

## KRYSTEXXA (PEGLOTICASE) INFUSION ORDERS

| **Physician Signature:   | Date:                         |                          |
|--|-------------------------------|--------------------------|
| Physician Name:  | Phone:                        | Fax:                     |
|  |                               |                          |
|  |                               |                          |
|  |                               |                          |
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|  |                               |                          |
| Additional Instructions:   |                               |                          |
|  |                               |                          |
| Patient advised to take antihistamine day before infusion  |                               |                          |
| Protocol Pre-Medication Orders: □Solu-Medrol 125mg IV □Antihistamine 25mg PO/IV  |                               |                          |
| Frequency: Every 2 weeks   |                               |                          |
| *Patient will be observed 1 hr post infusion   |                               |                          |
| ☐ Krystexxa (pegloticase) 8mg IV in 250ml of NS IV over 120 minutes  |                               |                          |
|  |                               |                          |
| KRYSTEXXA ORDERS   |                               |                          |
| J Code: J2507  |                               |                          |
| ☐ Chronic Arthropathy w/o mention of tophus (tophi) (ICD-10 Code:)   |                               |                          |
| Diagnosis: ☐ Chronic Gouty Arthropathy w/tophus (tophi) (ICD-10 Code:)   |                               |                          |
|  |                               |                          |
| Allergies:   | Patient Phone:                |                          |
| Patient Name:  | DOB:                          |                          |
| ratient must have Glucose-o-phosphate denydrogenase (G   | orb) deficiency screening pri | or to initiating therapy |
| *Patient must have Uric Acid level drawn 24-72 hours prior t  *Patient must have Glucose-6-phosphate dehydrogenase (G  |                               | or to initiating thorany |
| , and the second |                               |                          |
| ☐ Clinical/Progress Notes, Labs, Tests supporting primary dia ☐ Baseline Uric Acid < 6.0 mg/dl   | agnosis                       |                          |
| ☐ This signed order form from the provider ☐ Patient demographics & insurance information  |                               |                          |
| **REQUIRED INFORMATION**   |                               |                          |