

Medicare Part B Prior Authorization/Step Therapy Form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

This form applies to:	Medicare Part B	ing) 🗆 Non Urge	ant (atandard ravious)		
This request is:	Urgent (life threaten Urgent means the standard rev to regain maximum function.	o, —	,		ity
Neupoger	n® (filgrastim)			
Member Information	n				_
Last Name:		First Nar	me:		
				Gender:	
	n:				
Provider Informatio	n				
Requesting Provider: _		Phone:		Fax:	
Address:					
			Name:		
Provider Signature:		Date:	Date:		
Patient Dosing Info Date of last dose (if a	Neupogen 300 mcg/0.5mL [Neupogen 480 mcg/0.8mL [rmation: oplicable): upplicable)	Neupogen 480 m Total do Height:	eses/cycles/duration Weight:	requested: BSA:	
☐ Home infusion Ager		NPI:	Fax		-
Billing: Physician to buy an					
Facility to buy and b					
_ ,	/: 	NPI:	Fax:_		
ICD-10 Diagnosis Cod	de(s):	HCPCS	Code:		



Precertification Requirements

Step therapy (trial with the below listed drug[s]) is only applicable to members who are enrolled in a Medicare Advantage Prescription Drug (MAPD) plan and will not apply to members who are actively receiving treatment with the non-preferred drug (have a paid drug claim within the past 365 days).

Before this drug is covered, the patient must meet the following:

- 1. Must be used for a medically accepted indication and follow applicable NCD, LCD and/or LCA requirements².
- 2. Must first try Nivestym and Zarxio.

¹See Medically accepted indication section below

Additional information

When criteria are met, coverage duration is for 2 years. Dose will be approved according to the FDA-approved labeling or within accepted standards of medical practice.

Medically accepted indication¹

If no NCD, LCD, or LCA criteria² are available for the state in which the member is receiving services, Medicare Part B drugs will be reviewed for a medically accepted indication, defined in the Medicare Benefit Policy Manual Chapter 15 § 50:

A medically accepted indication for a drug that is not a part of an anti-cancer regimen is a use that is:

- approved by the Food and Drug Administration. (That is, the Food and Drug Administration has approved the drug for the diagnosis or condition for which it is being prescribed.)
- or supported by certain references, taking into consideration the major drug compendia (e.g. American Hospital Formulary Service-Drug Information, Micromedex DrugDex, Lexi-Drugs), authoritative medical literature, and/or accepted standards of medical practice.

National and Local Coverage Determination/Article (NCD, LCD, and LCA) Criteria²

Priority Health applies Medicare NCD, LCD, and LCA criteria for Part B drugs. The following apply to Neupogen: N/A

Precertification Documentation			
A. \	What condition is this drug being requested for? ☐ Febrile neutropenia prevention 1. Does the patient have a non-myeloid cancer and is receiving myelosuppressive anti-cancer drugs? ☐ Yes. ☐ No. Are you asking for an exception to this requirement? ☐ Yes. Rationale for exception:		
	Other:		

²See NCD, LCD, and LCA section below



B.	Has the patient tried Nivestym? Yes. No. Are you asking for an exception to this requirement? Yes. Rationale for exception: No
C.	Has the patient tried Zarxio? Yes. No. Are you asking for an exception to this requirement? Yes. Rationale for exception:
<u></u>	is with the olds Madisons Everetion Democrat (everetions to the above evitoris)
Do If y	iority Health Medicare Exception Request (exceptions to the above criteria) you believe one or more of the step therapy requirements should be waived? Yes No es, you must provide a statement explaining the medical reason why the exception should be approved.
	No Yes, because:
	he patient is currently using Neupogen, would changing the patient's current regimen likely result in adverse ects for the patient? No Yes, because: