

CYTOKINE AND CAM ANTAGONISTS PRIOR AUTHORIZATION FORM (form effective 2/17/2022)

Prior authorization guidelines for **Cytokine and CAM Antagonists** and **Quantity Limits/Daily Dose Limits** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<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:			Suite #:	City/state/zip:
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

STARTER PACK requested (name/strength):		MAINTENANCE 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 (e.g., rheumatologist, dermatologist, gastroenterologist, etc.)?	<input type="checkbox"/> Yes <i>If prescriber is not a specialist, submit documentation of consultation.</i> <input type="checkbox"/> No	

Is the beneficiary using a different Cytokine and CAM Antagonist?

- Yes
- No

If Yes to the above question:

Will discontinue use of that Cytokine and CAM Antagonist prior to starting the requested Cytokine and CAM Antagonist

OR

- Has a medical reason for concomitant use of both Cytokine and CAM Antagonists that is supported by peer-reviewed medical literature or national treatment guidelines,
- Is dependent on glucocorticoids in addition to a Cytokine and CAM Antagonist to prevent life-threatening complications,
- Has 2 or more autoimmune or autoinflammatory conditions for which a single Cytokine and CAM Antagonist is not sufficient

INITIAL requests

Complete the sections below that are applicable to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

DRUG

Requested drug is **NON-PREFERRED drug**:

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

DIAGNOSIS

For a **Cytokine and CAM Antagonists associated with an increased risk of infection according to the FDA-approved package labeling**:

screened for hepatitis B (surface antigen, surface antibody, and core antibody)

screened for tuberculosis

Adult-onset Still's disease:

Has predominantly systemic disease:

Has steroid-dependent disease

Tried and failed or has a contraindication or intolerance to systemic glucocorticoids

Has predominantly joint disease:

Tried and failed or has a contraindication or intolerance to conventional DMARDs (eg, MTX)

Ankylosing spondylitis & non-radiographic axial spondyloarthritis:

Tried and failed a 2-week trial of or has a contraindication or intolerance to 2 different oral NSAIDs

Atopic dermatitis (eczema):

Tried and failed or has a contraindication or an intolerance to:

For the face or skin folds, low-potency (or higher) topical corticosteroids

For other body areas, a topical corticosteroid with a potency appropriate for the beneficiary's age and affected area(s) of the body

Elidel (pimecrolimus) or Protopic (tacrolimus)

Phototherapy/photochemotherapy (e.g., PUVA, UVB light)

Systemic immunosuppressives (e.g., azathioprine, cyclosporine, methotrexate, mycophenolate)

Behçet's syndrome:

Has recurrent oral ulcers associated with Behçet's syndrome

Tried and failed or has a contraindication or intolerance to a topical corticosteroid (eg, triamcinolone dental paste)

Tried and failed a 3-month trial of or has a contraindication or intolerance to colchicine at maximally tolerated doses

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 intolerance to an induction course of corticosteroids

Tried and failed to maintain remission with or has a contraindication or intolerance to immunomodulators (eg, AZA, 6-MP, MTX)

Familial Mediterranean fever:

Tried and failed a 3-month trial of or has a contraindication or intolerance to colchicine at maximally tolerated doses

Giant cell arteritis:

Tried and failed or has a contraindication or intolerance to systemic glucocorticoids

Is at high risk for glucocorticoid-related complications

Has steroid-dependent disease

- 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 intolerance to topical clindamycin
 - Tried and failed or has a contraindication or intolerance to systemic antibiotics (eg, doxycycline, minocycline, tetracycline, clindamycin)
- 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 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 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 or mental functioning
 - Tried and failed or has a contraindication or intolerance to topical corticosteroids
 - Tried and failed or has a contraindication or intolerance to non-steroid topical medications (eg, anthralin, calcineurin inhibitor, tazarotene, etc)
 - Tried and failed or has a contraindication or intolerance to ultraviolet light therapy
 - Tried and failed a 3-month trial of or has a contraindication or intolerance to oral systemic medications (eg, MTX, cyclosporine, acitretin)
- Psoriatic arthritis:**
- Has severe disease
 - Has comorbid moderate-to-severe nail psoriasis
 - Has predominantly axial disease and/or enthesitis
 - Has predominantly peripheral disease:**
 - Tried and failed an 8-week trial of or has a contraindication or intolerance to conventional DMARDs (eg, AZA, leflunomide, MTX, SSZ)
- Rheumatoid arthritis:**
- Tried and failed a 3-month trial of or has a contraindication or intolerance to conventional DMARDs (eg, AZA, leflunomide, MTX, etc)
- Sarcoidosis:**
- Tried and failed or has a contraindication or intolerance to systemic glucocorticoids
 - Has steroid-dependent disease
 - Tried and failed or has a contraindication or intolerance to a conventional DMARD (eg, AZA, leflunomide, MTX, mycophenolate)
- 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 intolerance to an induction course of corticosteroids
 - Tried and failed to maintain remission with or has a contraindication or intolerance to immunomodulators (eg, AZA, cyclosporine, 6-MP, MTX)
- 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 intolerance to systemic, topical, intraocular, or periocular corticosteroids
 - Tried and failed or has a contraindication or intolerance to conventional systemic immunosuppressives (eg, AZA, MTX, MMF, etc)

RENEWAL requests

Check all of the following that apply to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- 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
 - New dose/frequency is within the FDA-approved recommended dosing OR is supported by national compendia or peer-reviewed medical literature
- Requested drug is OTEZLA (apremilast) or SILIQ (brodalumab):
 - Was recently reevaluated for behavioral and mood changes

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

Prescriber Signature:

Date:

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