

PHARMACY POLICY STATEMENT

Marketplace

DRUG NAME	Neupogen (Filgrastim)
BILLING CODE	For medical - J1442 For Rx - must use valid NDC
BENEFIT TYPE	Medical or Pharmacy
SITE OF SERVICE ALLOWED	Home/Office/Outpatient Hospital
COVERAGE REQUIREMENTS	Prior Authorization Required (Non-Preferred Product) Alternative preferred product include

Neupogen (Filgrastim) is a non-preferred product and will only be considered for coverage under the medical or pharmacy benefit when the following criteria are met:

Members must be clinically diagnosed with one of the following disease states and meet their individual criteria as stated.

ACUTE MYELOID LEUKEMIA (AML)

For initial authorization:

1. Member has diagnosis of AML documented in chart notes; AND
2. Member must have tried and failed treatment with Zarxio; AND
3. Medication is being used to reduce the time to neutrophil recovery and the duration of fever following induction or consolidation chemotherapy treatment; AND
4. Medication is being administered 24 hours after the last dose of chemotherapy until neutrophil recovery (1 day) or for a maximum of 35 days.
5. Dosage allowed: 1 mg/kg daily or continuous intravenous infusion.

If member meets all the requirements listed above, the medication will be approved for 3 months .

For reauthorization :

1. Member must be in compliance with all initial criteria; AND
2. Chart notes have been provided that show the member is stable or has shown improvement on Neupogen therapy.

If member meets all the reauthorization requirements above, the medication will be approved for an additional 12 months .

AUTOLOGOUS BONE MARROW TRANSPLANT (BMT)

For initial authorization:

1. Member has diagnosis of non-myeloid malignancy and is undergoing myeloablative chemotherapy followed by autologous BMT; AND
2. Member must have tried and failed treatment with Zarxio; AND
3. Medication is being used

If member meets all the requirements listed above, the medication will be approved for 3 months .

For reauthorization :

1. Member must be in compliance with all initial criteria; AND
2. Chart notes have been provided that show the member is stable or has shown improvement on Neupogen therapy.

If member meets all the reauthorization requirements above, the medication will be approved for an additional 12 months.

AUTOLOGOUS PERIPHERAL BLOOD PROGENITOR CELL (PBPC) MOBILIZATION

For initial authorization:

1. Medication is being used to mobilize autologous peripheral blood progenitor cells for collection by leukapheresis; AND
2. Member must have tried and failed treatment with Zarxio; AND
3. Medication is being administered for at least 4 days before first leukapheresis and continued until the last leukapheresis (X Q W L O D V X V W D € '7L srl the

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Appendix

Chemotherapy Regimens with a High Risk for Febrile Neutropenia (>

Cancer Type	Regimen
Acute Lymphoblastic Leukemia (ALL)	ALL induction regimens (see NCCN guidelines)
Bladder Cancer	0 9 \$ & P H W K R W U G E D M W L Q H G R [R U X E L F L Q F L V (metastatic)
Breast Cancer	Docetaxel + trastuzumab (metastatic or relapsed)
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	7 \$ & G R F H W D [H O c y c l o p h o s p h a m i d e (adjuvant)
Esophageal and Gastric Cancers	Docetaxel/cisplatin/fluorouracil
Hodgkin Lymphoma	% (\$ & 2 3 3 E O H R P \ F L Q H W R S R V L G H G R [R U X E L F L SURFDUED]LQH SUHGQLVRQH
Kidney Cancer	Doxorubicin/gemcitabine
Non-Hodgkin's Lymphoma	, & (L I R V I D P L G H F D U E R S O D W L Q - F H W O R S O V L S K I R P peripheral T- F H O O O \ P S K R P D V > 3 7 & / @ Q G O L Q H
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National Comprehensive Cancer Network (NCCN): Myeloid Growth Factors, 2016.

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Cancer Type	Regimen
Occult Primary Adenocarcinoma	[REDACTED]

	&LVSODWLQ HWRSRVLGH DGMXYDQW DGYDQFH
	&DUERSODWLQ SDFOLWD[HO DGMXYDQW DGYD
	Docetaxel (advanced/metastatic)

