

Prior Authorization Request Form for Colony Stimulating Factors

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.covernymeds.com/main/prior-authorization-forms/

OKTITOT authorization may be comp	neted at https://		i my measies.		
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	g request per for	m)			
Drug name and strength:	Dosage Interval (si	g):		Qty. per Day:	
IV. REQUIRED DOCUMENTION (Detailed medical record documentation demonstrating evidence for each					
item must be submitted with prior of	authorization rea	quest)			
Specify diagnosis & diagnosis code relevant to this request:					
specify diagnosis & diagnosis code relevant to this request.			Dx/Dx Code:		
Does the member have a history of a contraindication to the requested medication?			□ Yes		
			□ N		
			🗆 No		
Requests for all non-preferred Colony Stimulating Factor the member have a history of trial and failure of or contrained or intolerance to the preferred Colony Stimulating Factors? <i>I</i> <u>https://papdl.com/preferred-drug-list</u> for a list of preferred and preferred medications in this class.			🗆 No	Submit documentation of previous trials/failures, contraindications, and/or intolerances or current use.	
☐ If requesting for daily quantity en <u>Services/Pages/Quantity-Limits</u> information:					
SUBMIT MEDICAL RECORD INFORMATIC INITIAL REQUEST:					
If not prescribed by the following specialist, a hematologist or oncologist, please indicate a specialist consulted:					
For primary prophylaxis of chemotherapy-induced febrile neutropenia in patients with non-myeloid malignancies, one of the following:					
Will be receiving a chemotherapy regimen with an expected incidence of febrile neutropenia > 20% as defined by the National Comprehensive Cancer Network (NCCN):					
Has risk factors for developing febrile neutropenia as defined by the NCCN:					
For Neulasta (pegfilgrastim), will 14 days before and ending 24 hou				e medication during the period beginning therapy	
RENEWAL REQUIEST:					
Member has demonstrated tolerability and a positive clinical response based on the prescriber's assessment:					

IV. ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION :					
Appropriate clinical information to support the request on	Provider Signature:	Date:			
the basis of medical necessity must be submitted.		Date.			

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.)