



Colony Stimulating Factors:

Filgrastim (Neupogen®); Filgrastim-aafi (Nivestym™); Filgrastim-sndz (Zarxio®); Filgrastim-ayow (Releuko®); Tbo-Filgrastim (Granix®) (Subcutaneous/Intravenous)

Document Number: IC-0235

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I. Length of Authorization

Coverage will be provided for 4 months and may be renewed.

II. Dosing Limits

A. Quantity Limit (max daily dose) [NDC Unit]:

- Neupogen 300 mcg single-dose vial: 3 vials per 1 day
- Neupogen 300 mcg single-dose prefilled syringe (Single-Ject): 3 syringes per 1 day
- Neupogen 480 mcg single-dose vial: 3 vials per 1 day
- Neupogen 480 mcg single-dose prefilled syringe (Single-Ject): 3 syringes per 1 day
- Nivestym 300 mcg single-dose vial: 3 vials per 1 day
- Nivestym 300 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Nivestym 480 mcg single-dose vial: 3 vials per 1 day
- Nivestym 480 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Zarxio 300 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Zarxio 480 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Releuko 300 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Releuko 480 mcg single-dose prefilled syringe: 3 syringes per 1 day
- Releuko 300 mcg single-dose vial: 3 vials per 1 day
- Releuko 480 mcg single-dose vial: 3 vials per day
- Granix 300 mcg single-dose pre-filled syringe: 4 syringes per 1 day
- Granix 300 mcg single-dose vial: 4 vials per 1 day
- Granix 480 mcg single-dose pre-filled syringe: 3 syringes per 1 day
- Granix 480 mcg single-dose vial: 3 vials per 1 day



B. Max Units (per dose and over time) [HCPCS Unit]:

Severe Chronic Neutropenia (Congenital Neutropenia):

• 1380 billable units per day

BMT or PBPC or H-ARS:

• 1200 billable units per day

All other indications:

• 600 billable units per day

III. Initial Approval Criteria

Coverage is provided in the following conditions:

Bone Marrow Transplant (BMT) † ‡ Φ 1-6

Peripheral Blood Progenitor Cell (PBPC) mobilization and transplant † ‡ Ф 1-6,19,30,33,35-37

Prophylactic use in patients with solid tumors or non-myeloid malignancy $\dagger \ddagger {}^{1\text{-}7,9,10,12,13,15,17,27\text{-}29}$

- Patient is undergoing myelosuppressive chemotherapy with an expected incidence of febrile neutropenia of greater than 20% §; OR
- Patient is undergoing myelosuppressive chemotherapy with an expected incidence of febrile neutropenia of 10% to 20% **§ AND** one or more of the following co-morbidities:
 - Age >65 years receiving full dose intensity chemotherapy
 - Extensive prior exposure to chemotherapy
 - Previous exposure of pelvis, or other areas of large amounts of bone marrow, to radiation
 - Pre-existing neutropenia (ANC ≤ 1000/mm³)
 - Bone marrow involvement with tumor
 - Patient has a condition that can potentially increase the risk of serious infection (i.e., HIV/AIDS with low CD4 counts)
 - Recent surgery and/or open wounds
 - Poor performance status
 - Renal dysfunction (creatinine clearance <50 mL/min)
 - Liver dysfunction (elevated bilirubin >2.0 mg/dL)
 - Chronic immunosuppression in the post-transplant setting including organ transplant Note: Dose-dense therapy, in general, requires growth factor support to maintain dose intensity and schedule. In the palliative setting, consideration should be given to dose reduction or change in regimen.

Treatment of chemotherapy-induced febrile neutropenia ‡ 1-7,9,10,12,13,15,17,27-29

• Patient has been on prophylactic therapy with filgrastim or tho-filgrastim (*Note: therapy should not be used concomitantly with pegfilgrastim*); **OR**



- Patient has not received prophylactic therapy with a granulocyte colony stimulating factor;
 AND
 - Patient has one or more of the following risk factors for developing infection-related complications:
 - Sepsis Syndrome
 - Age greater than 65 years
 - Absolute neutrophil count [ANC] less than 100/mcL
 - Duration of neutropenia expected to be greater than 10 days
 - Pneumonia or other clinically documented infections
 - Invasive fungal infection
 - Hospitalization at the time of fever
 - Prior episode of febrile neutropenia

Patient who experienced a neutropenic complication from a prior cycle of the same chemotherapy \$\pmu\$ 1-7,9,10,12,13,15,17,27-29

<u>Note</u>: Dose-dense therapy, in general, requires growth factor support to maintain dose intensity and schedule. In the palliative setting, consideration should be given to dose reduction or change in regimen.

