

Outpatient COVID-19 Infusion Checklist

Positive COVID-19 test (home tests are not accepted)
☐ Identified symptom onset date: (within 7 days)
Patient demographic sheet
Labs within 3 months for GFT and ALT Adequate renal function, GFR ≥ 30ml/min and ALT < 10 times upper limits normal
☐ Signed and dated referral/order set
Fax to 540.741.2583
540.741.2583 f additional information is needed, a Health Link nurse will call the provider; otherwise, the patient will be contacted directly for scheduling.
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f additional information is needed, a Health Link nurse will call the provider; otherwise, the patient will be contacted directly for scheduling. Provider name:

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Allergies		

DO NOT USE FELT TIP PEN

FOR THOSE ORDERS WITH OPTIONS, ITEMS MUST BE MARKED OR THE ORDER IS NOT INITIATED.

Treatment: Remdesivir should be administered as soon as possible after positive viral test for SARS-CoV-2 and within 5 days of symptom onset in adults and pediatric patients 12 years of age and older weighing at least 40 kg who are at high risk for progressing to severe COVID-19 and/or hospitalization.

High risk is defined as patients who meet at least one of the following criteria:

AGE: 60 years or greater.

WEIGHT/ BMI Ages: 12-17 (Weight MUST be more than 88 lbs (40 kg) RESOURCE: Child and Teen BMI Calculator A BMI ≥ 85th percentile for their age

Ages: 18+ Body Mass Index (BMI) ≥ 30 RESOURCE: Adult BMI Calculator | Healthy Weight, Nutrition, and Physical Activity | CDC

MEDICAL CONDITIONS:

Cardiovascular disease (including congenital heart disease); Cerebrovascular disease

Chronic lung diseases (COPD, asthma, interstitial lung disease, cystic fibrosis and pulmonary hypertension)

Chronic Kidney Disease (CKD)

Chronic Liver Disease

Diabetes (type 1 or type 2)

Hypertension

Immunosuppressive disease:

Receiving Immunosuppressive treatment (chemotherapy, transplant immunosuppressants, etc.)

Neurodevelopmental disorders (I.e., CP) or other conditions that confer medical complexity (I.e., genetic or metabolic syndromes and severe congenital anomalies)

Sickle cell disease

Pregnancy

Technological dependence (e.g. tracheostomy, gastrostomy, or positive pressure ventilation not related to COVID-19)

Date:	N	Name:	
	(OUTPATIENT 'tentative' INFUSION DATE)		(ORDERING PHYSICIAN NAME)
INDIC	ATIONS: Indicated for the treatment of	COVID-1	19 administered daily for 3

DOSING & DILUTION CHART::

→ Dilute all doses mixed in 0.9% sodium chloride 100ml

Remdesivir 3 day course of therapy			Total Volume		
Medication	Dose (mg)	Number of Vials	IV Fluid		Infusion Rate and Duration
Remdesivir Day 1	200mg	2	100ml 0.9% NaCl	100 ml	Infused over 30 minutes
Remdesivir Day 2 and 3	100mg	1			

- DO NOT infuse other medications through the same IV line > Infusion must be completed within 6.5 hours of preparation
- After infusion is completed, flush line with 20 mL of normal saline
- Infuse using standard IV tubing with in-line, non-pyrogenic, low-protein-binding filter (pore size 0.2 microns to 0.22 microns)

IV ACCESS: (check one):
Monitor insertion site and maintain tubing, dressing and cap changes per hospital standard.

Start peripheral line

VITAL SIGNS: ☑ Baseline, then at infu	ision completion	i (ivionitor for art	d reactions each	n time)
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MANAGING INFUSION RELATED EVENTS: Adult: □ DiphenhydrAMINE (Benadryl) 50 mg IV PRN Infusion reaction

☐ Methylprednisolone (Solumedrol) 40 mg IV PRN infusion reaction (IV slow push over several minutes)

For Hypersensitivity (mild-moderate):

Stop infusion and notify physician for further orders

Monitor vital signs every 15min

☑ If infusion is stopped and then restarted, resume at 10mL/hr and follow rate advance per physician's directions. For Anaphylaxis: ☑ Stop infusion and notify physician ☑ Epinephrine (1:1,000 = 1mg/mL) → Pt. Wt. 0–15 kg _ mL (0.01mL/kg) IM Once PRN Anaphylaxis

