

Otezla (apremilast)

Member and Medication Information	
* indicates required field	
*Member ID:	*Member Name:
*DOB:	*Weight:
*Medication Name/ Strength:	
<input type="checkbox"/> Do Not Substitute. Authorizations will be processed for the preferred Generic/Brand equivalent unless specified.	
*Directions for use:	
Provider Information	
* indicates required field	
*Requesting Provider Name:	*Requesting Prescriber NPI:
Address:	
*Contact Person:	*Office Phone:
*Office Fax:	*Office Email:
Fax form and relevant documentation including: laboratory results, chart notes and/or updated provider letter to Pharmacy PA at 855-828-4992 , to prevent processing delays.	

PART I: Criteria for Approval: (All of the following criteria must be met) - **Then move to PART II**

1. Is the medication being prescribed by or in consultation with a provider specializing in the disease treatment? Yes No
2. Does the patient have any of the following diagnoses?: (check the applicable) Yes No
 - Oral Ulcers Associated with Behçet's Disease
 - Plaque Psoriasis (PsO)
 - Psoriatic Arthritis (PsA)
 - Other - Off Label or Compendia Use (specify): _____
3. Does the requested medication and diagnosis follow FDA-approved age, dosing, monitoring and contraindications? Yes No
4. Does the provider attest that the patient is not taking concurrent treatment or that the medication will not be used in combination with other TNF-inhibitor, biologic response modifier or other biologic DMARDs, Janus kinase Inhibitors, or Phosphodiesterase 4 inhibitor (i.e., apremilast, tofacitinib, baricitinib) as verified by prescriber attestation, patient medication fill history, or submitted documentation? Yes No

If answer is No, go to Part II, section 4

PART II: Select and fill out applicable sections:**Section 1: Additional criteria for Oral Ulcers Associated with Behçet's Disease** (All of the following criteria must be met)

1. Is the patient at least 18 years of age or older? Yes No
2. Does the patient have a diagnosis of Oral Ulcers Associated with Behçet's Disease demonstrated by description of baseline symptoms present in the chart notes? Yes No
3. Has the patient tried and failed a topical steroid for at least 7 days, unless intolerated or contraindicated? Yes No
4. Has the patient failed treatment of at least **ONE** conventional immunosuppressant therapy (e.g., systemic corticosteroids, interferon alfa, colchicine)? Yes No

UTAH MEDICAID PHARMACY PRIOR AUTHORIZATION REQUEST FORM

Section 2: Additional criteria for Plaque Psoriasis (PsO) (All of the following criteria must be met)

1. Is the patient at least 6 years of age or older? Yes No
2. Does the patient weigh at least 20 kg? Yes No
3. Has the patient been diagnosed with moderate to severe plaque psoriasis involving greater than 3% body surface area? Yes No
4. If less than 3% of the body is involved, is there scalp, palmar, foot, or groin involvement causing significant disability? Yes No
5. Has the patient tried and failed at least **ONE** of the following medications for at least 3 months, or has a contraindication to all 3, if clinically appropriate? Yes No
 - methotrexate
 - cyclosporine
 - acitretin therapy

Section 3: Additional criteria for Psoriatic Arthritis (PsA) (All of the following criteria must be met)

1. Is the patient at least 6 years of age or older? Yes No
2. Does the patient weigh at least 20 kg? Yes No
3. Does the patient have a diagnosis of active psoriatic arthritis demonstrated by description of baseline symptoms present in the chart notes? Yes No
4. Has the patient had an adequate trial and failure of at least **ONE** of the following disease-modifying antirheumatic drugs (DMARDs) for at least 3 months, unless all are contraindicated: methotrexate, leflunomide, sulfasalazine, azathioprine, if clinically appropriate? Yes No

Section 4: Other - Off Label or Compendia Use of FDA-Approved Drugs Additional Criterion:

1. Does the submitted clinical documentation show an adequate trial and failure of at least **ONE** FDA-labeled and Medicaid preferred medication, if applicable? Yes No
2. Does the provider attest that the requested drug is being used for a medically accepted indication that is supported by information from the appropriate *compendia** of current literature. Including at least (1) major multi-site study or three (3) smaller studies published in JAMA, NEJM, Lancet, or other peer review specialty medical journals in the most recent years? Yes No

* Compendia use must be recommended by generally accepted compendia such as American Hospital Formulary Service Drug Information (AHFS), United States Pharmacopeia-Drug Information (USP-DI), the Micromedex Information System, Pediatric and Neonatal Lexi-Drugs, or clinical guidelines.

Reauthorization Criteria:

1. Has the provider submitted updated chart notes demonstrating a positive clinical response? Yes No

Initial Authorization: Up to six (6) months

Reauthorization: Up to one (1) year

PROVIDER CERTIFICATION

I hereby certify this treatment is indicated, necessary and meets the guidelines for use.

Prescriber's Signature

Date