Acute Myeloid Leukemia (AML) † ‡ Φ 1-6,8,14,35

- Used in patients receiving induction/consolidation or re-induction chemotherapy; **OR**
- Used for relapsed or refractory disease

Bone Marrow Transplantation (BMT) failure or Engraftment Delay † ‡ 1-7,25,26,30,33,35-37

Severe Chronic Neutropenia † ‡ Φ 1-6,11

- Patient must have an absolute neutrophil count (ANC) < 500/mm³; AND
- Patient must have a diagnosis of one of the following:
 - o Congenital neutropenia; **OR**
 - o Cyclic neutropenia; **OR**
 - o Idiopathic neutropenia

Myelodysplastic Syndrome ‡ 6

- Patient has symptomatic anemia with no del(5q) mutation; AND
- Patient has lower risk disease (i.e., defined as IPSS-R [Very Low, Low, Intermediate]); AND
- Endogenous serum erythropoietin level of ≤500 mUnits/mL; AND
 - Patient has ring sideroblasts ≥15% (or ring sideroblasts ≥5% with an SF3B1 mutation); AND
 - Used in combination with an Erythropoiesis Stimulating Agent (ESA); OR



- Patient has ring sideroblasts <15% (or ring sideroblasts <5% with an SF3B1 mutation); AND
 - Used in combination with an ESA following no response (despite adequate iron stores) or erythroid response followed by loss of response to ESA alone

Patients acutely exposed to myelosuppressive doses of radiation (Hematopoietic Subsyndrome of Acute Radiation Syndrome [H-ARS]) $\dagger \ddagger \Phi$ ^{1-6,18}

Management of CAR T-cell related Toxicity ‡ 6

- Patient has been receiving therapy with CAR T-cell therapy (e.g., axicabtagene ciloleucel, brexucabtagene autoleucel, ciltacabtagene autoleucel, idecabtagene vicleucel, lisocabtagene maraleucel, tisagenlecleucel, etc.); AND
- Patient is experiencing neutropenia related to their therapy

Wilms Tumor (Nephroblastoma) ‡ 6

- Patient has favorable histology disease; AND
- Used in combination with a cyclophosphamide-based chemotherapy regimen (i.e., Regimen M or I only)
- † FDA-labeled indication(s); ‡ Compendia recommended indication(s); Φ Orphan Drug

§Febrile neutropenia is defined as: ⁷

- Temperature: a single temperature ≥38.3 °C orally or ≥38.0 °C over 1 hour; AND
- Neutropenia: <500 neutrophils/mcL or <1,000 neutrophils/mcL and a predicted decline to ≤500 neutrophils/mcL over the next 48 hours

§ Expected incidence of febrile neutropenia percentages for myelosuppressive chemotherapy regimens can be found in the NCCN Hematopoietic Growth Factors Clinical Practice Guideline at NCCN.org ⁷

IV. Renewal Criteria

Coverage may be renewed based upon the following criteria:

- Patient continues to meet indication-specific relevant criteria such as concomitant therapy requirements (not including prerequisite therapy), performance status, etc. identified in section III; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: splenic rupture, acute respiratory distress syndrome (ARDS), serious allergic reactions/anaphylaxis, sickle cell crisis, glomerulonephritis, leukocytosis, capillary leak syndrome, potential for tumor growth stimulation of malignant cells, aortitis, alveolar hemorrhage and hemoptysis, thrombocytopenia, cutaneous vasculitis, MDS/AML (when used for congenital neutropenia), etc.

