## **Myeloid Growth Factors**



Included Products: Fylnetra (pegfilgrastim-pbbk), Nyvepria (pegfilgrastim-apgf)

10/1/18 Note on product preference: The pharmacy formulary identifies preferred products with an emphasis on biosimilars (with exception for nonformulary Neulasta OnPro). The medical benefit currently allows access equally.

Myeloid growth factors (MGFs) are indicated for the prevention of neutropenic fever – not for the prevention of neutropenia itself. Neutropenia is an expected side effect of many antineoplastic drugs and chemotherapy regimens. MGFs reduce the duration of neutropenia, not the magnitude (known as the nadir). Clinical practice guidelines (NCCN, ESMO, ASCO) adopt the same stance regarding when prophylaxis with MGFs is appropriate using evidence-based recommendations. MGFs are also generally not recommended for the treatment of febrile neutropenia.

- Indicated when the risk of febrile neutropenia is > 20%
- Indicated when the risk of febrile neutropenia is 10-20% plus a risk factor
- Indicated when a patient experienced febrile neutropenia with a previous chemotherapy regimen
- MGF's may still be appropriate in cases where the risk of febrile neutropenia is
   10% but clinicians should be providing justification in these cases

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All Diagnoses					
Initial Criteria		If yes	If no		
1.	Is this request for Leukine?	Continue to #9.	Continue to #2.		
2.	Is this request for <b>prevention</b> of febrile neutropenia in patients undergoing myelosuppressive chemotherapy?	Continue to #10.	Continue to #3.		
3.	Is this request for the <b>treatment</b> of febrile neutropenia? <b>Note:</b> treatment of neutropenia <b>without fever</b> after chemotherapy is not an indication for G-CSF use and should be denied for appropriateness.	Continue to #4.	Continue to #6.		
4.	Is this request for filgrastim or any of its biosimilars?  Note: Pegfilgrastim is a long-acting, one-time injection so it is not appropriate for treating febrile neutropenia.	Continue to #5.	Do not approve. Not medically appropriate.		

5.	Does the patient have any of the following risk factors?  a. Expected hospital stay >10 days b. Profound neutropenia (<100 cells/uL) c. Age>65 d. Pneumonia or other clinically documented infection e. Sepsis syndrome f. Invasive fungal infections g. Prior episode of febrile neutropenia h. Developed fever after being hospitalized (i.e. hospital acquired infection)	Continue to #10.	Do not approve. Not medically necessary.
6.	Does the member have a diagnosis of neutropenia associated with Hepatitis C treatment?	Review for medical necessity.	Continue to #7.
7.	Does the member have acute radiation poisoning?	Continue to #10.	Continue to #8.
8.	Does the member have one of the following diagnoses/ procedures for approval of the medication?  a. Bone marrow transplant (allogenic or autologous).  b. Autologous peripheral blood progenitor cells (PBPC) transplant.  c. Severe chronic neutropenia.  d. AIDS.  e. Myelodysplastic syndromes.	Continue to #10.	Do not approve. Not medically appropriate.
9.	Is Leukine FDA indicated or supported by the NCCN for the requested use?	Continue to #10.	Do not approve.
10.	Is the requested product preferred or formulary on the benefit?	Continue to #12.	Continue to #11.
11.	Is there a reason the preferred product cannot be used on the benefit?	Continue to #12.	Do not approve and offer preferred alternatives.
12.	Approve for 12 months.		

Preferred Products						
Pharmacy Benefit	Brand	Generic				
Formulary	Fylnetra	pegfilgrastim-pbbk				
	Nivestym	filgrastim-aafi				
Non-formulary	Granix	tbo-filgrastim				
	Neupogen	filgrastim				
	Releuko	filgrastim-ayow				
	Zarxio	filgrastim-sndz				
	Fulphila	pegfilgrastim-jmdb				
	Neulasta	pegfilgrastim				
	Neulasta Onpro	pegfilgrastim				
	Nyvepria	pegfilgrastim-apgf				
	Rolvedon	eflapegrastim-xnst				
	Stimufend	pegfilgrastim-fpgk				
	Udenyca	pegfilgrastim-cbqv				
	Ziextenzo	pegfilgrastim-bmez				
	Leukine	sargramostim				