

KRYSTEXXA (pegloticase) PRIOR AUTHORIZATION FORM

Prior authorization guidelines for **Antihyperuricemics** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		total # of pgs: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street address:	
Beneficiary name:			Suite #:	City/state/zip:
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

Drug requested:	<input type="checkbox"/> Krystexxa 8 mg/ml vial	<input type="checkbox"/> Krystexxa _____
Directions:	Quantity:	Refills:
Diagnosis (<i>submit documentation</i>):	Dx code (required):	

ALL requests

Is Krystexxa being prescribed by or in consultation with a specialist?	<input type="checkbox"/> Yes <i>Submit documentation of consultation if applicable.</i> <input type="checkbox"/> No
Does the beneficiary have glucose-6-phosphate dehydrogenase (G6PD) deficiency?	<input type="checkbox"/> Yes <i>Submit documentation of G6PD screening for at-risk beneficiaries.</i> <input type="checkbox"/> No
Will the beneficiary be using Krystexxa concomitantly with any oral urate-lowering medications?	<input type="checkbox"/> Yes <i>Submit beneficiary's current complete medication list.</i> <input type="checkbox"/> No

INITIAL requests

Does the beneficiary have a history of trial and failure of maximally tolerated doses of xanthine oxidase inhibitors (e.g., allopurinol, febuxostat) as indicated by any of the following? <i>Check all that apply.</i> <input type="checkbox"/> Continues to have frequent gout flares (≥2 flares per year) <input type="checkbox"/> Has non-resolving subcutaneous tophi	<input type="checkbox"/> Yes <i>Submit documentation.</i> <input type="checkbox"/> No
Does the beneficiary have a recent uric acid level that is above goal (based on ACR guidelines) despite maximally tolerated doses of xanthine oxidase inhibitors (e.g., allopurinol, febuxostat)?	<input type="checkbox"/> Yes <i>Submit lab results.</i> <input type="checkbox"/> No
Does the beneficiary have a contraindication or an intolerance to maximally tolerated doses of xanthine oxidase inhibitors (e.g., allopurinol, febuxostat)?	<input type="checkbox"/> Yes <i>Submit documentation.</i> <input type="checkbox"/> No
Was the beneficiary counseled regarding the following? <i>Check all that apply.</i> <input type="checkbox"/> Appropriate dietary and lifestyle modifications <input type="checkbox"/> Discontinuation of other medications known to precipitate gout attacks	<input type="checkbox"/> Yes – <i>Submit documentation.</i> <input type="checkbox"/> No

RENEWAL requests

Did the beneficiary experience improvement in disease severity since initiating treatment with Krystexxa?	<input type="checkbox"/> Yes <i>Submit documentation of clinical response.</i> <input type="checkbox"/> No
---	---

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION

Prescriber Signature:	Date:
-----------------------	-------

Confidentiality Notice: The documents accompanying this telecopy may contain confidential information belonging to the sender. The information is intended only for the use of the individual named above. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or taking of any telecopy is strictly prohibited.