V. Dosage/Administration 1-5

| indication | | Dose | | |
|------------|-----------------------|----------------------|---------------------|--|
| | FILGRASTIM (Neupogen® | . Nivestvm™. Zarxio¹ | ™. Releuko® and | |



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| BMT/PBPC/H-ARS | 10 mcg/kg daily for up to 14 days |
|------------------------|-----------------------------------|
| Congenital Neutropenia | 6 mcg/kg twice daily |
| All other indications | 5 mcg/kg daily for up to 14 days |

VI. Billing Code/Availability Information

HCPCS Code:

- J1442 Injection, filgrastim (g-csf), excludes biosimilars, 1 mcg: 1 billable unit = 1 mcg
- Q5110 Injection, filgrastim-aafi, biosimilar, (Nivestym), 1 mcg: 1 billable unit = 1 mcg
- Q5101 Injection, filgrastim-sndz, biosimilar, (Zarxio), 1 mcg: 1 billable unit = 1 mcg
- J1447 Injection, tbo-filgrastim (Granix), 1 mcg: 1 billable unit = 1 mcg
- Q5125 Injection, filgrastim-ayow, biosimilar, (Releuko), 1 mcg; 1 billable unit = 1 mcg

NDC:

- Neupogen 300 mcg single-dose vial: 55513-0530-xx
- Neupogen 300 mcg single-dose prefilled syringe (SingleJect): 55513-0924-xx
- Neupogen 480 mcg single-dose vial: 55513-0546-xx
- Neupogen 480 mcg single-dose prefilled syringe (SingleJect): 55513-0209-xx
- Nivestym 300 mcg single-dose vial: 00069-0293-xx
- Nivestym 300 mcg single-dose prefilled syringe: 00069-0291-xx
- Nivestym 480 mcg single-dose vial: 00069-0294-xx
- Nivestym 480 mcg single-dose prefilled syringe: 00069-0292-xx
- Zarxio 300 mcg single-dose prefilled syringe: 61314-0318-xx
- Zarxio 480 mcg single-dose prefilled syringe: 61314-0326-xx
- Releuko 300 mcg single-dose prefilled syringe: 70121-1568-xx
- Releuko 480 mcg single-dose prefilled syringe: 70121-1570-xx
- Releuko 300 mcg single-dose vial: 70121-1569-xx
- Releuko 480 mcg single-dose vial: 70121-1571-xx
- Granix 300 mcg single-dose prefilled syringe: 63459-0910-xx
- Granix 480 mcg single-dose prefilled syringe: 63459-0912-xx
- Granix 300 mcg single-dose vial: 63459-0918-xx
- Granix 480 mcg single-dose vial: 63459-0920-xx

VII. References

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- 7. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) Hematopoietic Growth Factors. Version 2.2023. National Comprehensive Cancer Network, 2023. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed March 2023.
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Appendix 1 – Covered Diagnosis Codes

| ICD-10 | ICD-10 Description | |
|---|--|--|
| C64.1 | Malignant neoplasm of right kidney, except renal pelvis | |
| C64.2 | Malignant neoplasm of left kidney, except renal pelvis | |
| C64.9 | Malignant neoplasm of unspecified kidney, except renal pelvis | |
| C92.00 | Myeloid leukemia not having achieved remission | |
| C92.02 | Myeloid leukemia in relapse | |
| C92.50 | Acute myelomonocytic leukemia not having achieved remission | |
| C92.52 | Acute myelomonocytic leukemia in relapse | |
| C92.60 | Acute myeloid leukemia with 11q23-abnormality not having achieved remission | |
| C92.62 | Acute myeloid leukemia with 11q23-abnormality in relapse | |
| C92.A0 | Acute myeloid leukemia with multilineage dysplasia not having achieved remission | |
| C92.A2 | Acute myeloid leukemia with multilineage dysplasia in relapse | |
| C93.00 | Acute monoblastic/monocytic leukemia not having achieved remission | |
| C93.02 | Acute monoblastic/monocytic leukemia in relapse | |
| C93.10 Chronic myelomonocytic leukemia, not having achieved remission | | |
| D46.0 Refractory anemia without ring sideroblasts, so stated | | |
| D46.1 | Refractory anemia with ring sideroblasts | |
| D46.20 | Refractory anemia with excess of blasts, unspecified | |
| D46.21 | Refractory anemia with excess of blasts 1 | |
| D46.4 | Refractory anemia, unspecified | |
| D46.9 | Myelodysplastic syndrome, unspecified | |
| D46.A | Refractory cytopenia with multilineage dysplasia | |
| D46.B | Refractory cytopenia with multilineage dysplasia and ring sideroblasts | |
| D46.Z | Other myelodysplastic syndrome | |
| D61.81 | Pancytopenia | |
| D70.0 | Congenital agranulocytosis | |
| D70.1 | Agranulocytosis secondary to cancer chemotherapy | |
| D70.2 | 0.2 Other drug-induced agranulocytosis | |
| D70.4 | 0.4 Cyclic neutropenia | |
| D70.9 | Neutropenia, unspecified | |



