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Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

Nivestym™ Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Acute myeloid leukemia (AML) following induction or consolidation chemotherapy <input type="checkbox"/> Bone marrow transplant (BMT)/stem cell transplant <input type="checkbox"/> HIV-related neutropenia <input type="checkbox"/> Prophylaxis of febrile neutropenia <input type="checkbox"/> Severe chronic neutropenia (SCN) <input type="checkbox"/> Treatment of febrile neutropenia <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical Information: Select if Nivestym is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Hematologist/oncologist <input type="checkbox"/> Infectious disease specialist Please specify the duration of therapy: _____					
For bone marrow transplant (BMT)/stem cell transplant, also answer the following: Select the procedure for which Nivestym is being used: <input type="checkbox"/> For patients with non-myeloid malignancies undergoing myeloablative chemotherapy followed by autologous or allogeneic BMT <input type="checkbox"/> For mobilization of hematopoietic progenitor cells into the peripheral blood for collection by leukapheresis <input type="checkbox"/> For peripheral stem cell transplant (PSCT) patients who have received myeloablative chemotherapy					
For HIV-related neutropenia, also answer the following: Is the absolute neutrophil count (ANC) \leq 1,000 cells/mm ³ ? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For prophylaxis of febrile neutropenia, also answer the following: Select if Nivestym will be used for prophylaxis of febrile neutropenia (FN) due to the following: <input type="checkbox"/> Patient is receiving National Cancer Institute's Breast Intergroup, INT C9741 dose dense chemotherapy protocol for primary breast cancer (doxorubicin, cyclophosphamide, and paclitaxel) <input type="checkbox"/> Patient is receiving a dose-dense chemotherapy regimen for which the incidence of FN is unknown <input type="checkbox"/> Patient is receiving a chemotherapy regimen associated with > 20% incidence of FN <input type="checkbox"/> Patient is receiving a chemotherapy regimen associated with 10-20% incidence of FN <input type="checkbox"/> Patient has one or more risk factors associated with chemotherapy-induced infection, FN, or neutropenia <input type="checkbox"/> Patient is receiving myelosuppressive anti-cancer drugs associated with neutropenia <input type="checkbox"/> Patient has a history of FN or dose-limiting event during a previous course of chemotherapy (secondary prophylaxis)					

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Nivestym™ Prior Authorization Request Form (Page 2 of 2)

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For severe chronic neutropenia, also answer the following:

Does the patient have severe chronic neutropenia (i.e., congenital, cyclic, and idiopathic neutropenias with chronic ANC ≤ 500 cells/mm³)? Yes No

For treatment of febrile neutropenia, also answer the following:

Has the patient received or is receiving myelosuppressive anticancer drugs associated with neutropenia? Yes No

Does the patient have febrile neutropenia and is at high risk for infection-associated complications? Yes No

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received.

For urgent or expedited requests please call 1-800-711-4555.

This form may be used for non-urgent requests and faxed to 1-800-527-0531.

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