

**PATIENT DEMOGRAPHICS**

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Phone: \_\_\_\_\_  
 Address: \_\_\_\_\_ City/ST/Zip: \_\_\_\_\_  
 Allergies: \_\_\_\_\_  NKDA Weight: \_\_\_\_\_  lbs  kg Height: \_\_\_\_\_  in  cm  
**Patient Status:**  New to Therapy  Dose or Frequency Change  Order Renewal

**INSURANCE INFORMATION: Please attach copy of insurance card (front and back).**

**DIAGNOSIS\***

**\*ICD 10 Code Required**

- Idiopathic chronic gout (M1A.00-M1A.09), ICD10 \_\_\_\_\_
- Lead-induced chronic gout (M1A.10-M1A.19), ICD10 \_\_\_\_\_
- Drug-induced chronic gout (M1A.20-M1A.29), ICD10 \_\_\_\_\_
- Other: \_\_\_\_\_, ICD10 \_\_\_\_\_
- Chronic gout due to renal impairment (M1A.30-M1A.39), ICD10 \_\_\_\_\_
- Other secondary chronic gout (M1A.40-M1A.49), ICD10 \_\_\_\_\_
- Chronic gout, unspecified (M1A.9XX0-M1A.9XX1), ICD10 \_\_\_\_\_

**INFUSION ORDERS**

MEDICATION	DOSE	DIRECTIONS/DURATION
Krystexxa® (Pegloticase)	8 mg	Infuse IV over 2 hours once every 2 weeks x 1 year <input type="checkbox"/> Notify physician if uric acid >6 mg/mL before infusing <input type="checkbox"/> Observe patient for 1 hour after completion of infusion

**Is patient currently receiving therapy above from another facility?**

NO  YES

If yes, Facility Name: \_\_\_\_\_

Date of last treatment: \_\_\_\_\_ Date of next treatment: \_\_\_\_\_

**PRE-MEDICATION ORDERS**

- Acetaminophen 650mg PO 30 minutes prior to infusion
- Diphenhydramine 25mg/50mg PO 30 minutes prior to infusion
- Methylprednisolone 100mg IV 30 minutes prior to infusion
- Other: \_\_\_\_\_

**LAB ORDERS**

- Labs to be drawn by:**  Infusion Center  Referring Physician
- No labs ordered at this time
  - Serum uric acid – baseline and prior to each infusion with results
  - Other: \_\_\_\_\_

**REFERRING PHYSICIAN INFORMATION**

Physician Signature: \_\_\_\_\_ Date: \_\_\_\_\_  
 Physician Name: \_\_\_\_\_ Provider NPI: \_\_\_\_\_ Specialty: \_\_\_\_\_  
 Address: \_\_\_\_\_ City/ST/Zip: \_\_\_\_\_  
 Contact Person: \_\_\_\_\_ Phone #: \_\_\_\_\_ Fax #: \_\_\_\_\_  
 Email Where Follow Up Documentation Should Be Sent: \_\_\_\_\_

**REQUIRED CLINICAL DOCUMENTATION**

**Please attach medical records: Initial H&P, current MD progress notes, medication list, and labs/test results to support diagnosis.**

**Clinical Information, select all that apply:**

- The patient has refractory chronic gout as evidenced by the following:
  - History of at least 2 gout flares in the previous 12 months
  - At least 1 gout tophus
  - Gouty arthritis
  - Other: \_\_\_\_\_
- The patient has a baseline serum uric acid of  $\geq 6$  mg/dL prior to initiating Krystexxa.
- The patient does not have Glucose-6-phosphate dehydrogenase (G6PD) deficiency.
- Krystexxa will be co-administered with weekly oral methotrexate unless there is a documented contraindication.

**LAB and TEST RESULTS (required)**

- Baseline serum uric acid level
- Glucose-6-phosphate dehydrogenase (G6PD) test
- Other: \_\_\_\_\_

**PRIOR FAILED THERAPIES FOR CHRONIC GOUT**

Medication Failed: \_\_\_\_\_ Dates of Treatment: \_\_\_\_\_ Reason for D/C: \_\_\_\_\_  
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