| ICD-10 | ICD-10 Description | |
|----------|---|--|
| T45.1X5A | Adverse effect of antineoplastic and immunosuppressive drugs initial encounter | |
| T45.1X5D | Adverse effect of antineoplastic and immunosuppressive drugs subsequent encounter | |
| T45.1X5S | Adverse effect of antineoplastic and immunosuppressive drugs sequela | |
| T66.XXXA | Radiation sickness, unspecified, initial encounter | |
| T66.XXXD | Radiation sickness, unspecified, subsequent encounter | |
| T66.XXXS | Radiation sickness, unspecified, sequela | |
| W88.1 | Exposure to radioactive isotopes | |
| W88.8 | Exposure to other ionizing radiation | |
| Z41.8 | Encounter for other procedures for purposes other than remedying health state | |
| Z48.290 | Encounter for aftercare following bone marrow transplant | |
| Z51.11 | Encounter for antineoplastic chemotherapy | |
| Z51.12 | Encounter for antineoplastic immunotherapy | |
| Z51.89 | Encounter for other specified aftercare | |
| Z52.001 | Unspecified donor, stem cells | |
| Z52.011 | Autologous donor, stem cells | |
| Z52.091 | Other blood donor, stem cells | |
| Z76.89 | Persons encountering health services in other specified circumstances | |
| Z94.81 | Bone marrow transplant status | |
| Z94.84 | Stem cells transplant status | |

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD), Local Coverage Determinations (LCDs) and Local Coverage Articles (LCAs) may exist and compliance with these policies is required where applicable. They can be found at: https://www.cms.gov/Medicare-Coverage-Database/search.aspx. Additional indications may be covered at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCD/LCA):

| Jurisdiction(s): J,M | NCD/LCD Document (s): A56748 | |
|--|------------------------------|--|
| https://www.cms.gov/medicare-coverage-database/new-search/search- | | |
| results.aspx?keyword=a56748&areaId=all&docType=NCA%2CCAL%2CNCD%2CMEDCAC%2CTA%2CM | | |
| <u>CD%2C6%2C3%2C5%2C1%2CF%2CP</u> | | |

| Jurisdiction(s): N (9) | NCD/LCD Document (s): A57789 |
|------------------------|------------------------------|
|------------------------|------------------------------|



 $\frac{https://www.cms.gov/medicare-coverage-database/new-search/search-results.aspx?keyword=a57789\&areaId=all\&docType=NCA%2CCAL%2CNCD%2CMEDCAC%2CTA%2CMCD%2C6%2C3%2C5%2C1%2CF%2CP$

| | Medicare Part B Administrative Contractor (MAC) Jurisdictions | | |
|--------------|---|---|--|
| Jurisdiction | Applicable State/US Territory | Contractor | |
| E (1) | CA, HI, NV, AS, GU, CNMI | Noridian Healthcare Solutions, LLC | |
| F (2 & 3) | AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ | Noridian Healthcare Solutions, LLC | |
| 5 | KS, NE, IA, MO | Wisconsin Physicians Service Insurance Corp (WPS) | |
| 6 | MN, WI, IL | National Government Services, Inc. (NGS) | |
| H (4 & 7) | LA, AR, MS, TX, OK, CO, NM | Novitas Solutions, Inc. | |
| 8 | MI, IN | Wisconsin Physicians Service Insurance Corp (WPS) | |
| N (9) | FL, PR, VI | First Coast Service Options, Inc. | |
| J (10) | TN, GA, AL | Palmetto GBA, LLC | |
| M (11) | NC, SC, WV, VA (excluding below) | Palmetto GBA, LLC | |
| L (12) | DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA) | Novitas Solutions, Inc. | |
| K (13 & 14) | NY, CT, MA, RI, VT, ME, NH | National Government Services, Inc. (NGS) | |
| 15 | KY, OH | CGS Administrators, LLC | |



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- Qualified sign language interpreters.
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- Qualified interpreters.
- Information written in other languages.

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Nondiscrimination Grievance Coordinator

Aspirus Health Plan, Inc.

