

CYTOKINE AND CAM ANTAGONISTS PRIOR AUTHORIZATION FORM *(form effective 1/8/2024)*

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

STARTER PACK requested (drug name / strength / formulation [pen, syringe, tablet, etc.]):		MAINTENANCE product/packaging requested (drug name / strength / formulation [pen, syringe, tablet, etc.]):	
Quantity per fill:	Refills:	Quantity per fill:	Refills:
Directions:		Directions:	
Diagnosis (<u>submit documentation</u>):		Dx code (<u>required</u>):	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 BIMZELX (bimekizumab), 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 other 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

1. **ALL diagnoses:**
 - Screened for hepatitis B virus infection (surface antigen, surface antibody, and core antibody)
 - Screened for tuberculosis
2. **Adult-onset Still's disease (AOSD):**
 - Has predominantly systemic AOSD AND:
 - Has steroid-dependent AOSD
 - Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
 - Has predominantly joint AOSD AND:
 - 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 colchicine at maximally tolerated doses or has a contraindication or an intolerance to colchicine
6. **Crohn's disease (CD):**
 - Has moderate-to-severe CD AND:
 - 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, 6-MP, MTX)
 - Has CD that is associated with high-risk or poor prognostic features
 - Has achieved remission with the requested medication AND:
 - Will be using the requested medication as maintenance therapy to maintain remission
7. **Familial Mediterranean fever:**
 - Tried and failed a 3-month trial of colchicine at maximally tolerated doses or has a contraindication or an intolerance to colchicine
8. **Generalized pustular psoriasis (GPP):**
 - Request is for Spevigo (spesolimab) AND:
 - Beneficiary has received a single dose of Spevigo (spesolimab) for the current GPP flare AND:
 - Continues to experience moderate to severe GPP flare symptoms since the previous dose
 - Beneficiary has not received a dose of Spevigo (spesolimab) for the current GPP flare AND:
 - Is experiencing a moderate to severe GPP flare that warrants rapid stabilization or improvement
9. **Giant cell arteritis (GCA):**
 - Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
 - Is at high risk for glucocorticoid-related complications
 - Has steroid-dependent GCA
10. **Gout flares:**
 - Tried and failed maximally tolerated doses of or has a contraindication or an intolerance to NSAIDs
 - Tried and failed maximally tolerated doses of or has a contraindication or an intolerance to colchicine
 - Tried and failed maximally tolerated doses of or has a contraindication or an intolerance to corticosteroids

Has a medical reason why repeated courses of corticosteroids are not appropriate

11. Hidradenitis suppurativa (HS):

- Has Hurley stage II or stage III HS
- 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)

12. Juvenile idiopathic arthritis (JIA):

- Has systemic JIA with active systemic features
- Has JIA associated with any of the following:
 - Positive anti-CCP antibodies
 - Presence of joint damage
 - High disease activity
 - Positive rheumatoid factor
 - 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 AND:
 - Tried and failed a 2-week trial of or has a contraindication or an intolerance to oral NSAIDs

13. 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 psoriasis
- 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)

14. Polymyalgia rheumatica (PMR):

- Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids
- Has steroid-dependent PMR

15. Psoriatic arthritis (PsA):

- 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 PsA, dactylitis, and/or enthesitis
- Has severe PsA
- Has comorbid moderate-to-severe nail psoriasis
- Has comorbid active inflammatory bowel disease

16. 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)

17. Sarcoidosis:

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

18. Ulcerative colitis (UC):

- Has moderate-to-severe UC
- Has UC 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)
- Has achieved remission with the requested medication AND:

Will be using the requested medication as maintenance therapy to maintain remission

19. Uveitis (non-infectious):

- Has comorbid juvenile idiopathic arthritis
- Has comorbid Behçet's syndrome
- Has steroid-dependent uveitis
- 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)

20. 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 BIMZELX (bimekizumab), 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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