

Division: Pharmacy Policy	Subject: Prior Authorization Criteria
Original Development Date: Original Effective Date: Revision Date:	May 10, 2024

Zoryve® (roflumilast)

LENGTH OF AUTHORIZATION: Up to one year

REVIEW CRITERIA:

CREAM:

- Patient must be ≥ 6 years of age.
- Patient must have a documented diagnosis of plaque psoriasis, including intertriginous areas, with 2% to 20% involvement of body surface area (BSA).
- Patient has had an inadequate response, intolerance, or contraindication to a minimum 4-week trial duration on at least 1 of the following (clinical documentation demonstrating prior treatment failures must be provided):
 - Preferred topical corticosteroids; OR
 - o Calcipotriene.
- Medication is prescribed by, or in consultation with a dermatologist.
- Prescriber attests that the patient does not have moderate to severe liver impairment (Child-Pugh B or C).

FOAM:

- Patient must be ≥ 9 years of age.
- Patient must have a documented diagnosis of seborrheic dermatitis.
- Patient has had an inadequate response, intolerance, or contraindication to a minimum 4-week trial duration on at least 1 of the following (clinical documentation demonstrating prior treatment failures must be provided):
 - O Topical antifungals (e.g., ciclopirox, ketoconazole, etc.); **OR**
 - o Preferred topical corticosteroids.
- Medication is prescribed by, or in consultation with a dermatologist.
- Prescriber attests that the patient does not have moderate to severe liver impairment (Child-Pugh B or C).

Note: The propellants in Zoryve[®] foam are flammable. Patients should be advised to avoid fire, flame, and smoking during and immediately following application.

CONTINUATION OF THERAPY

- Patient met initial review criteria.
- Documentation of improved clinical response.
- Dosing is appropriate as per labeling or is supported by compendia.





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DOSING AND ADMINISTRATION:

- Refer to product labeling at https://www.accessdata.fda.gov/scripts/cder/daf/
- Available as a 0.3% cream and foam

