## COLONY STIMULATING FACTORS PRIOR AUTHORIZATION FORM







(form effective 1/6/2025)

Fax to PerformRx<sup>SM</sup> at **1-855-851-4058**, or to speak to a representative call **1-888-674-8720**.

PRIOR AUTHORIS	ATION DEGLIES	TINEODMATI	ON					
PRIOR AUTHORIZ  ☐ New request ☐ Re	enewal request	Total # of pages:						
Name of office contact:				LTC fa	TC facility contact/phone:			
			Oontdot o	3110110 111	anibor.	LIOIU	omity contact phone.	
PATIENT INFORM	ATION			Dotion	ID #-		DOB:	
Patient name: Street address:				Patient ID #:			DOB.	
	City/atata/ain	state/zip: Phone:						
Apt #:	City/state/zip:				Priorie:			
PRESCRIBER INFO	ORMATION							
Specialty:				N	IPI:		State license #:	
Street address:								
Suite #:	City/state/zip:							
Phone:			Fax:					
CLINICAL INFORM	MATION							
Medication requested:								
Preferred:			Non-Preferred:					
☐ Fulphila (pegfilgrastim-jmdb) Syringe			☐ Fylnetra (pegfilgrastim-pbbk) Syringe					
☐ Granix (tbo-filgrastim) Syringe				☐ Leukine (sargramostim) Vial				
☐ Granix (tbo-filgrastim) Vial			□ Neulasta (pegfilgrastim) Onpro					
☐ Neupogen (filgrastim) Syringe			☐ Neulasta (pegfilgrastim) Syringe					
☐ Neupogen (filgrastim) Vial			☐ Nivestym (filgrastim-aafi) Syringe					
☐ Releuko (filgrastim-ayow) Syringe			□ Nivestym (filgrastim-aafi) Vial					
☐ Releuko (filgrastim-ayow) Vial			☐ Nyvepria (pegfilgrastim-apgf) Syringe					
					vedon (eflapegrastim-xnst) Syrin			
				☐ Stir	nufend (pegfilgrastim-fpgk) Syrir	nge		
				□ Ude	enyca (pegfilgrastim-cbvq) Autoir	njector		
					enyca (pegfilgrastim-cbvq) Onboo	•		
					enyca (pegfilgrastim-cbvq) Syring	ge		
					xio (filgrastim-sndz) Syringe			
				□ Zie	ktenzo (pegfilgrastim-bmez) Syrii	nge		
Dosage form (e.g., vial, sy	ringe, kit, etc.):					!	Strength:	
Dose/route/frequency:				(	Quantity:	I	Refills:	
Diagnosis (submit docum	entation):					I	Ox code <u>(required)</u> :	
Beneficiary's height:	in. / cm	1	Beneficiary's weight:	-	lbs / kg		BSA (Leukine only):	m²



NITIAL REQUESTS
Complete all sections that apply to the beneficiary and this request.  Check all that apply and submit documentation for each item.
☐ Has recent results of a CBC with differential (submit copy of results) ☐ Is or will be receiving chemotherapy.  List chemotherapy regimen: ☐ 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 beneficiary's diagnosis (Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.)  List medications tried:
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:
6. <u>Treatment of febrile neutropenia:</u> ☐ 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:
5. 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:
6. Acute myeloid leukemia:  CSF will be used following induction chemotherapy  CSF will be used following consolidation chemotherapy  other:
<ul> <li>✓. Hematopoietic syndrome of acute radiation syndrome:</li> <li>☐ Suspected or confirmed exposure to a radiation dose &gt;2 gray (Gy)</li> </ul>
3. <u>Severe chronic neutropenia — specify type:</u> □ congenital neutropenia □ cyclic neutropenia □ idiopathic neutropenia □ Experiencing symptoms of neutropenia
PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber signature:

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.

Coverage by AmeriHealth First.

KF\_244041800-4 Page 2 of 2