

**CYTOKINE AND CAM ANTAGONISTS PRIOR AUTHORIZATION FORM** *(form effective 1/9/2023)*

Prior authorization guidelines for Cytokine and CAM Antagonists and Quantity Limits/Daily Dose Limits are available on the DHS Pharmacy Services website at <https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	# of pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street address:	
Beneficiary name:			City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:	

**CLINICAL INFORMATION**

<b>STARTER PACK</b> requested (name/strength):		<b>MAINTENANCE</b> product/packaging requested (name/strength):	
Quantity per fill:	Refills:	Quantity per fill:	Refills:
Directions:		Directions:	
Diagnosis ( <i>submit documentation</i> ):		Dx code ( <i>required</i> ):	Beneficiary weight:
Is the beneficiary currently being treated with the requested medication?		<input type="checkbox"/> Yes – date of last dose: _____ <i>Submit documentation.</i> <input type="checkbox"/> No	
Is the requested medication prescribed by or in consultation with a specialist (eg, rheumatologist, dermatologist, gastroenterologist, etc)?		<input type="checkbox"/> Yes <i>If prescriber is not a specialist, submit documentation of consultation.</i> <input type="checkbox"/> No	

**Complete all sections that apply to the beneficiary and this request.**  
**Check all that apply and submit documentation for each item.**

**INITIAL requests**

**DRUG**

- Requested drug is NON-PREFERRED on the Statewide PDL:**  
 Tried and failed or has a contraindication or intolerance to the preferred drugs in this class approved or medically accepted for the beneficiary's condition (*Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.*)
- Requested drug is OTEZLA (apremilast) or SILIQ (brodalumab):**  
 Was evaluated for history of prior suicide attempt, bipolar disorder, or major depressive disorder
- Requested drug is an oral JAK inhibitor (eg, Olumiant [baricitinib], Rinvoq [upadacitinib], Xeljanz [tofacitinib]):**  
 Tried and failed at least one TNF blocker or another biologic as recommended in the JAK inhibitor's package labeling  
 Has a contraindication or an intolerance to TNF blockers or other biologics as recommended in the JAK inhibitor's package labeling

**DIAGNOSIS**

- ALL diagnoses:**  
 Screened for hepatitis B virus infection (surface antigen, surface antibody, and core antibody)  
 Screened for tuberculosis

2. **Adult-onset Still's disease:**
  - Has predominantly systemic disease:
    - Has steroid-dependent disease
    - Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
  - Has predominantly joint disease:
    - Tried and failed or has a contraindication or an intolerance to conventional DMARDs (eg, MTX)
3. **Alopecia areata:**
  - Has alopecia universalis
  - Has >50% scalp involvement or alopecia totalis
  - Has alopecia that causes significant disability or impaired physical, mental, or psychosocial functioning
  - Has a current episode of alopecia areata that has lasted at least 6 months
4. **Ankylosing spondylitis & non-radiographic axial spondyloarthritis:**
  - Tried and failed a 2-week trial of or has a contraindication or an intolerance to 2 different oral NSAIDs
5. **Behçet's syndrome:**
  - Has recurrent oral ulcers associated with Behçet's syndrome
  - Tried and failed or has a contraindication or an intolerance to a topical corticosteroid (eg, triamcinolone dental paste)
  - Tried and failed a 3-month trial of or has a contraindication or an intolerance to colchicine at maximally tolerated doses
6. **Crohn's disease:**
  - Has moderate-to-severe disease
  - Has disease that is associated with high-risk or poor prognostic features
  - Tried and failed to achieve remission with or has a contraindication or an intolerance to an induction course of corticosteroids
  - Tried & failed to maintain remission with or has a contraindication or an intolerance to conventional immunomodulators (eg, AZA, 6-MP, MTX)
7. **Familial Mediterranean fever:**
  - Tried and failed a 3-month trial of or has a contraindication or an intolerance to colchicine at maximally tolerated doses
8. **Giant cell arteritis:**
  - Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
  - Is at high risk for glucocorticoid-related complications
  - Has steroid-dependent disease
9. **Hidradenitis suppurativa (HS):**
  - Has Hurley stage II or stage III disease
  - Is a candidate for or has a history of surgical intervention for HS
  - Tried and failed a 3-month trial of or has a contraindication or an intolerance to topical clindamycin
  - Tried and failed or has a contraindication or an intolerance to systemic antibiotics (eg, doxycycline, minocycline, tetracycline, clindamycin)
10. **Juvenile idiopathic arthritis:**
  - Has systemic disease with active systemic features
  - Has disease associated with any of the following:
    - Positive anti-CCP antibodies
    - Presence of joint damage
    - High disease activity
    - Positive rheumatoid factor
    - At high risk of disabling joint damage
    - Involvement of high-risk joints (cervical spine, hip, wrist)
  - Tried and failed a 3-month trial of or has a contraindication or an intolerance to conventional DMARDs (eg, MTX)
  - Has active sacroiliitis and/or enthesitis:
    - Tried and failed a 2-week trial of or has a contraindication or an intolerance to oral NSAIDs
11. **Plaque psoriasis:**
  - Has a BSA of  $\geq 3\%$  that is affected
  - Has involvement of critical areas of the body (eg, skin folds, face, genitals)

