

## INSTRUCTIONS FOR COMPLETING THIS FORM

Please fill out all sections on all pages completely and legibly. If the following information is not complete, correct, or legible, the prior authorization (PA) process can be delayed. Use one form per member, per drug, please. This form, along with other PA forms can be found at: <https://welcome.optumrx.com/tenncare/landing>.

## MEMBER INFORMATION

Member Last Name: \_\_\_\_\_  
 Member First Name: \_\_\_\_\_  
 Member ID: \_\_\_\_\_  
 Date of Birth (MM/DD/YYYY): \_\_\_\_\_ Sex:  Male  Female  
 Street Address: \_\_\_\_\_  
 City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_  
 Phone Number: \_\_\_\_\_

## PRESCRIBER INFORMATION

Prescriber Last Name: \_\_\_\_\_  
 Prescriber First Name: \_\_\_\_\_  
 Prescriber NPI: \_\_\_\_\_ Prescriber DEA: \_\_\_\_\_  
 Specialty: \_\_\_\_\_ Office Phone: \_\_\_\_\_ Office Fax: \_\_\_\_\_  
 Supervising Physician and DEA (if applicable): \_\_\_\_\_  
 Office Street Address: \_\_\_\_\_  
 City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_  
 Is the prescriber a single patient contract holder for this patient?  Yes  No  
 Is the prescriber a TennCare provider with a TN Medicaid ID?  Yes  No

## MEDICATION AND DISPENSING INFORMATION

Please refer to the [Preferred Drug List](#) or see below for Preferred and Non-Preferred Topical Immunomodulators.

With the exception of the “[Branded Drugs to be Classified as Generics List](#),” TennCare is a mandatory generic program in accordance with state law (TCA 53-10-205). Approval of Non-Preferred agents requires trial and failure, contraindication, or intolerance of two (2) preferred agents, unless otherwise indicated on the [Preferred Drug List \(PDL\)](#).

### Antipsoriatics, Topical

Preferred Drugs	Non-Preferred Drugs
calcipotriene cream <sup>PA, QL</sup>	calcipotriene ointment <sup>PA, QL</sup>
calcipotriene foam <sup>PA, QL</sup>	Calcitrene® ointment <sup>PA, QL</sup>
calcipotriene scalp soln <sup>PA</sup>	calcitriol ointment <sup>PA, QL</sup>
Taclonex® <sup>PA</sup>	calcipotriene/betamethasone <sup>PA, QL</sup>
tazarotene 0.1% cream <sup>PA</sup>	Enstilar® <sup>PA, QL</sup>
	Sorilux® <sup>PA, QL</sup>
	tazarotene 0.1% gel <sup>PA, QL</sup>
	Vtama® <sup>PA, QL</sup>
	Zoryve® 0.3% cream <sup>PA, QL</sup>

This document and others if attached contain information that is privileged, confidential, and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of Optum Rx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing, or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Member Last Name: \_\_\_\_\_ DOB: \_\_\_\_\_

### Atopic Dermatitis Agents, Topical

Preferred Drugs	Non-Preferred Drugs
Eucrisa <sup>®</sup> PA, QL	Anzupgo <sup>®</sup> PA, QL
pimecrolimus <sup>QL</sup>	Opzelura <sup>®</sup> PA, QL
tacrolimus <sup>QL</sup>	Zoryve <sup>®</sup> 0.15% cream <sup>PA, QL</sup>

### Antiseborrheic Agents

Preferred Drugs	Non-Preferred Drugs
selenium sulfide 2.5% lotion <sup>QL</sup>	Ovace Plus <sup>®</sup>
	Ovace Plus Wash <sup>®</sup>
	selenium sulfide shampoo
	sodium sulfacetamide gel, liquid, shampoo
	sodium sulfacetamide /urea pads
	Zoryve <sup>®</sup> 0.3% topical foam <sup>PA, QL</sup>

