McLeod Health

The Choice for Medical Excellence

COVID-19 OUTDATIENT INFLISION Referral Form

Patient Name:				Date of Birth:	Preferred Location: ☐Florence
MH Employee ID (if applicable):			(if applicable):		
Dia	gnosis:	COVID	-19 - ICD 10: U07.1 Treatment of	mild to moderate COVID-19 in patie	ents at risk of progression to severe COVID-19.
Dat	e of <u>syr</u>	nptom	onset (must be within the last	t 7-10 days):	Date of Positive Test:
201	/ID Vac	cinatio	on Status: Unvaccinated Vacci	nated ≥ 6 months ago □ Vaccinate	ed or boosted < 6 months ago
/er	ify whic	ch eligi	bility requirements your patien	nt meets (if this is not compl	ete the patient CANNOT be scheduled):
	Check	Tier	Risk Group		
		1A			o mount an adequate immune response to onditions, regardless of vaccination status
-		1B	Vaccinated pregnant person withou Unvaccinated individuals at the high o Aged ≥ 75 years o Aged ≥ 65 years with any clinical i	est risk of severe disease, defined a	s at least one of the following:
-		2	Vaccinated pregnant person with a Unvaccinated individuals at risk of so o Aged ≥ 65 years o Aged < 65 years with any clinical of	evere disease not included in Tier 1	AND meeting at least one of the following:
-		3A	Patients in this tier are only eligible	for IV remdesivir at this time due t	o limited sotrovimab supply:
			Vaccinated individuals without a bo o Aged ≥ 75 years o Aged ≥ 65 years with any clinical i	-	defined as at least one of the following:
=		3B	Patients in this tier are only eligible	for IV remdesivir at this time due t	o limited sotrovimab supply:
			Vaccinated individuals with a boostory Aged ≥ 75 years Aged 65-74 years with any clinica		ined as at least one of the following:

TREATMENT OPTIONS: Please indicate which outpatient infusion therapy the patient is being referred for: By checking a specific therapy below, the referring provider affirms the patient is eligible for this specific therapy AND the patient consents to receiving the specific therapy. ☐ Sotrovimab only Sotrovimab 500 mg IV x 1 dose ☐ Remdesivir only – contraindicated if eGFR<30 mL/min or in known hepatic disease/ALT>10x upper limit of normal Remdesivir 200 mg IV on day 1, followed by 100 mg IV daily x 2 days Baseline BUN/SCr & LFTs (if not provided from last 48-72 hours with referral) ☐ Either Sotrovimab or Remdesivir If EITHER sotrovimab or remdesivir are acceptable, the infusion center will attempt to provide sotrovimab FIRST, but if sotrovimab is unavailable, the infusion center will attempt to provide 3-day course of IV remdesivir instead. Note, patients will only receive ONE outpatient infusion therapy ☐ I have informed the patient that the infusion center will only contact the patient if they are eligible to receive COVID-19 infusion treatment and there are doses available. ☐ I have discussed the risks and benefits of this treatment(s) under the prescribing information and/or Emergency Use Authorization with the patient, and they have given verbal consent for the treatment(s). Referring Provider Signature: Provider Name & Contact Information:

Please note patients are scheduled based on appointment availability, drug allocation, and patient criteria including tier and symptom onset date. All requests may not be able to be accommodated. Referring provider will be notified if the patient cannot be scheduled. If you have any questions, please contact the COVID Call Center at 843-777-2919 and press 5.

References:

Table 1. Severe Immune Compromising Conditions

- Patients within 1 year of receiving B-cell depleting therapies (e.g., rituximab, ocrelizumab, ofatumumab, alemtuzumab)
- Patients receiving Bruton tyrosine kinase inhibitors
- Chimeric antigen receptor T cell (CAR-T) recipients
- Post-hematopoietic cell transplant recipients who have chronic graft versus host disease (GVHD) or who are taking immunosuppressive medications for another indication
- Patients with hematologic malignancies who are on active therapy
- Lung transplant recipients
- Patients who are within 1 year of receiving a solid-organ transplant (other than lung transplant)
- Solid-organ transplant recipients with recent treatment for acute rejection with T or B cell depleting agents
- Severe combined immunodeficiencies
- Untreated HIV with a CD4 cell count <50 cells/mm3

Table 2. Clinical Risk Factors for Progression to Severe COVID-19

- Adult patient BMI > 35 or pediatric patient > 85th percentile
- Immune compromising condition other than Tier 1 severe immune compromising condition
- Cancers other than Tier 1 hematologic malignancies
- Diabetes mellitus, type 1 and 2
- Chronic kidney disease
- Chronic lung disease
- Neurodevelopmental disorders

- Cerebrovascular disease
- Chronic liver disease,
- Heart conditions
- Hypertension
- Medical-related technology dependence (e.g. tracheostomy, gastrostomy)
- Sickle cell disease
- Smoking, current and former