- Has psoriasis that causes significant disability or impaired physical, mental, or psychosocial functioning
- Has moderate-to-severe nail disease
- Tried and failed a 4-week trial of or has a contraindication or an intolerance to topical corticosteroids
- Tried and failed an 8-week trial of or has a contraindication or an intolerance to non-steroid topical medications (eg, anthralin, calcineurin inhibitor, tazarotene, etc)
- Tried and failed or has a contraindication or an intolerance to ultraviolet light therapy
- Tried & failed a 3-month trial of or has a contraindication/intolerance to conventional systemic medications (eg, MTX, cyclosporine, acitretin)

**12. Psoriatic arthritis:**

- Tried and failed an 8-week trial of or has a contraindication or an intolerance to conventional DMARDs (eg, AZA, leflunomide, MTX, SSZ)
- Has predominantly axial disease, dactylitis, and/or enthesitis
- Has severe disease
- Has comorbid moderate-to-severe nail psoriasis
- Has comorbid active inflammatory bowel disease

**13. Rheumatoid arthritis:**

- Tried and failed a 3-month trial of or has a contraindication or an intolerance to conventional DMARDs (eg, AZA, leflunomide, MTX, etc)

**14. Sarcoidosis:**

- Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
- Has steroid-dependent disease
- Tried and failed or has a contraindication or an intolerance to a conventional DMARD (eg, AZA, leflunomide, MTX, mycophenolate)

**15. Ulcerative colitis:**

- Has moderate-to-severe disease
- Has disease associated with multiple poor prognostic factors
- Tried and failed to achieve remission with or has a contraindication or an intolerance to an induction course of corticosteroids
- Tried and failed to maintain remission with or has a contraindication or an intolerance to conventional immunomodulators (eg, AZA, cyclosporine, 6-MP, MTX)

**16. Uveitis (non-infectious):**

- Has comorbid juvenile idiopathic arthritis
- Has comorbid Behçet's syndrome
- Has steroid-dependent disease
- Tried and failed or has a contraindication or an intolerance to systemic, topical, intraocular, or periocular corticosteroids
- Tried and failed or has a contraindication or an intolerance to conventional systemic immunosuppressives (eg, AZA, MTX, MMF, etc)

**17. Other diagnosis:** \_\_\_\_\_

- List other treatments tried (including start/stop dates, dose, outcomes): \_\_\_\_\_

**RENEWAL requests**

- Experienced an improvement in disease severity or level of functioning since starting therapy with the requested medication
- Is prescribed an increased dose or more frequent administration of the requested medication
- Requested drug is OTEZLA (apremilast) or SILIQ (brodalumab):
  - Was recently reevaluated for behavioral and mood changes

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION**

Prescriber Signature:

Date:

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