



**HEALTH PARTNER PLANS  
PRIOR AUTHORIZATION REQUEST FORM**

Neulasta

Phone: 215-991-4300

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

**PLEASE NOTE: Any information (patient, prescriber, drugs, labs) left blank, illegible or not attached WILL delay the review process.**

<p><b>Patient Name:</b> HPP Member Number: Date of Birth: Address: City, State ZIP: Patient Primary Phone: Line of Business: <input type="checkbox"/> Medicaid <input type="checkbox"/> CHIP</p>	<p><b>Prescriber Name:</b> Fax: _____ Phone: _____ Office Contact: NPI: _____ Promise ID: _____ <i>Prescriber PA PROMISe ID:</i> Address: City, State ZIP: <b>Specialty/facility name (if applicable):</b></p>
--	--

Expedited/Urgent

**Drug name:**  
**Strength:**  
**Days Supply:**  
**Number of Refills:**  
**Directions / SIG:**

*HPP's maximum approval time is 12 months but may be less depending on the drug.*

<p><b>Please attach any pertinent medical history including labs and information for this member that may support approval.</b></p> <p><b>Please answer the following questions and sign.</b></p>
<p>Q1. Primary Prophylaxis of Febrile Neutropenia: Is patient receiving myelosuppressive chemotherapy? Include chemotherapy regimen. <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>
<p>Q2. Primary Prophylaxis of Febrile Neutropenia: Is patient at an approximately 20% or higher risk for febrile neutropenia? <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>
<p>Q3. Primary Prophylaxis of Febrile Neutropenia: Is patient at an approximately 10% or higher for febrile neutropenia AND has ONE of the following risk factors (such as age &gt;65 years, renal dysfunction, liver dysfunction, recent surgery and/or open wounds, bone marrow involvement by tumor, persistent neutropenia, prior radiation therapy)? Provide supporting documentation? <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>
<p>Q4. Primary Prophylaxis of Febrile Neutropenia: Is patient receiving dose-dense or high-dose chemotherapy? <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>
<p>Q5. Secondary Prophylaxis of Febrile Neutropenia: Is patient receiving myelosuppressive chemotherapy with a history of Febrile Neutropenia during previous course of chemotherapy (for which primary prophylaxis was not received)? Please include treatment plan. <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>
<p>Q6. Secondary Prophylaxis of Febrile Neutropenia: Have a CBC (complete blood count with differential including ANC been obtained? (please attached the copy of the lab result). <input type="checkbox"/> Yes                      <input type="checkbox"/> No</p>

This telecopy transmission confidential information belonging to the sender that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or action taken in reference to the contents of this document is strictly prohibited. If you have receive this telecopy in error, please notify the sender to arrange for the return of this document.



HEALTH PARTNER PLANS  
PRIOR AUTHORIZATION REQUEST FORM

Neulasta

Phone: 215-991-4300

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

**PLEASE NOTE: Any information (patient, prescriber, drugs, labs) left blank, illegible or not attached WILL delay the review process.**

Patient Name:

Prescriber Name:

Q7. Hematopoietic Subsyndrome of Acute Radiation Syndrome: Has the patient acutely exposed to myelosuppressive doses of radiation?

Yes                       No

Q8. Hematopoietic Subsyndrome of Acute Radiation Syndrome: Have a CBC (complete blood count with differential including ANC)) and platelet count had been obtained? (please attach a copy of the lab result).

Yes                       No

Q9. Requested Duration:

3 months

Q10. Additional Information:

\_\_\_\_\_  
Prescriber Signature

\_\_\_\_\_  
Date

Updated 2018