

**PATIENT INFORMATION**

PATIENT NAME: \_\_\_\_\_ PHONE: \_\_\_\_\_

DATE OF BIRTH: \_\_\_\_\_ SEX:  M  F HEIGHT: \_\_\_\_\_ WEIGHT: \_\_\_\_\_  LBS  KG

ALLERGIES: \_\_\_\_\_ PREFERRED CLINIC: \_\_\_\_\_

REFERRAL STATUS:  NEW REFERRAL  ORDER CHANGE  ORDER RENEWAL

**DIAGNOSIS & CLINICAL DOCUMENTATION**

\*PLEASE COMPLETE ICD-10 FOR SPECIFIC DIAGNOSIS

M1A. \_\_\_\_\_ Chronic Gout

ICD-10 CODE: \_\_\_\_\_ DESCRIPTION: \_\_\_\_\_

Is patient prescribed methotrexate or other immunomodulation therapy?  YES  NO

Date the patient started methotrexate or other immunomodulation therapy: \_\_\_\_\_

**REQUIRED DOCUMENTATION**

Insurance Information  List of Medications  Tried & failed Therapies  Most recent History & Physical  G6PD  Baseline serum uric acid

**MEDICATION ORDER**

KRYSTEXXA® (pegloticase) 8 mg / 50 mL IV over a minimum of 2 hours every 2 weeks

LABS

Obtain serum uric acid level (sUA) within 72 hours prior to each infusion  
\*Dose will be held if 2 consecutive serum uric acid levels are above 6 mg/dL

REFILL X 12 MONTHS UNLESS OTHERWISE NOTED HERE: \_\_\_\_\_

**PRE-MEDICATIONS**

**REQUIRED**

Acetaminophen: 650 mg  PO  IV  
 Diphenhydramine: 25mg  PO  IV  
 Methylprednisolone: 125mg  PO  IV

**OTHER:** \_\_\_\_\_  PO  IV

Patient to be observed for 60 minutes following each administration. Administer per protocol.  
In the event of an adverse reaction occurring in the infusion clinic, utilize the Immersiv Health adverse reaction protocol.

**PRESCRIBER INFORMATION**

PROVIDER NAME: \_\_\_\_\_ NPI #: \_\_\_\_\_

EMAIL: \_\_\_\_\_ PHONE: \_\_\_\_\_ FAX: \_\_\_\_\_

ADDRESS (INCLUDE CITY, STATE, ZIP): \_\_\_\_\_

SUPERVISING PHYSICIAN: \_\_\_\_\_ CONTACT NAME: \_\_\_\_\_  
(IF APPLICABLE)

SIGNATURE: \_\_\_\_\_ DATE: \_\_\_\_\_  
(NO STAMPS)

SUBSTITUTION PERMITTED

DISPENSE AS WRITTEN