

Krystexxa® (pegloticase) Standard Plan of Treatment

PATIENT DEMOGRAPHICS:

Date of Referral:	Patient's Phone:
Patient Name:	Address:
Date of Birth:	City, State, Zip:
Height in inches:	Weight: LB or KG
Gender:	Allergies:
	See list NKDA

DIAGNOSIS: (PLEASE COMPLETE 2ND AND 3RD DIGITS TO COMPLETE ICD 10 FOR BILLING)

M1A. _____ - Chronic gout, without tophi
M1A. _____ - Chronic gout, with tophi
M10. _____ - Idiopathic gout
_____ - Other:

REQUESTED DOCUMENTATION:

1	Insurance information	IF NO:	IF YES:
2	Most recent History & Physical	PLEASE STATE	LAST INFUSION DATE:
3	Full medication list	REQUIRED WASHOUT	NEXT INFUSION DATE:
4	Tried and failed therapies	FROM PREVIOUS	IF ORDER CHANGE:
5	Baseline serum uric acid level	THERAPY:	Continue current order until insurance approved
6	G6PD serum level		
7	Specify if patient is prescribed prophylaxis for gout flare:	8	Specify if patient is prescribed methotrexate or other immunomodulation therapy:

MEDICATION ORDERS:

NOTE: Patient may be ineligible to receive Krystexxa® if patient has a diagnosis of G6PD or has new or worsening symptoms of CHF. If appropriate, it is recommended that Krystexxa® be coadministered with methotrexate and Folic Acid.

PREMEDICATION TO BE ADMINISTERED 30 MINUTES PRIOR TO ADMINISTRATION AS SELECTED

Manufacturing guidelines suggest the administration of IV corticosteroids and antihistamine prior to administration of Krystexxa®.

IV	Diphenhydramine	25mg	50mg		PO	Acetaminophen	325mg	500mg	650mg	1000mg
	Methylprednisolone	40mg	125mg	Other:		Famotidine	20mg	40mg		
	Famotidine	20mg	40 mg			Diphenhydramine	25mg	50mg		
	Other:					Fexofenadine	60mg	180mg		
						Cetirizine	10mg			
						Loratadine	10mg			
						Other:				

MEDICATION/DOSE:

☒ Krystexxa® (pegloticase) 8 mg in 250ml NS IV to infuse over 2 hours

Monitor patient for one (1) hour post infusion completion.

FREQUENCY:

☐ Dosing every 2 weeks

☐ Other: _____

LAB PARAMETERS: (Pharmacist to perform clinical lab monitoring)

Serum uric acid level preferred 48 hours prior to each infusion. Hold infusion if 2 consecutive levels are above 6 mg/dL or unknown.

Please ensure all lab work is faxed to Palmetto Infusion Services

SPECIAL/LAB ORDERS:

☐ _____

If 2 doses (4 weeks) of therapy are missed, then referring provider must give written clearance to resume therapy or treatment will be discontinued.

☒ Refills x 12 months unless noted otherwise here:

LINE USE/CARE ORDERS:

- ☒ Start PIV/Access CVC
- ☒ Flush device per facility standard flushing procedure

ADVERSE REACTION & ANAPHYLAXIS ORDERS:

Administer acute infusion and anaphylaxis medications per Palmetto Infusion standing adverse reaction orders, which can be found at our website or scan here.



PRESCRIBER INFORMATION:

PROVIDER NAME:	PHONE:
ADDRESS:	FAX:
CITY, STATE, ZIP:	NPI:

PRESCRIBER SIGNATURE: (No stamp signatures)

DATE:

Dispense as written/Brand medically necessary	Substitution permitted
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