→ Pt. Wt. **15-29 kg** \square At MWH \rightarrow Initiate Code (Dial 55); At SH \rightarrow Initiate Code (Dial 55) --(PER PATIENT WEIGHT)--= 0.15 mg (0.15 mL) IM Once PRN Anaphylaxis Monitor vital signs every 15 minutes

→ Pt. Wt. 30 kg or more = 0.3 mg (0.3 mL) IM Once PRN Anaphylaxis

DISCHARGE: If NO Signs and Symptoms of reaction, discharge 60 minutes after infusion completion

Scanned:			
Clerical Associate:	Date	Time	Physician Signature
RN/LPN:	Date	inne	i flysician dignature



Patient Identification

Remdesivir **Outpatient Infusion Orderset**



Allergies		

DO NOT USE FELT TIP PEN

FOR THOSE ORDERS WITH OPTIONS, ITEMS MUST BE MARKED OR THE ORDER IS NOT INITIATED.

Treatment: Bebtelovimab should be administered as soon as possible after positive viral test for SARS-CoV-2 and within 7 days of symptom onset in adults and pediatric patients 12 years of age and older weighing at least 40 kg who are at high risk for progressing to severe COVID-19 and/or hospitalization.

High risk is defined as patients who meet at least one of the following criteria:

- Have a body mass index (BMI) ≥25, or if age 12-17 have BMI ≥85th percentile for their age and gender based on CDC growth charts.
- Pregnancy

Bebtelovimab

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Outpatient Infusion Orderset

- Have chronic kidney disease
- Have diabetes
- Have immunosuppressive disease
- Are currently receiving immunosuppressive treatment
- Are ≥65 years of age
- Have chronic lung diseases (ex. COPD, moderate to severe asthma, interstitial lung disease, cystic fibrosis, pulmonary hypertension)
- Have sickle cell disease
- Have neurodevelopmental disorders (ex. Cerebral palsy) or other conditions that confer medical complexity (ex. Genetic or metabolic syndromes and severe congenital anomalies)
- Have a medical-related technological dependence (ex. Tracheostomy, gastrostomy, or positive pressure ventilation not related to COVID
 19)
- Other medical conditions or factors (ex. Race or ethnicity) may also place individual patients at risk for progression to severed COVID 19 and authorization of bebtelovimab under the EUA is not limited to the medical conditions or factors listed above.

(OUTPATIENT 'tenta	ative' INFUSION DATE)	Name:(ORDERIN	G PHYSICIAN NAME)		
NDICATIONS: Indica	ted for the treatm	ent of COVID-19			
DOSING::					
				Total Volume	
Medication	Dose (mg)	Number of Vials			Infusion Rate and Duration
Bebtelovimab	175mg	1 of 2ml		2ml	IV Push over at least 30 sec.
V ACCESS: (check □ Start /ITAL SIGNS: ☑	one): Moniton peripheral line Baseline, then a	t injection completion	v-protein-binding filter (por ain tubing, dressing and ca (Monitor for drug reaction	p changes per hospital s	standard.
For Hypersensitivi		☐ Me	, ,		n reaction on reaction (IV slow push over several minutes)
For Anaphylaxis: ☑ ☑ At MWH → Initi ☑ Monitor vital sig	Epinephrine (1:1,0 ate Code (Dial 55); Ans every 15 minutes	200 = 1mg/mL) → Pt. Wt. 0- At SH → Initiate Code (Dial 5	15 kg =mL (0.0 (SE)(PER PATIENT WEIGHT) → Pt. Wt. 30	1mL/kg) IM Once PRN Ana → Pt. Wt. 15-29 kg : 0 kg or more = 0.3 mg (0.3	phylaxis = 0.15 mg (0.15 mL) IM Once PRN Anaphylaxis mL) IM Once PRN Anaphylaxis
DISCHARGE: If NO	Signs and Symp	otoms of reaction, discl	harge 60 minutes afte	er injection completion	
Scanned:		X	Χ	Х	
Clerical Associate RN/LPN:		Date	Time	_	Physician Signature
Dahtalasimah	4	Mary Washingt	on Healthcare	Patient I	dentification

NAME: DOB: