## Prior Authorization

## AETNA BETTER HEALTH OF ILLINOIS FAMILY HEALTH PLAN (MEDICAID) Neulasta (IL88)

This fax machine is located in a secure location as required by HIPAA regulations.

Complete/review information, sign and date. Fax signed forms to Aetna Better Health Illinois Medicaid at **1-844-242-0908**. Please contact Aetna Better Health Illinois Medicaid at **1-866-212-2851** with questions regarding the Prior Authorization process.

When conditions are met, we will authorize the coverage of Neulasta (IL88).

Please note that all authorization requests will be reviewed as the AB rated generic (when available) unless states otherwise.

Drug Name (select	from list of drugs shown)			
Neulasta (pegfilgrastin	1)			
Quantity	Frequency		Strength	
Route of Administration	n Expected Length of th	nerapy		
Patient Information	n			
Patient Name:				
Patient ID:		_		
Patient Group No.:		_		
Patient DOB:		_		
Patient Phone:		_		
Prescribing Physic	ian			
Physician Name:	<u> </u>			
Physician Phone:				
Physician Fax:				
Physician Address:				
City, State, Zip:				
Diagnosis:	ICD Code:			
	priate answer for each question.			
при				
1. Has Aetna Better	Health authorized this medication in th	e Y	Ν	
past for this patier	nt (i.e., previous authorization is on file			
under Aetna Bette	er Health)?			
[If yes, skip to que	estion 9.1			
in yes, ship to que	.5.1011 5.1			
2. Is the patient an a	adult or an adolescent who weighs	Υ	N	
greater than or ed	•	•		
[If no, no further q	uestions.]			
3. Is therapy prescri	bed by a hematologist and/or	Υ	N	
oncologist?	,			
[]{	westings 1			
[If no, no further q	uestions.]			

[If no, no further questions.]  5. Is the request for primary prophylaxis of chemotherapy-induced neutropenia? Please document # of chemotherapy cycles:  [If no, no further questions.]  6. Does the patient meet ONE of the following conditions? YN  Patient is receiving a myelosuppressive chemotherapy regimen that has an expected incidence of febrile neutropenia greater than or equal to 17% and chemotherapy cycle of greater than 14 days \ Patient is at high risk for neutropenic complications (e.g., age greater than 65 years, pre-existing neutropenia, infection/open wounds, renal impairment, liver dysfunction, poor nutritional status, other serious co-morbidities)  [If no, no further questions.]  7. Will Neulasta be administered during the period between 14 days before and 24 hours after the administration of cytotoxic chemotherapy?  [If yes, no further questions.]  8. Will Neulasta be used concurrently with radiation therapy, mitomycin C, antimetabolites (e.g., 5-fluorouracil, cytosine arabinoside) or chemotherapeutic agents that have a delayed myelosuppressive effects (e.g., nitrosoureas)?  [No further questions.]  9. Has a recent ANC demonstrated a response to therapy? YN Please document date lab drawn and ANC value:	N	Υ	Does the patient have a documented diagnosis of a nonmyeloid malignancy?	4.
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Comments:	N	Υ	·	9.
			Comments:	C
				_

Date

Prescriber (Or Authorized) Signature