



COLONY STIMULATING FACTORS PRIOR AUTHORIZATION FORM

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/>

Prior authorization guidelines for **Colony Stimulating Factors** and **Quantity Limits/Daily Dose Limits** are available on the PA Health & Wellness website at <https://www.pahealthwellness.com/providers/pharmacy.html>

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

CLINICAL INFORMATION

Drug requested:	Strength:	Dosage form (e.g., vial, syringe, kit, etc.):	
Dose/route/frequency:		Quantity:	Refills:
Diagnosis (<i>submit documentation</i>):		DX code (<i>required</i>):	
Member's height: _____ ins / cms	Member's weight: _____ lbs / kg	BSA (<i>Leukine only</i>): _____ m ²	

Complete the sections below that are applicable to the member and this request and SUBMIT DOCUMENTATION for each item.

- Has recent results of a CBC with differential
- Is or will be receiving chemotherapy
- Is or will be receiving radiation therapy:
 - Dates or planned dates of radiation: _____
- For a NON-PREFERRED Colony Stimulating Factor (CSF):**
 - Has a history of trial and failure of or a contraindication or an intolerance to the preferred Colony Stimulating Factors that are approved or medically accepted for treatment of the member's diagnosis (*Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.*): _____
- Prophylaxis of chemotherapy-induced febrile neutropenia:**
 - Has at least 1 of the following risk factors for the development of febrile neutropenia:
 - Age >65 years
 - Recent surgery
 - History of febrile neutropenia
 - Poor liver or kidney function
 - Current infection or open wound

- Previous chemotherapy or radiation
- Cardiovascular disease
- Poor nutritional or performance status
- other: _____

Receiving or will receive a chemotherapy regimen with an expected incidence of neutropenia >20%

For pegfilgrastim (Neulasta, Udenyca, etc.):

Last date of chemo: _____

Planned administration date: _____

Next expected chemo date: _____

Treatment of febrile neutropenia:

- Received or is receiving myelosuppressive anticancer drugs associated with neutropenia
- Is at high risk for infection-related complications

Bone marrow transplant:

Has a non-myeloid malignancy and is undergoing myeloablative chemotherapy to be followed by bone marrow transplant

Planned transplant date: _____

Has non-Hodgkin's lymphoma, acute lymphoblastic leukemia, or Hodgkin's lymphoma and had an autologous bone marrow transplant

Transplant date: _____

Stem cell transplant:

- Is planned for autologous peripheral stem cell transplant
- Is planned for allogeneic peripheral stem cell transplant
- Will be using the requested medication in combination with plerixafor (*also complete Mozobil prior authorization form*) or another stem cell mobilizer

Planned leukapheresis date: _____

Planned transplant date: _____

Had an autologous or allogeneic peripheral stem cell transplant

Transplant date: _____

Acute myeloid leukemia:

- CSF will be used following induction chemotherapy
- CSF will be used following consolidation chemotherapy
- other: _____

Hematopoietic syndrome of acute radiation syndrome:

Suspected or confirmed exposure to a radiation dose >2 gray (Gy)

Severe chronic neutropenia – specify type: congenital neutropenia cyclic neutropenia idiopathic neutropenia

Experiencing symptoms of neutropenia

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO 844-205-3386

Prescriber 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.)