PO Box 1890

Southampton, PA 18966-9998

Phone: 1-866-631-5404 (TTY: 711)

Fax: 763-847-4010

Email: customerservice@aspirushealthplan.com

You can file a *grievance* in person or by mail, fax, or email. If you need help filing a *grievance*, the Nondiscrimination Grievance Coordinator is available to help you.

You can also file a civil rights complaint with the U.S. Department of Health and Human Services, Office for Civil Rights, electronically through the Office for Civil Rights Complaint Portal, available at https://ocrportal.hhs.gov/ocr/portal/lobby.jsf, or by mail or phone at:

U.S. Department of Health and Human Services

200 Independence Avenue, SW

Room 509F, HHH Building

Washington, D.C. 20201

1.800.368.1019, 800.537.7697 (TDD)

Complaint forms are available at http://www.hhs.gov/ocr/office/file/index.html. This notice is available at Aspirus Health Plan, Inc.'s website: https://aspirushealthplan.com/webdocs/70021-AHP-NonDiscrim_Lang-Assist-Notice.pdf.

Language Assistance Services

Albanian: KUJDES: Nëse flitni shqip, për ju ka në dispozicion shërbime të asistencës gjuhësore, pa pagesë. Telefononi në 1-800-332-6501 (TTY: 711). (مراكبة العربية، فإن خدمات المساعدة اللغوية متاحة لك مجاناً الصم والبك ، Arabic

French: ATTENTION: Si vous parlez français, des services d'aide linguistique vous sont proposés gratuitement. Appelez le 1-800-332-6501 (ATS: 711).

German: ACHTUNG: Wenn Sie Deutsch sprechen, stehen Ihnen kostenlos sprachliche Hilfsdienstleistungen zur Verfügung. Rufnummer: 1-800-332-6501 (TTY: 711).

Hindi: यान द: य द आप िहंदी बोलते ह तो आपके िलए म्. त.म. भाषा सहायता सेवाएं उपल ध ह। 1-800-332-6501 (TTY: 711) पर कॉल कर।

Hmong: LUS CEEV: Yog tias koj hais lus Hmoob, cov kev pab txog lus, muaj kev pab dawb rau koj. Hu rau 1-800-332-6501 (TTY: 711).

Korean: 주의: 한국어를 사용하시는 경우, 언어 지원 서비스를 무료로 이용하실 수 있습니다.1-800-332-6501 (TTY: 711)번으로 전화해 주십시오.

Polish: UWAGA: Jeżeli mówisz po polsku, możesz skorzystać z bezpłatnej pomocy językowej. Zadzwoń pod numer1-800-332-6501 (TTY: 711).

Russian: ВНИМАНИЕ: Если вы говорите на русском языке, то вам доступны бесплатные услуги перевода. Звоните 1-800-332-6501 (телетайп: 711)

Spanish: ATENCIÓN: si habla español, tiene a su disposición servicios gratuitos de asistencia lingüística. Llame al1-800-332-6501 (TTY: 711).

Tagalog: PAUNAWA: Kung nagsasalita ka ng Tagalog, maaari kang gumamit ng mga serbisyo ng tulong sa wika nangwalang bayad. Tumawag sa 1-800-332-6501 (TTY: 711).

Traditional Chinese: 注意: 如果您使用繁體中文, 您可以免費獲得語言援助服務。請 致電 1-800-332-6501 (TTY: 711)

Vietnamese: CHÚ Ý: Nếu bạn nói Tiếng Việt, có các dịch vụ hỗ trợ ngôn ngữ miễn phí dành cho bạn. Gọi số 1-800-332-6501 (TTY: 711).

Pennsylvania Dutch: Wann du Deitsch (Pennsylvania German / Dutch) schwetzscht, kannscht du mitaus Koschte ebbergricke, ass dihr helft mit die englisch Schprooch. Ruf selli Nummer uff: Call 1-800-332-6501 (TTY: 711).

Lao: ໂປດຊາບ: ຖ້າວ່າ ທ່ານເວົ້າພາສາ ລາວ, ການບໍລິການຊ່ວຍເຫຼືອດ້ານພາສາ,ໂດຍບໍ່ເສັຽຄ່າ, ແມ່ນມີພ້ອມໃຫ້ທ່ານ. ໂທຣ 1-800-332-6501 (TTY: 711).