

Patient Name: _____

DOB: _____

Age: _____

REGEN-COV™ (CASIRIVIMAB + IMDEVIMAB) SUBCUTANEOUS INJECTION ORDERS

Intravenous infusion is strongly recommended. Subcutaneous injection is an alternative route of administration when intravenous infusion is not feasible and would lead to delay in treatment.

Drug Allergies:	Weight (40 kg or more):
	Date of suspected exposure or symptom onset:

Indication/Diagnoses:

Z20.828 Contact with and (suspected) exposure to other viral communicable diseases

U7.1 COVID-19 infection

Other (include ICD-10 code(s) and description(s): _____)

Prescriber must indicate *all* of the following requirements have been met:

Patient/caregiver has been given the Fact Sheet for Patients and Parents/Caregivers

Patient/caregiver has been informed of alternatives to receiving REGEN-COV™

Patient/caregiver has been informed that REGEN-COV™ is an unapproved product that is authorized for use under an Emergency Use Authorization.

<input checked="" type="checkbox"/> Withdraw a total dose of casirivimab 600 mg/5 mL AND imdevimab 600 mg/5 mL into FOUR syringes: <ul style="list-style-type: none"> • TWO syringes, each containing casirivimab 300 mg/2.5 mL; and, • TWO syringes, each containing imdevimab 300 mg/2.5 mL. 	<p>Repeat dosing for ongoing exposure (to begin four weeks after initial dose):</p> <input type="checkbox"/> Withdraw a total dose of casirivimab 300 mg/2.5 mL AND imdevimab 300 mg/2.5 mL into TWO syringes: <ul style="list-style-type: none"> • ONE syringe containing casirivimab 300 mg/2.5 mL; and, • ONE syringe containing imdevimab 300 mg/2.5 mL. <input type="checkbox"/> Repeat every four weeks for duration of ongoing exposure.
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Consecutively administer each syringe **subcutaneously** using a 25- or 27-gauge needle in a different injection site (thigh, back of arm, or abdomen except for 2 inches around navel), spacing injections apart and avoiding skin that is tender, damaged, bruised, or scarred.

Post-treatment:

- Monitor patient for hypersensitivity reaction for a period of 60 minutes following injections.**
- If adverse reaction occurs, treat per orders/protocol as clinically indicated.
- Record vital signs immediately following injections and prior to discharge.
- Provide patient with discharge instructions.
- Send record of treatment to prescriber at fax number below.

Prescriber Name (print): _____ Fax: _____

Prescriber signature: _____ Date: _____