Referral Form for UVA COVID-19 Clinic Antibody Infusion for Adults with COVID-19
Fax to 434.243.9800

Eligible patients can be referred to the UVA COVID-19 clinic for intravenous infusion of investigational monoclonal antibody therapy under an FDA Emergency Use Authorization.

ELIGIBILITY:

Outpatients with COVID-19 with mild-to-moderate symptoms for fewer than 10 days, without new or increasing oxygen requirement, and at greater risk for developing more serious symptoms that may require hospitalization:

- A body mass index (a body-fat measurement based on height and weight) of 35 or higher
- Chronic kidney disease
- Diabetes
- Diseases that weaken the body's immune system, such as rheumatoid arthritis and multiple sclerosis
- Receiving another treatment that weakens the immune system
- Age 65 and older
- Age 55 and older with cardiovascular disease, high blood pressure, chronic obstructive pulmonary disease (COPD) or other chronic respiratory diseases

REFERRAL:

To initiate a referral for monoclonal antibody at UVA, complete this form and fax to **434.243.9800**. The COVID clinic infusion team will review the referral form upon receipt and contact the patient to coordinate services as soon as possible. A telehealth visit will be scheduled prior to infusion, and patients prioritized for infusion will receive their monoclonal antibodies during an appointment at the UVA COVID-19 Clinic. Infusion appointments typically take 3-4 hours. Infusions are administered Mondays to Fridays each week. As therapy should be administered as soon as possible after a positive test result and within 10 days of symptom onset, it is recommended that patient referrals are made as soon as possible to allow time for COVID-19 clinic clinician review and scheduling. You can call the COVID-19 clinic 434.982.6843 with questions about referrals.

DEMOGRAPHIC INFORMATION:

Date of referral	
Patient Name	
Patient DOB	
Preferred Language	
Address	
Best Contact number	

PERTINENT MEDICAL INFORMATION:

Please include any additional information re: patient's health history and medication history. You may free text here or you attach a document that includes current problem list, health history (major surgeries, major illnesses), current medication list, and medication/food allergies.

MONOCLONAL ANTIBODY ELIGIBILITY CRITERIA:

Age: _____ years BMI: _____ kg/ m2 weight: _____ kg

SARS-CoV2 Test Result (please include copy of result): ☐ PCR ☐ Antigen ☐ positive ☐ negative ☐ pending Date: _____

SARS-CoV2 symptom onset date: _____ [Note: Antibody therapy is approved for patients with mild to moderate COVID symptoms. Asymptomatic patients likely will not benefit and should not be referred. Patients with severe symptoms should seek emergency medical attention]

SpO2: ____ ☐ On RA ☐ On chronic O2 therapy – Baseline O2 Flow rate: _____

Has the patient required an increase in O2 flow rate since becoming symptomatic with COVID? ☐ Yes ☐ No

High Risk for Severe COVID Illness (check all that apply):

□ Age ≥ 65 y/o
□ BMI ≥ 35
□ CKD Disease Stage ____ Baseline [Cr]_____
□ Diabetes Mellitus
□ Immunosuppressive Disease (e.g. leukemia, lymphoma, HIV if CD4 < 200, etc.) / Specify: _____
□ Immunosuppressive Treatment (e.g. chronic steroid, chemotherapeutic, immunomodulator) / Specify: _____
□ Age ≥ 55 y/o *and:* □ Cardiovascular Disease / Specify (e.g. CAD, cardiomyopathy, arrhythmia, CHF): _____□ HTN
□ COPD □ Other Chronic Respiratory Disease (e.g. Pulmonary Sarcoid, Pulmonary Fibrosis) / Specify: _____

REFERRING PROVIDER AGREEMENTS:

I, the referring provider, am the patient's PCP or other continuity provider and have arranged for the patient to follow up with me/my designee following Antibody infusion. Or I am an ED or Urgent Care provider who will update the patient's PCP about his/her Antibody infusion in order to arrange follow up. If the patient does not have a PCP, I will refer him/her to an appropriate provider and ensure that follow up has been arranged.

□ **Indicates Provider Agreement**

I, the referring provider, have communicated to the patient or parent/caregiver, as appropriate, information consistent with the "Fact Sheet for Patients, Parents and Caregivers" (sheets will be provided in clinic) including:

1. FDA has authorized the emergency use of monoclonal antibodies for the treatment of mild to moderate COVID-19 in those who are at high risk for progressing to severe COVID-19 and/or hospitalization
2. The potential risks and benefits are unknown
3. The patient or parent/caregiver has the option to accept or refuse this treatment and alternatives were discussed
4. Treated patients should continue to self-isolate and use infection control measures according to CDC guidelines.

□ **Indicates Provider Agreement**

I, the referring provider, have advised or will advise the patient that if his/her clinical status declines by the time of the infusion appointment, the treatment may no longer be appropriate for him/her. The patient's clinical status will be re-evaluated at the infusion center at the appointment time. If the patient is deemed in need of hospital care, s/he will be referred immediately.

□ **Indicates Provider Agreement**

The COVID Clinic staff will communicate with the referring provider regarding such matters as treatment inappropriateness for patient, ultimate completion of treatment for patient, adverse events, etc..

Name of Referring Site:
Name of Referring Provider:
Address:
Point of Contact:
Phone Number:
Fax Number:
Preferred mode of contact: □ Phone □ Fax

RESOURCES:

There are two Antibody treatments on our formulary. Patients will be scheduled for one or the other treatment based on availability of medications and logistics.
Information about both medications, Casirivimab+Imdevimab or Bamlanivimab, including Fact Sheets and Manufacturer Instructions/Package Inserts for Healthcare Providers and for Patients/Parents/Care Givers, can be found at https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization#coviddrugs (scroll to section on Drugs and Biologic Products).