



Prior Authorization Request Form for Antihyperuricemics

FAX this completed form to (844) 205-3386

OR Mail requests to: Pharmacy Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720

OR Prior authorization may be completed at <https://www.covermymeds.com/main/prior-authorization-forms/>

I. PROVIDER INFORMATION		II. MEMBER INFORMATION	
Prescriber Name:	Member Name:		
Prescriber Specialty:	Identification #:		
NPI:	Group #:		
Office Contact Name:	Date of Birth:		
Fax #:	Medication Allergies:		
Phone #:			
III. DRUG INFORMATION (One drug request per form)			
Drug name and strength:	Dosage Interval (sig):	Qty. per Day:	
IV. REQUIRED DOCUMENTATION (Detailed medical record documentation demonstrating evidence for each item must be submitted with prior authorization request)			
Specify diagnosis & diagnosis code relevant to this request:		Dx/Dx Code: _____	
Requests for non-preferred Xanthine Oxidase Inhibitor: Does the member have a history of trial and failure of or contraindication or intolerance to the preferred Xanthine Oxidase Inhibitor? Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred medications in this class.	<input type="checkbox"/> Yes <input type="checkbox"/> No	Medications Taken (start and end date and dose): _____ _____ _____	
Requests for non-preferred single-ingredient Colchicine: Does the member have a history of trial and failure of or contraindication or intolerance to the preferred single-ingredient Colchicine? Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred medications in this class.	<input type="checkbox"/> Yes <input type="checkbox"/> No	Medications Taken (start and end date and dose): _____ _____ _____	
Requests for non-preferred Antihyperuricemic: Does the member have a history of trial and failure of or contraindication or intolerance to the preferred single-ingredient Antihyperuricemic? Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred medications in this class.	<input type="checkbox"/> Yes <input type="checkbox"/> No	Medications Taken (start and end date and dose): _____ _____ _____	
Does the member have a history of contraindication to the prescribed medication?	<input type="checkbox"/> Yes <input type="checkbox"/> No		
<input type="checkbox"/> If requesting for daily quantity exceeding daily limit (Refer to https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx), please provide supporting information: _____			
SUBMIT MEDICAL RECORD INFORMATION FOR EACH APPLICABLE ITEM.			
KRYSTEXXA:			
<input type="checkbox"/> If not prescribed by one of the following specialist rheumatologist or endocrinologist, please indicate a specialist consulted: _____			
<input type="checkbox"/> Recent uric acid level above goal based on American College of Rheumatology guidelines: _____ (submit labs)			
<input type="checkbox"/> One of the following:			
<input type="checkbox"/> Continues to have frequent gout flares (≥ 2 flares/year)			
<input type="checkbox"/> Has non-resolving subcutaneous tophi			
<input type="checkbox"/> Krystexxa will not be used concomitantly with oral urate-lowering agents			

- Member was counseled regarding the following:
 - Appropriate dietary and lifestyle modifications
 - Discontinuation of other medications know to precipitate gout attacks

KRYSTEXXA RENEWAL REQUEST:

- Documentation of improvement in disease severity since initiating Krystexxa, as evidenced by: _____
- If not prescribed by one of the following specialist rheumatologist or endocrinologist, please indicate a specialist consulted: _____
- Does not have a history of a contraindication to Krystexxa
- Krystexxa will not be used concomitantly with oral urate-lowering agents

IV. ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION :

Appropriate clinical information to support the request on the basis of medical necessity must be submitted.	Provider Signature:	Date:
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Pharmacy Department will respond via fax or phone within 24 hours.
 Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)