

KRYSTEXXA (PEGLOTICASE) INFUSION ORDERS

A Carelon Company

P: 877.365.5566 | F: 855.889.2946

PATIENT INFORMATION: Fax completed form, insurance information, and clinical documentation to 855.889.294					
Patient Name: DOB: Phone: Patient Status: New to Therapy Continuing Therapy Next Treatment Date:					
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MEDICAL INFORMATION					
Diagnosis: Chronic Gouty Arthropathy w/tophus (tophi) Chronic Arthopathy w/o mention of tophus (tophi) Other:					
ICD-10 Code:					
Weight: lbs Allergies:					
THERAPY ORDER					
Krystexxa Dose: 8mg IV in 250mL of NS IV over 120 minutes *Patient will be observed 1 hour post infusion					
Frequency: Every 2 weeks					
Refills: 1 year Other:					
If you would like Paragon to dispense the methotrexate, please check appropriate box: Methotrexate 15mg PO weekly x1 year (to begin 4 weeks prior to Krystexxa)					
Protocol Pre-Medication Orders: Solu-Medrol 125mg IV, Benadryl 25mg PO/IV *Patient advised to take antihistamine day before infusion					
Lab Orders: Serum uric acid 24-72 hours prior to infusion G6PD serum level (required prior to first dose) Other lab orders:					
Labs: Required labs to be drawn by 🛛 Infusion Center 🗆 Referring Provider					

Other orders: ____

	ur services, you are authorizing <i>Paragon I</i> prescription insurance companies, and to			horization and spe	cialty pharmacy designated
Provider Name:		_ Signature:	-	[Date:
Provider NPI:	Phone:	Fax:	Date: Contact Person:		
□ Opt out of Paragor	n selecting site of care (if	checked, please list site	e of care):		
PREFERRED LOC	ATION				
City:	State:	View our loca	ations here:		
	ended to be delivered only to the name red addressee, you should not disseminate				





COMPREHENSIVE SUPPORT FOR KRYSTEXXA (PEGLOTICASE) THERAPY

A Carelon Company

PATIENT INFORMATION:

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Patient Name:	DOB:
REQUIRED DOCUMENTATION FOR REFERRAL PROC	ESSING & INSURANCE APPROVAL
Include signed and completed order (MD/prescriber	to complete page 1)
Include patient demographic information and insura	nce information
Include patient's medication list	
Krystexxa Service Request form	
Supporting clinical notes (H&P) to support primary of	diagnosis
*Product information suggests the co-administration of weekly oral methotrex	ate 15mg and folic acid or folinic acid
supplementation. Krystexxa alone may be used in patients where methotrexa	te is contraindicated or not clinically appropriate.
If co-administering with methotrexate, start weekly methotrexate and folic or	folinic acid supplementation at least 4 weeks prior to
initiating, and throughout treatment with Krystexxa*	
☐ Will the patient co-administer methotrexate or o	ther immunomodulation therapy?
☐ Yes ☐ No If yes, which drug?	
\square Documentation of frequency and date of flares in	n the last 18 months (either attach
or document here):	
Has the patient tried and failed Allopurinol/Ulori	c, Colchicine, or Probenecid?
Yes No If yes, which drug(s)?	
Labs attached, including:	
Baseline serum uric acid (required)	
G6PD serum level (required)	
\Box It is recommended that patients discontinue oral ura	ate-lowering medications before
starting Krystexxa	
Other medical necessity:	

Paragon Healthcare will complete insurance verification and submit all required documentation for approval to the patient's insurance company for eligibility. Our team will notify you if any additional information is required. We will review financial responsibility with the patient and refer him/her to any available co-pay assistance as needed. Thank you for the referral.

Please fax all information to (855) 889-2946 or call (877) 365-5566 for assistance

PARAGONHEALTHCARE.COM

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