

Request for Prior Authorization Select Topical Agents

FAX Completed Form To 1 (800) 574-2515 Provider Help Desk 1 (877) 776-1567

(PLEASE PRINT – ACCURACY IS IMPORTANT)

IA Medicaid Member ID #	Patient name	Patient name	
Patient address			
Provider NPI	Prescriber name	Prescriber name	
Prescriber address			Fax
Pharmacy name	Address		Phone
Prescriber must complete all info	ormation above. It must be legible,	correct, and complete or fo	orm will be returned.
Pharmacy NPI	Pharmacy fax	NDC	
I. Request adheres to all F contraindications, warning 2. Patient has a diagnosis a. Request is for reduction b. Patient has a diagnosis a. Request is for reductive were 3. Patient has a diagnosis a. Request is for reduction b. Patient has document of the properties of the properties of the properties of the patient has a diagnosis a. Request is for reduction b. Patient has a diagnosis a. Request is for reduction b. Patient has failed c. Patient has document of the properties of the pr	of seborrheic dermatitis; and oflumilast 0.3% foam; and umentation of an adequate trial and corticosteroid (scalp-medium to all for a minimum of 4 consecutive of mild to moderate atopic derma oflumilast 0.15% cream or tapinared to respond to good skin care arumentation of an adequate trial and corticosteroid for a minimum of 2 umentation of an adequate trial and an adequa	ted drug and indication, intions, and use in specific pent estimated to affect ≤ 2 of 1% cream; and therapy failure of combined a preferred topical vitaminal therapy failure of combining potency or nonscalpweeks; or titis; and regular use of emollient and therapy failure with one consecutive weeks; or and therapy failure with a topical vitaminal therapy failure with a topical vitaminal therapy failure with a topical vitaminal vitam	cluding age, dosing, populations; and 0% of the body surface area; and ination therapy with a preferred in D analog for a minimum of 4 dination therapy with a low potency) and preferred s; and a preferred medium to high spical immunomodulator for a
Non-Preferred			
□ Vtama □ Zoryve			
Strength	Usage Instructions	Quantity	Day's Supply
Diagnosis:			
□ Plaque Psoriasis			
_	ency Topical Corticosteroid Tri		

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Preferred Topical Vitamin D Analog Trial:				
Drug name & dose:	Trial dates:			
Failure reason:				
Is affected area estimated to affect ≤ 20% body surface area? ☐ Yes	□ No			
□ Seborrheic Dermatitis				
Preferred Topical Corticosteroid Trial: Scalp Nonscalp Drug name & dose: Failure reason:				
Failure reason:				
Preferred Topical Antifungal Trial: Drug name & dose: Failure reason:				
☐ Mild to moderate atopic dermatitis				
Preferred Medium to High Potency Topical Corticosteroid Trial:				
Drug name & dose:				
Preferred Topical Immunomodulator Trial:				
Drug name & dose:				
Has patient failed to respond to good skin care and regular use of emollients? ☐ Yes ☐ No				
Medical or contraindication reason to override trial requirements:				
ttach lab results and other documentation as necessary.				
	f submission			

IMPORTANT NOTE: In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Health and Human Services, that the member continues to be eligible for Medicaid.

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