



# Prior Authorization Request Form for Cytokine and CAM Antagonists

**FAX this completed form to (844) 205-3386**

**OR Mail requests to: Pharmacy Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720**

**OR Prior authorization may be completed at <https://www.covermymeds.com/main/prior-authorization-forms/>**

<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:	
Member name:			City/state/zip:	
Member ID#:	DOB:	Phone:	Fax:	

## CLINICAL INFORMATION

<b>STARTER PACK requested (drug name / strength / formulation [pen, syringe, tablet, etc.]):</b>		<b>MAINTENANCE product/packaging requested (drug name / strength / formulation [pen, syringe, tablet, etc.]):</b>	
Quantity per fill:	Refills:	Quantity per fill:	Refills:
Directions:		Directions:	
Diagnosis ( <i>submit documentation</i> ):		Dx code ( <i>required</i> ):	Member weight:
Is the member 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 member 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 member's condition: \_\_\_\_\_
- 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**

**ALL diagnoses:**

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

**Adult-onset Still's disease (AOSD):**

- Has predominantly systemic AOSD AND one of the following:
  - Has steroid-dependent AOSD
  - Tried and failed systemic glucocorticoids: \_\_\_\_\_
  - Has a contraindication or an intolerance to systemic glucocorticoids: \_\_\_\_\_
- Has predominantly joint AOSD AND one of the following:
  - Tried and failed conventional DMARDs (eg, MTX): \_\_\_\_\_
  - Has a contraindication or an intolerance to conventional DMARDs (eg, MTX): \_\_\_\_\_

**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

**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: \_\_\_\_\_
- OR
- Has a contraindication or an intolerance to 2 different oral NSAIDs: \_\_\_\_\_

**Behçet's syndrome:**

- Has recurrent oral ulcers associated with Behçet's syndrome
- One of the following:
  - Tried and failed a topical corticosteroid (eg, triamcinolone dental paste): \_\_\_\_\_
  - Has a contraindication or an intolerance to a topical corticosteroid (eg, triamcinolone dental paste): \_\_\_\_\_
- One of the following:
  - Tried and failed a 3-month trial of colchicine at maximally tolerated doses: \_\_\_\_\_
  - Has a contraindication or an intolerance to colchicine: \_\_\_\_\_

**Crohn's disease (CD):**

- Has moderate-to-severe CD AND:
  - One of the following:
    - Tried and failed to achieve remission with an induction course of corticosteroids: \_\_\_\_\_
    - Has a contraindication or an intolerance to an induction course of corticosteroids: \_\_\_\_\_
    - Tried and failed to maintain remission with conventional immunomodulators (eg, AZA, 6-MP, MTX): \_\_\_\_\_
    - 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

**Familial Mediterranean fever:**

- Tried and failed a 3-month trial of colchicine at maximally tolerated: \_\_\_\_\_
- Has a contraindication or an intolerance to colchicine: \_\_\_\_\_

**Generalized pustular psoriasis (GPP):**

- Request is for Spevigo (spesolimab) AND:
  - Member 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
  - Member 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

**Giant cell arteritis (GCA):**

- Tried and failed or has a contraindication or an intolerance to systemic glucocorticoids: \_\_\_\_\_
- OR
- Has a contraindication or an intolerance to systemic glucocorticoids: \_\_\_\_\_
- Is at high risk for glucocorticoid-related complications: \_\_\_\_\_
- Has steroid-dependent GCA

**Gout flares:**

- One of the following:
- Tried and failed maximally tolerated doses of NSAIDs: \_\_\_\_\_
  - Has a contraindication or an intolerance to NSAIDs: \_\_\_\_\_
- One of the following:
- Tried and failed maximally tolerated doses of colchicine: \_\_\_\_\_
  - Has a contraindication or an intolerance to colchicine: \_\_\_\_\_
- One of the following:
- Tried and failed maximally tolerated doses of corticosteroids: \_\_\_\_\_
  - Has a contraindication or an intolerance to corticosteroids: \_\_\_\_\_
- 
- Has a medical reason why repeated courses of corticosteroids are not appropriate: \_\_\_\_\_

**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 topical clindamycin: \_\_\_\_\_
- Has a contraindication or an intolerance to topical clindamycin: \_\_\_\_\_
- Tried and failed systemic antibiotics (eg, doxycycline, minocycline, tetracycline, clindamycin): \_\_\_\_\_
- Has a contraindication or an intolerance to systemic antibiotics (eg, doxycycline, minocycline, tetracycline, clindamycin): \_\_\_\_\_

**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 conventional DMARDs (eg, MTX): \_\_\_\_\_
- 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 oral NSAIDs: \_\_\_\_\_
  - Has a contraindication or an intolerance to oral NSAIDs: \_\_\_\_\_

**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 topical corticosteroids: \_\_\_\_\_
- OR
- Has a contraindication or an intolerance to topical corticosteroids: \_\_\_\_\_
- Tried and failed an 8-week trial of non-steroid topical medications (eg, anthralin, calcineurin inhibitor, tazarotene, etc): \_\_\_\_\_
- OR
- Has a contraindication or an intolerance to non-steroid topical medications (eg, anthralin, calcineurin inhibitor, tazarotene, etc): \_\_\_\_\_

**Polymyalgia rheumatica (PMR):**

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

**Psoriatic arthritis (PsA):**

- Tried and failed an 8-week trial of conventional DMARDs (eg, AZA, leflunomide, MTX, SSZ): \_\_\_\_\_
- 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

**Rheumatoid arthritis:**

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

**Sarcoidosis:**

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

**Ulcerative colitis (UC):**

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

**Uveitis (non-infectious):**

- Has comorbid juvenile idiopathic arthritis
- Has comorbid Behçet's syndrome
- Has steroid-dependent uveitis
- Tried and failed systemic, topical, intraocular, or periocular corticosteroids: \_\_\_\_\_

Has a contraindication or an intolerance to systemic, topical, intraocular, or periocular corticosteroids: \_\_\_\_\_

Tried and failed conventional systemic immunosuppressives (eg, AZA, MTX, MMF, etc): \_\_\_\_\_

Has a contraindication or an intolerance to conventional systemic immunosuppressives (eg, AZA, MTX, MMF, etc): \_\_\_\_\_

**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

**ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION**

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO (844) 205-3386**

**Prescriber Signature:**

**Date:**

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Pharmacy Department will respond via fax or phone within 24 hours.

Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)