



Fax your completed order to 877-734-1157

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_/\_\_\_\_/\_\_\_\_  
 Date of Last Infusion: \_\_\_\_/\_\_\_\_/\_\_\_\_ Height \_\_\_\_\_ Weight \_\_\_\_\_  
 Infusion Location: (state and site) \_\_\_\_\_

### Krystexxa® (pegloticase) Infusion Orders

Diagnosis (please provide ICD-10 code in space provided):		
_____ <small>(ICD-10)</small>	_____ <small>(ICD-10)</small>	_____ <small>(description)</small>
<input type="checkbox"/> G6PD Results _____		<input type="checkbox"/> Baseline uric acid level & date _____

- Nursing Orders:**
- Hold infusion pending provider notification if:
    - Uric acid level greater than 6 mg/dL for 2 consecutive treatments (lab orders below)
    - Patient has had more than 4 weeks between treatments (due to increased risk for adverse reaction)
    - Patient reports continued use of uric acid lowering agents (allopurinol, febuxostat, probenecid, etc.)
    - Hypertension (170/90 or symptomatic)
  - Remind patient flares may occur during first 6 months of therapy
  - Monitor vital signs every 30 minutes during infusion
  - If infusion-related reaction occurs, stop infusion, and initiate Hypersensitivity Reaction Management Policy/Protocol as clinically indicated

- Labs:**
- Obtain serum uric acid level prior to each infusion (or may use result obtained within 48 hrs prior to infusion).
  - Other: \_\_\_\_\_ Frequency: \_\_\_\_\_

**Pre-medications (to be administered once 30 minutes prior to infusion):**

<input type="checkbox"/> Tylenol 500 mg orally	<input type="checkbox"/> Loratadine 10 mg orally
<input type="checkbox"/> Solu-Medrol 125 mg IVP	<input type="checkbox"/> Must have Benadryl _____ IV or PO per PI

**Dosing:**

- Krystexxa 8mg IV every 2 weeks with weekly oral methotrexate 15mg and daily folic acid 1mg\*\*
- Methotrexate contraindicated and patient is on Krystexxa Monotherapy 8mg IV every 2 weeks

- \*\*Begin weekly Methotrexate and Folic Acid 4 weeks prior to the start of Krystexxa infusions.\*\* Observation Period:**
- Monitor patient for hypersensitivity reaction for a period of 60 minutes following each infusion
  - Record vital signs prior to discharge
  - Frequency:  Every 2 weeks       Other: \_\_\_\_\_

Additional Orders:

Provider Name (print) \_\_\_\_\_ Date: \_\_\_\_\_

Provider Signature: \_\_\_\_\_ Time: \_\_\_\_\_

Reviewed 4/13/23. Order valid for one year unless otherwise indicated. IV solutions/diluents may be substituted as allowed per manufacturer's instructions as necessitated by product availability.