

Patient Name: \_\_\_\_\_

DOB: \_\_\_\_\_

Age: \_\_\_\_\_

## EVUSHELD™ (tixagevimab + cilgavimab) Orders for Pre-Exposure Prophylaxis

<b>Drug Allergies:</b>	<b>Weight</b> (at least 40 kg):
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<b>Indication/Diagnoses:</b> <input type="checkbox"/> <b>Z28.04</b> Immunization not carried out because of patient allergy to vaccine or component <input type="checkbox"/> <b>D84.9</b> Immunodeficiency, unspecified <input type="checkbox"/> <b>Other</b> (include ICD-10 code(s) and description(s):
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<b>Prescriber must indicate <i>all</i> of the following requirements have been met:</b> <input type="checkbox"/> Patient/caregiver has been given the Fact Sheet for Patients and Parents/Caregivers <input type="checkbox"/> Patient/caregiver has been informed that EVUSHELD™ is an unapproved product that is authorized for use under an Emergency Use Authorization.
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<input checked="" type="checkbox"/> Using aseptic technique, prepare <b>TWO</b> separate syringes: <input checked="" type="checkbox"/> ONE syringe containing <b>1.5 mL of tixagevimab</b> ; and, <input checked="" type="checkbox"/> ONE syringe containing <b>1.5 mL of cilgavimab</b> <input checked="" type="checkbox"/> Consecutively administer each syringe <u>intramuscularly</u> in different injection sites, preferably one in each of the gluteal muscles.
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**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:** \_\_\_\_\_