

**Zoryve Topicals (roflumilast)**  
**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.

**URGENT**

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

**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](http://PRIMETHERAPEUTICS.COM/NOPP)**

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

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

MEDICATION OR MEDICAL DISPENSING INFORMATION			
<b>MEDICATION NAME:</b>			
<b>DOSE/STRENGTH:</b>	<b>FREQUENCY:</b>	<b>LENGTH OF THERAPY/REFILLS:</b>	<b>QUANTITY:</b>
<input type="checkbox"/> <b>NEW THERAPY</b> <input type="checkbox"/> <b>RENEWAL</b> <b>IF RENEWAL: DATE THERAPY INITIATED:</b>			
<b>DURATION OF THERAPY (SPECIFIC DATES):</b>			

*Continued on next page*

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1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?		
<input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
<b>MEDICATION/THERAPY</b> (SPECIFY DRUG NAME AND DOSAGE):	<b>DURATION OF THERAPY</b> (SPECIFY DATES):	<b>RESPONSE/REASON FOR FAILURE/ALLERGY:</b>
<b>2. LIST DIAGNOSES:</b>		<b>ICD-10:</b>
<input type="checkbox"/> Plaque Psoriasis <input type="checkbox"/> Plaque Psoriasis including intertriginous areas <input type="checkbox"/> Moderate to Severe Seborrheic dermatitis <input type="checkbox"/> Mild to Moderate Atopic Dermatitis  <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s):		
3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.		
Is patient going to be using drug in combination with a clinical trial? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Is the prescriber a dermatologist? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Is the request for : Zoryve 0.3% Cream? <input type="checkbox"/> Yes <input type="checkbox"/> No      Zoryve 0.3% FOAM? <input type="checkbox"/> Yes <input type="checkbox"/> No      Zoryve 0.15% Cream? <input type="checkbox"/> Yes <input type="checkbox"/> No		
<u>For diagnosis of Psoriasis, please answer the following:</u> Is the request for Zoryve 0.3% CREAM? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Is the psoriasis affecting 2% - 20% of body surface area? <input type="checkbox"/> Yes <input type="checkbox"/> No (documentation required)		
Has the patient had a trial and failure to at least two of the following four topical therapies (topical corticosteroid, topical vitamin D analog, topical calcineurin inhibitor, anthralin) <input type="checkbox"/> Yes <input type="checkbox"/> No (documentation required for drugs, dates, directions and therapy length)		
Is the patient receiving therapy with Otezla (apremilast) tablets or any other systemic immunomodulating agent? <input type="checkbox"/> Yes <input type="checkbox"/> No (documentation required if answer is yes)		
<u>For diagnosis of Seborrheic dermatitis, please answer the following:</u> Is the request for Zoryve 0.3% FOAM? <input type="checkbox"/> Yes <input type="checkbox"/> No		
Patient has a diagnosis of moderate or severe seborrheic dermatitis? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide documentation.		
Patient has had seborrheic dermatitis for at least 3 months? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide documentation.		



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Patient has tried at least 3 different topical treatments for seborrheic dermatitis, such as topical steroids, sulfur/sulfacetamide products, antifungals, selenium sulfide products, zinc pyrithione products and/or topical calcineurin inhibitor products?  Yes  No Please provide documentation with dates of service.

For diagnosis of mild to moderate Atopic dermatitis, please answer the following:

Is the request for Zoryve 0.15% CREAM?  Yes  No

Has patient had atopic dermatitis for at least 3 months?  Yes  No Please provide documentation.

Has the patient tried at least at least 2 different topical steroids?  Yes  No Please provide documentation.

If the patient has not tried at least 2 different topical steroids, has the patient tried at least one topical steroid AND one topical calcineurin inhibitor (tacrolimus or pimecrolimus)?  Yes  No Please provide documentation.

Renewal Information:

Is the prescriber a dermatologist?  Yes  No

Has the member shown improvement in condition over baseline?  Yes  No (documentation required)

Is the patient receiving therapy with Otezla (apremilast) tablets, Sotyktu(deucravacitinib) or any other systemic immunomodulating agent?  Yes  No (documentation required if answer is yes)

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

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**Please note:** Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

**ATTESTATION:** 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:** \_\_\_\_\_

**CONFIDENTIALITY NOTICE:** 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

Attn: CP-4201

P.O. Box 64811

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St. Paul, MN 55164-0811  
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