## Bamlanivimab Consent Casirivimab & Imdevimab Consent

Patient Name:	
DOB:	<u>.</u>
Date:	•
I authorize the intravenous administration of Bamlar the infusion nursing staff of the COVID INFUSION C	nivimab OR Casinivimab & Imdevimab as determined by medication availability by ENTER under the supervision of the CIC physicians.
not been approved by the FDA but have been author	ocional antibodies for the purpose of treating COVID-19. These medications have brized for emergency use to treat mild to moderate COVID-19 in adults and S-CoV-2 viral testing who are 12 years of age and older weighing at least 40 kg, COVID-19 and/or hospitalization.
Allergic reaction — fever, chills, low blood pressure, throat, rash including hives, itching, headache, naus getting any medication by vein may include brief pa	th Bamlanivimab and Casirivimab/Imdevimab may include, but are not limited to: changes in heartbeat, shortness of breath, wheezing, swelling of lips, face or sea, vomiting, sweating, muscle aches, dizziness, or shivering. The side effects of in, bleeding, bruising of the skin, soreness, swelling, and possible infection at the during the infusion or be delayed until after I leave the facility.
I authorize the CIC infusion nursing staff to adminis reaction. If these symptoms occur after I leave this staff at CIC.	ter medications as deemed medically necessary at the time of the infusion facility, I will seek emergency care as needed, and I will notify the infusion nursing
	nivimab and Casirivimab/Imdevimab. Not many people have been given and unexpected side effects may happen. These drugs are still being studied so ne.
of SARS-CoV-2. Similarly, Bamlanivimab and Casirivi	devimab could interfere with your body's own ability to fight off a future infection imab/Imdevimab may reduce your body's immune response to a vaccine for SARS-address these possible risks. Talk to your healthcare provider if you have any
obtained. I have received the Fact Sheet for Patie Bamlanivimab OR Casirivimab/Imdevimab for	although no guarantee or assurance has been made as to the results to be ents, Parents and Caregivers Emergency Use Authorization (EUA) of r Coronavirus Disease 2019 (COVID-19). I have read and understand this b OR Casirivimab/Imdevimab intravenous treatment.
Patient Signature:	Date:
Printed Name:	DOB:
Witnessed by	Date