## SIBLEY MEMORIAL HOSPITAL- JOHNS HOPKINS MEDICINE

## **EVUSHELD ORDER FORM FOR PRE-EXPOSURE PROPHYLAXIS**

(complete	all	sections	legibly)
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	(complete all sections legibly)
Date &Time	Patient Name
DOB	Phone number
Allergy:	Address:
Name and pho	ne number of nearest relative
Previously adm	nitted to SMH or JHHS entity:YesNoUnknown
Outpa	tient request Inpatient request (SM#)
Evusheld (tixa	gevimab and cilgavimab) is available to patients who meet emergency use authorization (EUA) criteria.
<ul> <li>he/she/they submission</li> <li>If the require via secure by phone of Epic/Beaco</li> <li>Patient will be patient to schere</li> <li>Please note Explored</li> </ul>	esting provider is a <u>Sibley-based oncologist or Community-based oncologist privileged at Sibley</u> , we enters the order directly into Epic/Beacon. No pre-screening by clinical pharmacist or paper order form a required. esting provider is a <u>non-oncologist privileged at Sibley</u> , he/she/they must complete this form, and email email to the <u>SMHEvusheldAdmin@jhmi.edu OR</u> fax to pharmacy department at (202) 537-0072 followed all at (202) 537-4171. The Infusion Center clinical pharmacist will review then enter the order into on within 72 hours of receipt of request. escheduled on a daily basis excluding weekends and holidays. Sibley Infusion Center staff will call the hule administration of the therapy in the non-urgent setting. rusheld (tixagevimab and cilgavimab) supply and staffing resources may be limited and can impact patients we are able to treat.
<ul><li>Weight of</li><li>Received p</li></ul>	ria and Data Requirements for Patient: (Must complete each item as appropriate) patient is >/= 40Kg : Yes No; if no, note the patient's weight rior infusion of monoclonal antibodies for COVID-19 infection: YesNo; If yes, please e name and when received
• Received of if received	r scheduled to receive COVID-19 vaccination YesNo; if yes, when Evusheld should be administered at least 2 weeks after vaccination
<ul> <li>I confirm I</li> <li>Pat</li> <li>Pat</li> <li>19.</li> </ul>	ID-19 infection: Yes (if yes, when) No Evusheld is authorized for use in patients: Yes No ients must NOT be currently infected or having symptoms with COVID-19 <b>AND</b> ients must not have had a known recent exposure (within 14 days) to an individual infected with COVID- ditionally, patients must fall into one of the two categories below to qualify for treatment: Patients with moderate to severe immune compromise due to a medical condition or receipt of immunosuppressive medications for treatment, and who may not mount an adequate immune response to COVID-19 vaccination. Please specify
	Patients in whom vaccination with any available COVID-19 vaccine, according to the approved or authorized schedule, is not recommended due to a history of severe adverse reaction (e.g., severe allergic reaction) to a COVID-19 vaccine(s) and/or COVID19 vaccine component(s).

Initial Dosing: Tixagevimab 300 mg and cilgavimab 300 mg as a single dose  Repeat Dosing: For patients who initially received tixagevimab 150 mg and cilgavimab 150 mg (previous approved dose)  If initial dose was ≤3 months ago: administer a follow-up dose of tixagevimab 150 mg and cilgavima mg If initial dose was >3 months ago: administer tixagevimab 300 mg and cilgavimab 300 mg  M.D ()	Dosing:	EVUSHELD ORDER	R FORM FOR H (complete all se		OPHYLAXIS
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For Pharmacy/Antimicrobial Stewardship Team Use		Please Print Name	-		
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RPh Date/Time		R	RPh	Date/Time	

Updated: 7/19/22