



**Fax completed prior authorization request form to 844-802-1412 or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts.**

All requested data must be provided. **Incomplete forms or forms without the chart notes will be returned.**

Aetna Better Health®

Pharmacy Coverage Guidelines are available at <https://www.aetnabetterhealth.com/illinois-medicaid>

## Colony Stimulating Factors Pharmacy Prior Authorization Request Form

**Do not copy for future use. Forms are updated frequently.**

**REQUIRED: Medical records, including labs and weight or body surface area (BSA), to support diagnosis are required to be submitted**

Member Information					
Member Name (first & last):	Date of Birth:	Gender:		Height:	
		<input type="checkbox"/> Male	<input type="checkbox"/> Female		
Member ID:	City:	State:		Weight:	
Prescribing Provider Information					
Provider Name (first & last):	Specialty:	NPI#		DEA#	
Office Address:	City:	State:		Zip Code:	
Office Contact:	Office Phone		Office Fax:		
Dispensing Pharmacy Information					
Pharmacy Name:		Pharmacy Phone:		Pharmacy Fax:	
Requested Medication Information					
Preferred Agents:	<input type="checkbox"/> Leukine		<input type="checkbox"/> Neupogen		
Non-Preferred Agents:	<input type="checkbox"/> Zarxio	<input type="checkbox"/> Nivestym	<input type="checkbox"/> Granix	<input type="checkbox"/> Fulphila	
	<input type="checkbox"/> Neulasta Onpro	<input type="checkbox"/> Neulasta	<input type="checkbox"/> Ziextenzo	<input type="checkbox"/> Udenyca	
<input type="checkbox"/> Other, please specify:					
Are there any contraindications to formulary medications?				<input type="checkbox"/> Yes	<input type="checkbox"/> No
If yes, please specify:				<input type="checkbox"/> New request	
<input type="checkbox"/> RENEWAL ONLY (check that apply):	<input type="checkbox"/> Request that	<input type="checkbox"/> Response to therapy	<input type="checkbox"/> Recent ANC, CBC, and/or platelet counts	<input type="checkbox"/> For Chemotherapy induced neutropenia: Recent ANC showing response to therapy	
Directions for Use:	Strength:		Dosage Form:		
	Quantity:	Day Supply:	Duration of Therapy/Use:		
What medication(s) has member tried and failed for this diagnosis? Please specify:					
Medication request is NOT for an FDA approved, or compendia-supported diagnosis (circle one): Yes No		Diagnosis:		ICD-10 Code:	
Turn-Around Time for Review					
<input type="checkbox"/> Standard – (24 hours)		<input type="checkbox"/> <b>Urgent</b> – waiting 24 hours for a standard decision could seriously harm life, health, or ability to regain maximum function, you can ask for an expedited decision.			
		Signature: _____			
Clinical Information					

Will requested medication be used concomitantly with radiation AND chemotherapy?				<input type="checkbox"/> Yes	<input type="checkbox"/> No
Will requested medication be administered at appropriate time after chemotherapy OR radiation?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Will requested medication be used in combination with other myeloid growth factors?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
<b>Chemotherapy-Induced Febrile Neutropenia</b>					
<input type="checkbox"/> <b>PRIMARY Prophylaxis</b>					
Member is receiving chemotherapy for NON-myeloid cancer AND meets ONE of the following (check that apply):			<input type="checkbox"/> Chemotherapy regimen is given after bone marrow transplant		
			<input type="checkbox"/> Chemotherapy regimen has >20% risk of febrile neutropenia		
			<input type="checkbox"/> Chemotherapy regimen has 10%-20% risk of febrile neutropenia AND ANY of the following risk factors for febrile neutropenia:		
			<input type="checkbox"/> Age > 65 years	<input type="checkbox"/> Persistent neutropenia	
			<input type="checkbox"/> Prior chemo or radiation	<input type="checkbox"/> Renal dysfunction CrCl < 50	
			<input type="checkbox"/> Bone marrow involvement by tumor	<input type="checkbox"/> Liver dysfunction (bilirubin > 2.0)	
			<input type="checkbox"/> Recent surgery and/or open wounds		
<input type="checkbox"/> <b>SECONDARY Prophylaxis</b>					
Has member previously experienced febrile neutropenia from same chemotherapy regimen?				<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> <b>TREATMENT</b>					
Has member received long acting CSF for prophylaxis?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	If received Zarxio, Nivestym, Neupogen, or Granix, will there be continuation with same agent?	<input type="checkbox"/> Yes	<input type="checkbox"/> No <input type="checkbox"/> N/A
Prophylactic therapy with a CSF was not received AND risk factors for poor outcome resulting from febrile neutropenia are present (check that apply):			<input type="checkbox"/> Age > 65		<input type="checkbox"/> Current infection
			<input type="checkbox"/> Sepsis		<input type="checkbox"/> Hospitalized at onset of fever
			<input type="checkbox"/> Severe neutropenia (ANC < 100/mcL)		<input type="checkbox"/> Prior episode of febrile neutropenia
<input type="checkbox"/> <b>Other Indications</b>					
Check below box for applicable diagnosis:	Member has one of the following:		<input type="checkbox"/> Evidence of inadequate bone marrow reserve		
<input type="checkbox"/> Severe chronic congenital neutropenia			<input type="checkbox"/> High risk for developing serious bacterial infection		
<input type="checkbox"/> Idiopathic Neutropenia			<input type="checkbox"/> Current bacterial infection		
Check below box for applicable diagnosis:					
<input type="checkbox"/> Neutropenia related to HIV	<input type="checkbox"/> Prescribed by or in consultation with an Infectious Disease Specialist, Hematologist, or HIV Specialist				
<input type="checkbox"/> Neutropenia related to drug therapy - ganciclovir or zidovudine induced:					
<input type="checkbox"/> Neupogen – Zarxio – Nivestym may also be approved for the following indications:	Member has ONE of the following (check that apply):				
	<input type="checkbox"/> AML in members receiving induction or consolidation chemotherapy				
	<input type="checkbox"/> Mobilization of hematopoietic progenitor cells before autologous stem cell transplant				
	<input type="checkbox"/> Mobilization of hematopoietic progenitor cells in donor before allogenic stem cell transplant				
	<input type="checkbox"/> Treatment of acute radiation exposure in members who receive myelosuppressive doses of radiation at a dose of 2 gray (Gy)				
	<input type="checkbox"/> MDS or aplastic anemia in a member with ANC <500				

<input type="checkbox"/> Leukine may also be approved for the following indications:	Member has ONE of the following (check that apply):
	<input type="checkbox"/> AML after induction chemotherapy for ages 55 years or older
	<input type="checkbox"/> Bone marrow transplant failure or engraftment delay
	<input type="checkbox"/> Myeloid reconstitution after autologous bone marrow transplant for Hodgkin's disease, non-Hodgkin's lymphoma or ALL
	<input type="checkbox"/> Before and after autologous peripheral blood stem cell transplantation
	<input type="checkbox"/> Members acutely exposed to myelosuppressive doses of radiation, administer once daily as subcutaneous injection

Additional information the prescribing provider feels is important to this review. Please specify below or submit medical records

Signature affirms that information given on this form is true and accurate and reflects office notes.

Prescribing Provider's Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**Please note: Incomplete forms or forms without the chart notes will be returned.**  
 Medical records, including labs and weight or body surface area (BSA), to support diagnosis are required to be submitted.  
 Standard turnaround time is 24 hours. You can call 866-329-4701 to check the status of a request.