

# KRYSTEXXA (PEGLOTICASE) ORDER FORM

## PATIENT INFORMATION

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

Mobile Number: \_\_\_\_\_ Patient Weight: \_\_\_\_\_

Allergies: \_\_\_\_\_

## DIAGNOSIS (Provider must specify)

Chronic Gout, ICD 10: M1A. \_\_\_\_\_

Other: \_\_\_\_\_

## PROVIDER INFORMATION

Provider Name (print name): \_\_\_\_\_ Provider NPI: \_\_\_\_\_

Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Contact Name: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

Email Address: \_\_\_\_\_

**Prerequisites to treatment** – ensure the following information is complete and attached with referral:

- Demographics     Labs and tests supporting diagnosis     Office/progress notes

## PRE-MEDICATIONS PER KRYSTEXXA PROTOCOL

IV corticosteroid:

Hydrocortisone 200 mg

Methylprednisolone 40 mg

Methylprednisolone 125 mg

Other: \_\_\_\_\_ Dose: \_\_\_\_\_ Route: \_\_\_\_\_

Antihistamine:

\_\_\_\_ mg Diphenhydramine  
 PO  IVP

\_\_\_\_ mg Fexofenadine PO

\_\_\_\_ mg Loratadine PO

Analgesic:

1000 mg Acetaminophen PO

Other: \_\_\_\_\_

## MEDICATION

MEDICATION	DOSE	ROUTE	FREQUENCY
Krystexxa	<input type="checkbox"/> 8 mg	<input type="checkbox"/> IV	<input type="checkbox"/> Every 2 weeks

New Start Therapy     Continuation of Therapy    Date of last dose (if applicable): \_\_\_\_\_

## LABS / SPECIAL INSTRUCTIONS