# Nemluvio (nemolizumab) **Prior Authorization Request Form**

Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

#### MEMBER'S LAST NAME: \_\_\_\_\_ MEMBER'S FIRST NAME: \_\_\_\_\_

Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

MEMBER INFORMATION	
LAST NAME:	FIRST NAME:
PHONE NUMBER:	DATE OF BIRTH:
STREET ADDRESS:	
CITY:	STATE: ZIP CODE:
PATIENT INSURANCE ID NUMBER:	

MALE FEMALE HEIGHT (IN/CM): \_\_\_\_\_ WEIGHT (LB/KG): \_\_\_\_ ALLERGIES: \_\_\_\_\_

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: PRIMETHERAPEUTICS.COM/NOPP

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): AUTHORIZED REPRESENTATIVE'S PHONE NUMBER:

PRESCRIBER INFORMATION	
LAST NAME:	FIRST NAME:
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:
NPI NUMBER:	DEA NUMBER:
PHONE NUMBER:	FAX NUMBER:
STREET ADDRESS:	
CITY:	STATE: ZIP CODE:
REQUESTER (if different than prescriber):	OFFICE CONTACT PERSON:

MEDICATION	DICDENCING INFODM	
WEDICATION	DISPENSING INFORM	

MEDICATION NAME:				
DOSE/STRENGTH:	FREQUENCY:		LENGTH OF	QUANTITY:
			THERAPY/REFILLS:	
NEW THERAPY	RENEWAL	IF REI	NEWAL: DATE THERAPY I	NITIATED:
DURATION OF THERAPY	(SPECIFIC DATES)	):		
Continued on next page				

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MEMBER'S LAST NAME:	MEMBER'S FIRST N	IAME:
1. HAS THE PATIENT TRIED ANY	OTHER MEDICATIONS FOR THIS	CONDITION?
<b>YES</b> (if yes, complete below)	NO	
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	<b>DURATION OF THERAPY</b> (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:
2. LIST DIAGNOSES:		ICD-10:
<ul> <li>Moderate to severe atopic derma</li> <li>Prurigo Nodulatis</li> <li>Other diagnosis:</li> </ul>	atitis ICD-10 Code(s):	
3. REQUIRED CLINICAL INFORMATO SUPPORT A PRIOR AUTHORIZ	ATION: PLEASE PROVIDE ALL REL ZATION.	EVANT CLINICAL INFORMATION
Will the patient be using the drug	as a part of the clinical trial? 🗌 Ye	es 🗌 No
following: ?  Yes No Allergist Immunologist Dermatologist	d by one of the following or in con	
TNF inhibitors, JAK inhibitors, IL-	used in combination with another i -4 inhibitors)?	mmunomodulatory agent (e.g.,
	used in combo w Cibinqo (abrociti ra (ruxolitinib), or Adbry (tralokinur	
Has the patient tried and failed a 3	3-month trial of Dupixent (Please s	ubmit documentation)?
For diagnosis of Atopic Dermatiti Has the patient had the diagnosis Please submit documentation.	<u>s</u> , answer the following: of atopic dermatitis for at least 12	? months? 🗌 Yes 🗌 No
Does the patient have atopic dern	natitis on at least 10% or more of t umentation.	heir body surface area?
Please submit documentation.	different topical steroids? 🗌 Yes	
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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:
-	lifferent topical steroids, has the patient tried at least one topical urin inhibitor (tacrolimus or pimecrolimus)?
	lifferent topical steroids, has the patient tried at least one topical )?
-	lifferent topical steroids, has the patient tried at least one topical ?
-	lifferent topical steroids, has the patient tried at least one topical
For a diagnosis of Prurigo Nodu	laris, please answer the following:
Has patient had chronic pruritus	lasting ≥6 weeks?
Does patient have history and/or and scars)?	r signs of repeated scratching, picking, or rubbing (eg, excoriations se submit documentation.
Does patient have presence of m Please submit documentation.	nultiple pruriginous lesions, including firm nodules? 🗌 Yes 🗌 No
Has the patient tried at least 2 di Please submit documentation.	fferent medium-to-super-potent topical steroids? 🗌 Yes 🗌 No
-	e medium-to-super-potent topical steroid AND one topical calcineurin limus)?
· · ·	en in combination with phototherapy AND at least one medium-to- Yes 🗌 No Please submit documentation.
Has patient tried excimer laser A No Please submit documentation	ND at least one medium-to-super-potent topical steroid?
Has patient tried cryotherapy AN documentation.	ID intralesional steroids? 🗌 Yes 🗌 No Please submit



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MEMBER'S LAST NAME: MEMBER'S FIRST NAME:
Has patient tried at least one oral DMARD such as methotrexate or cyclosporine?  Yes No Please submit documentation.
Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?
Renewal Request: Does the patient continue to have positive clinical response with continued use of Nemluvio (nemolizumab)?  Yes  No Please submit documentation.
Is the medication being prescribed by one of the following or in consultation with one of the following: ?
Will Nemluvio (nemolizumab) be used in combo w Cibinqo (abrocitinib), Olumiant (baracitinib), RinvoqER (upadacitinib), Opzelura (ruxolitinib), or Adbry (tralokinumab), Dupixent (dupilumab) or Ebglyss (lebrikizumab)?  Yes No
Will Nemluvio (nemolizumab) be used in combination with another immunomodulatory agent (e.g., TNF inhibitors, JAK inhibitors, IL-4 inhibitors)?
<b>Please note:</b> Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.
<b>ATTESTATION:</b> I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.
Prescriber Signature or Electronic I.D. Verification: Date:
<b>CONFIDENTIALITY NOTICE:</b> The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX) and arrange for the return or destruction of these documents.
FAX THIS FORM TO: 800-424-7640 MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program
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Attn: CP-4201 P.O. Box 64811 St. Paul, MN 55164-0811 Phone: 877-228-7909