Drug Name: \_\_\_\_\_ Drug strength: \_\_\_\_\_

Drug Formulation: \_\_\_\_\_ Dosing Frequency: \_\_\_\_\_

Quantity: \_\_\_\_\_ Day Supply: \_\_\_\_\_

Directions for use: \_\_\_\_\_

Location(s) on body: \_\_\_\_\_ Duration of treatment: \_\_\_\_\_

### CLINICAL CRITERIA

1. What is the diagnosis?
  - Atopic dermatitis
  - Eczema
  - Chronic Hand Eczema (CHE)
  - Plaque Psoriasis
  - Nonsegmental Vitiligo
  - Seborrheic dermatitis
  - Other: \_\_\_\_\_
2. Please provide the ICD-10 code for the diagnosis: \_\_\_\_\_
3. Has the patient tried any other agents in the past for this condition?  Yes  No
  - a. If yes, please list below.
    - i. Drug: \_\_\_\_\_ Dose: \_\_\_\_\_  
Frequency: \_\_\_\_\_ Duration: \_\_\_\_\_  
Result:  Currently taking  Failure/intolerance, reason: \_\_\_\_\_  
 Contraindication: \_\_\_\_\_
    - ii. Drug: \_\_\_\_\_ Dose: \_\_\_\_\_  
Frequency: \_\_\_\_\_ Duration: \_\_\_\_\_  
Result:  Currently taking  Failure/intolerance, reason: \_\_\_\_\_  
 Contraindication: \_\_\_\_\_
    - iii. Drug: \_\_\_\_\_ Dose: \_\_\_\_\_  
Frequency: \_\_\_\_\_ Duration: \_\_\_\_\_  
Result:  Currently taking  Failure/intolerance, reason: \_\_\_\_\_  
 Contraindication: \_\_\_\_\_
4. If the request is for **Opzelura<sup>®</sup>**, please answer this question.
  - a. Is the patient immunocompromised?  Yes  No
  - b. Is the patient breastfeeding?  Yes  No
  - c. Will Opzelura be used for short-term and/or non-continuous chronic treatment of atopic dermatitis?  
 Yes  No

*Question 4 continued on next page.*

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- d. Does the prescriber attest to all of the following?  Yes  No
  - i. Patient is not concurrently taking biologic agents (i.e., adalimumab, anakinra, etanercept, rituximab, tocilizumab, infliximab, abatacept) OR potent immunosuppressants (i.e., azathioprine, cyclosporine);
  - ii. Benefits of using this agent outweigh the risks of heart-related events (heart attack, stroke, blood clots, etc.) or cardiovascular risk factors.
  - iii. Risk of malignancy has been considered, and it has been determined that JAK Kinase inhibitor therapy is appropriate
- e. If it is a continuation of therapy, did the patient have a positive response to therapy (e.g. reduction in symptoms (itch, rash), re-pigmentation)?  Yes  No
- 5. If the request is for **Zoryve**<sup>®</sup>, please answer this question.
  - a. Does the patient have moderate to severe liver impairment (Child-Pugh B or C)?  Yes  No
  - b. If it is a continuation of therapy, did the patient have a positive response to therapy (e.g. reduction in itch, rash, inflammation)?  Yes  No
- 6. If the request is for **Vtama**<sup>®</sup>, please answer this question.
  - a. If it is a continuation of therapy, did the patient have a positive clinical response to therapy as evidenced by one of the following?
    - i. Reduction in the body surface area (BSA) involvement from baseline  Yes  No
    - ii. Improvement in symptoms (e.g., pruritic, inflammation) from baseline  Yes  No

Note: Please submit documentation to support the positive clinical response.
- 7. If the request is for **Anzupgo**<sup>®</sup>, please answer this question.
  - a. Is the patient taking other Janus kinase (JAK) inhibitors?  Yes  No
  - b. Is the patient taking any other potent immunosuppressants (e.g., azathioprine, cyclosporine)?  Yes  No
- 8. Please include any other information pertinent to this request: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

## **PRESCRIBER SIGNATURE**

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By signature, the prescriber confirms the above information is accurate and verifiable by patient records.

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

## **INSTRUCTIONS FOR SUBMISSION**

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PA requests can be submitted to Optum Rx via fax (866-434-5523), by phone (866-434-5524), or by electronic PA request, such as CoverMyMeds. For questions, please call 1-866-434-5524. Optum Rx will provide a response within 24 hours upon receipt.