



**Monoclonal Antibody**  
**CASIRIVIMAB & IMDEVIMAB Injection**  
For Accordia Health Monoclonal Antibody Clinic

**FAX COVER SHEET**

<b>To:</b>	Accordia Health Monoclonal Antibody Clinic	<b>From:</b>	
<b>Fax:</b>	251-281-2437	<b>Sender Fax:</b>	
<b>Phone:</b>	251-305-4550	<b>Sender Phone:</b>	
<b>Subject:</b>	Patient Information for appointment	<b>Date:</b>	

**No. Pages:**

**Comments:**

The following documents are REQUIRED to send:

- Physician Orders – signed, MUST be complete and include: patient ALLERGIES; date of COVID Positive test; and date of symptoms
- Patient Consent – signed by patient and provider
- Patient demographics sheet with updated phone number, last 4 of SS# and insurance information.

Accordia Health will call the patient within 24 hours to schedule appointment –

Patients on Friday should be prepared to receive injections on Monday if appropriate.

Please be sure all paperwork above is included in the fax for timely appointment scheduling.



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### PHYSICIAN'S ORDERS

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_

Weight: \_\_\_\_\_ Height: \_\_\_\_\_ ALLERGIES/Reactions: \_\_\_\_\_

Physician's Orders	
Status	Outpatient
Indication	<p>Mild to Moderate COVID-19 infection in patients 12 yr. &amp; older and weigh at least 40 kg with a high risk of progression to severe COVID-19 and/or hospitalization. Date of SARS-CoV-2 positive test result: ___/___/___ Date of symptom onset: ___/___/___</p> <p><b>High Risk Criteria:</b></p> <ul style="list-style-type: none"> <li><input type="checkbox"/> Age ≥ 65 yr</li> <li><input type="checkbox"/> Pregnancy</li> <li><input type="checkbox"/> Chronic Kidney Disease</li> <li><input type="checkbox"/> Diabetes</li> <li><input type="checkbox"/> Chronic lung disease (COPD, asthma, interstitial lung disease, cystic fibrosis, and pulmonary hypertension)</li> <li><input type="checkbox"/> Cardiovascular disease (including congenital heart disease) or hypertension</li> <li><input type="checkbox"/> Neurodevelopmental disorders (cerebral palsy) or other conditions that confer medical complexity</li> <li><input type="checkbox"/> Having medical-related technological dependence (tracheostomy, gastrostomy, or PPV (non-COVID))</li> <li><input type="checkbox"/> Other medical conditions or factors that place patients at high risk for progressing to severe COVID-19 (example: race or ethnicity)</li> <li><input type="checkbox"/> Obesity or being overweight (adults BMI &gt; 25; age 12-17 ≥ 85<sup>th</sup> percentile)</li> <li><input type="checkbox"/> Immunosuppressive disease or immunosuppressive treatment</li> <li><input type="checkbox"/> Sick cell disease</li> </ul>
Consent and Patient Information	<p>Review with patient the Fact Sheet for Patients, Parents and Caregivers Emergency Use Authorization (EUA) of Casirivimab &amp; Imdevimab for Coronavirus Disease 2019. Verify patient consent for Casirivimab &amp; Imdevimab obtained and in chart prior to administration.</p>
Clinical Care Orders	Vital signs at baseline.
	<p>Inspect vial for discoloration or particulate matter. <b>Do not expose to direct heat. Do not shake the vial.</b> Using 4 syringes, withdraw 2.5mL of Casirivimab &amp; Imdevimab into each syringe for subcutaneous injection Administer immediately (if not possible store at room temp for no more than 4 hours) If medication refrigerated, allow to equilibrate to room temp for about 20 min prior to administration. Administer the subcutaneous injections consecutively, each at a different injection site (thigh, back of upper arm, or abdomen [except for 2 inches around the navel]) The waistline should be avoided. It is recommended using different quadrants of the abdomen or upper thighs or back of the upper arms to space apart <b>DO NOT</b> inject into skin that is tender, damaged, bruised, or scarred</p>
	Vital signs every 15 minutes after injection. Clinically monitor patients for at least one hour
	Common reactions: fever, chills, nausea, headache, bronchospasm, difficulty breathing, arrhythmia, chest pain or discomfort, weakness, altered mental status, hypotension, hypertension, angioedema, throat irritation, rash including urticarial, pruritus, myalgia, vasovagal reactions (pre-syncope, syncope), dizziness fatigue and diaphoresis.
Treatment and Medications for an injection reaction	<ul style="list-style-type: none"> <li><input type="checkbox"/> Oxygen via nasal cannula or non-rebreather mask PRN difficulty breathing or decreased O2 saturation</li> <li><input type="checkbox"/> Bag-Valve Mask breathing with Oxygen PRN for apnea or slow, shallow breathing</li> <li><input type="checkbox"/> Diphenhydramine 25 mg IM once PRN for an injection reaction</li> <li><input type="checkbox"/> Dexamethasone 4 mg IM once PRN for an injection reaction</li> <li><input type="checkbox"/> Acetaminophen 650 mg PO once PRN for an injection reaction</li> <li><input type="checkbox"/> Epinephrine 0.3 mg INTRAMUSCULAR (EpiPen 0.3 mg auto-injector) once PRN for an injection reaction</li> </ul> <p>Call Physician if patient develops anaphylaxis symptoms. Call EMS/ambulance for transport to hospital</p>
Other	Report any adverse events/reactions to AltaPointe Health Systems Compliance Department and via RL-6.
Patient Instructions	Continue to self-isolate and use infection control procedures (social distance, wear mask, wash hands) per CDC Notify Healthcare provider for worsening symptoms.

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

Printed Provider Name: \_\_\_\_\_ Phone: \_\_\_\_\_

Collaborating Physician Signature (if applicable) \_\_\_\_\_



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**PATIENT CONSENT**

I have received and reviewed the Patient Fact Sheet for Casirivimab/Imdevimab.

I have been informed of alternative treatments.

I understand that this drug is not approved by the FDA, and that it has been authorized for emergency use only.

I consent to receive Casirivimab/Imdevimab treatment under Emergency Use Authorization (EUA) for COVID-19.

PATIENT Name: \_\_\_\_\_

Signature: \_\_\_\_\_

Date: \_\_\_\_ / \_\_\_\_ / \_\_\_\_      Time: \_\_\_\_\_

**HEALTHCARE PROFESSIONAL ATTESTATION**

I have explained the treatment to the patient/authorized representative and have answered all questions about this treatment to the best of my ability.

Printed Name/Credentials: \_\_\_\_\_

Signature: \_\_\_\_\_

Date: \_\_\_\_ / \_\_\_\_ / \_\_\_\_      Time: \_\_\_\_\_