease, Cystic Fibrosis, or Pulmonary Fibrosis

Monitoring: Vital Signs Baseline, 5 minutes after infusion started then every 30 minutes

**Observation period:** Monitor patient for hypersensitivity reaction for a period of 60 minutes following infusion **0.9% Sodium Chloride Bolus:** (to be infused during observation period): 
250 mL NS 1,000 mL NS 1,000 mL NS

Provider name (print):	Date:
Provider signature:	Time:

Revised 1/19/2021. Order valid for one year unless otherwise indicated. IV solutions/diluents may be substituted as allowed per manufacturer's instructions as necessitated by product availability.