



Oregon Health & Science University  
Hospital and Clinics Provider's Orders

PO7071



ADULT AMBULATORY INFUSION ORDER  
**Pegloticase (KRYSTEXXA) Infusion**

Page 1 of 3

ACCOUNT NO.  
MED. REC. NO.  
NAME  
BIRTHDATE

*Patient Identification*

**ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK (✓) TO BE ACTIVE.**

Weight: \_\_\_\_\_ kg      Height: \_\_\_\_\_ cm

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

Diagnosis Code: \_\_\_\_\_

Treatment Start Date: \_\_\_\_\_ Patient to follow up with provider on date: \_\_\_\_\_

**\*\*This plan will expire after 365 days at which time a new order will need to be placed\*\***

**GUIDELINES FOR ORDERING**

1. Send **FACE SHEET** and **H&P** or most recent chart note.
2. **Within 48 hours prior to each treatment, uric acid level must be obtained and results must be provided to the infusion clinic.** Anaphylaxis reactions have occurred. Risk of an infusion reaction is increased if patient uric acid is greater than 6 mg/dL. Discontinue treatment if levels exceed 6 mg/dL for 2 consecutive levels.
3. Prior to treatment initiation, Glucose-6-phosphate dehydrogenase (G6PD) serum test results must be included with these orders. Contraindication for G6PD deficiency, due to the risk of hemolysis and methemoglobinemia.
4. If patient misses 2 consecutive treatments (4 weeks), provider must approve continuing therapy or treatment will be discontinued.
5. **Patient must be given prescription for an EPINEPHrine auto-injector (EPIPEN) and instructed to bring one to each infusion appointment.**
6. Discontinue use of oral antihyperuricemic agents prior to initiating and during course of therapy
7. Gout Flares: Begin prophylaxis using nonsteroidal anti-inflammatory agents (NSAID) or colchicine, unless contraindicated, beginning at least 1 week before initiation of pegloticase and continuing for at least 6 months. An increase in gout flares is frequently observed. Gout flare-ups during treatment do not warrant discontinuation of therapy.
8. Congestive Heart Failure: Exercise caution in patients who have congestive heart failure and monitor patients closely following infusion.

**PRE-MEDICATIONS:** (Administer 30 minutes prior to infusion)

- acetaminophen (TYLENOL) tablet, 650 mg, oral, ONCE, every visit
- diphenhydrAMINE (BENADRYL) injection, 25 mg, intravenous, ONCE, every visit
- loratadine (CLARITIN) tablet, 10 mg, oral, AS NEEDED, every visit  
**(Choose as alternative to diphenhydrAMINE if patient does not have a driver)**
- methylPREDNISolone sodium succinate (SOLU-MEDROL), 40 mg, intravenous, ONCE, every visit

**MEDICATIONS:**

- pegloticase (KRYSTEXXA) 8 mg in 250 mL sodium chloride 0.9 %, intravenous, over 2 hours, ONCE, every visit

**Interval:**

- Every 2 weeks for \_\_\_\_\_ doses
- Every 2 weeks until discontinued



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#### **NURSING ORDERS:**

1. TREATMENT PARAMETERS - Hold treatment and notify provider:
  - a. If G6PD results are not available prior to initiation
  - b. If uric acid level is not obtained within 48 hours prior to each treatment or if uric acid is greater than 6 mg/dL (Treatment should be discontinued if 2 or more consecutive uric acid levels are greater than 6 mg/dL)
  - c. If patient misses 2 consecutive treatments (4 weeks). Provider must approve continuing therapy or treatment will be discontinued
2. VITAL SIGNS – Monitor vital signs prior to pegloticase infusion, one hour into infusion, and at end of infusion.
3. The diluted pegloticase solution should be protected from light and used within 4 hours of dilution. Prior to administration, allow the diluted solution to reach room temperature. Do not warm to room temperature using any form of artificial heating.
4. Monitor patient closely for infusion reactions during pegloticase infusion and for 1 hour after the infusion. Advise patient that delayed hypersensitivity reactions may occur. For patients with heart failure, exacerbations can occur. Educate patient on signs and symptoms of infusion reaction, including skin rash, redness of skin, difficulty breathing, flushing, chest discomfort, chest pain, and rash.
5. Explain to patient that gout flares may initially increase when starting treatment, and medications to help reduce flares may need to be taken regularly for the first few months after therapy is started. Advise patient to continue therapy even if there are flares.
6. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, dec clotting (alteplase), and/or dressing changes.

#### **AS NEEDED MEDICATIONS:**

1. acetaminophen (TYLENOL) tablet, 650 mg, oral, EVERY 4 HOURS AS NEEDED for headache, fever, chills or malaise
2. diphenhydrAMINE (BENADRYL) capsule, 50 mg, oral, EVERY 4 HOURS AS NEEDED for itching

#### **HYPERSENSITIVITY MEDICATIONS:**

1. NURSING COMMUNICATION – If hypersensitivity or infusion reactions develop, temporarily hold the infusion and notify provider immediately. Administer emergency medications per the Treatment Algorithm for Acute Infusion Reaction (Policy HC-PAT-133-GUD). Refer to algorithm for symptom monitoring and continuously assess as grade of severity may progress.
2. diphenhydrAMINE (BENADRYL) injection, 25-50 mg, intravenous, AS NEEDED x1 dose for hypersensitivity reaction
3. EPINEPHrine HCl (ADRENALIN) injection, 0.3 mg, intramuscular, AS NEEDED x1 dose for hypersensitivity reaction
4. hydrocortisone sodium succinate (SOLU-CORTEF) injection, 100 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity reaction
5. famotidine (PEPCID) 20 mg, intravenous, AS NEEDED x1 dose, for hypersensitivity reaction



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**By signing below, I represent the following:**

I am responsible for the care of the patient (*who is identified at the top of this form*);

I hold an active, unrestricted license to practice medicine in:  Oregon  \_\_\_\_\_ (*check box that corresponds with state where you provide care to patient and where you are currently licensed. Specify state if not Oregon*);

**My physician license Number is # \_\_\_\_\_ (MUST BE COMPLETED TO BE A VALID PRESCRIPTION);** and I am acting within my scope of practice and authorized by law to order Infusion of the medication described above for the patient identified on this form.

**Provider signature:** \_\_\_\_\_ **Date/Time:** \_\_\_\_\_

**Printed Name:** \_\_\_\_\_ **Phone:** \_\_\_\_\_ **Fax:** \_\_\_\_\_

OLC Central Intake Nurse:

Phone: 971-262-9645 (providers only) Fax: 503-346-8058

**Please check the appropriate box for the patient's preferred clinic location:**

**Beaverton**

OHSU Knight Cancer Institute  
15700 SW Greystone Court  
Beaverton, OR 97006  
Phone number: 971-262-9000  
Fax number: 503-346-8058

**NW Portland**

Legacy Good Samaritan campus  
Medical Office Building 3, Suite 150  
1130 NW 22nd Ave.  
Portland, OR 97210  
Phone number: 971-262-9600  
Fax number: 503-346-8058

**Gresham**

Legacy Mount Hood campus  
Medical Office Building 3, Suite 140  
24988 SE Stark  
Gresham, OR 97030  
Phone number: 971-262-9500  
Fax number: 503-346-8058

**Tualatin**

Legacy Meridian Park campus  
Medical Office Building 2, Suite 140  
19260 SW 65th Ave.  
Tualatin, OR 97062  
Phone number: 971-262-9700  
Fax number: 503-346-8058

Infusion orders located at: [www.ohsuknight.com/infusionorders](http://www.ohsuknight.com/infusionorders)