



Fax completed prior authorization request form to 800-854-7614 or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts.

All requested data must be provided. **Incomplete forms or forms without the chart notes will be returned**

Pharmacy Coverage Guidelines are available at www.mercycareaz.org/providers/complecare-forproviders/pharmacy

Cytokines and Cell Adhesion Molecule (CAM) Antagonists Pharmacy Prior Authorization Request Form

Do not copy for future use. Forms are updated frequently.

REQUIRED: Office notes, labs and medical testing relevant to request showing medical justification are required to support diagnosis

Member Information

Member Name (first & last):	Date of Birth:	Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Height:
Member ID:	City:	State:	Weight:

Prescribing Provider Information

Provider Name (first & last):	Specialty:	NPI#	DEA#
Office Address:	City:	State:	Zip Code:
Office Contact:	Office Phone	Office Fax:	

Dispensing Pharmacy Information

Pharmacy Name:	Pharmacy Phone:	Pharmacy Fax:
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Requested Medication Information

Preferred Agents:	<input type="checkbox"/> Enbrel	<input type="checkbox"/> Humira	<input type="checkbox"/> Otezla	<input type="checkbox"/> Xeljanz IR		
Non-Preferred Agents:	<input type="checkbox"/> Actemra	<input type="checkbox"/> Arcalyst	<input type="checkbox"/> Cosentyx	<input type="checkbox"/> Taltz	<input type="checkbox"/> Ilaris	<input type="checkbox"/> Ilumya
	<input type="checkbox"/> Kineret	<input type="checkbox"/> Siliq	<input type="checkbox"/> Orencia	<input type="checkbox"/> Renflexis	<input type="checkbox"/> Tremfya	<input type="checkbox"/> Tysabri
	<input type="checkbox"/> Olumiant	<input type="checkbox"/> Remicade	<input type="checkbox"/> Xeljanz XR	<input type="checkbox"/> Cimzia	<input type="checkbox"/> Skyrizi	<input type="checkbox"/> Simponi Aria
	<input type="checkbox"/> Simponi	<input type="checkbox"/> Stelara	<input type="checkbox"/> Inflectra	<input type="checkbox"/> Other, specify:		

Medication request is NOT for an FDA- approved, or compendia-supported diagnosis (circle one): Yes No	Diagnosis:	ICD-10 Code:
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Are there any contraindications to formulary medications? (if yes, specify):	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> New request	<input type="checkbox"/> Continuation of therapy
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Directions for Use:	Strength:	Dosage Form:	
	Quantity:	Day Supply:	Duration of Therapy/Use:

What medication(s) has the member tried and failed for this diagnosis? Please specify below.

Turn-Around Time for Review

<input type="checkbox"/> Standard – (24 hours)	<input type="checkbox"/> Urgent – waiting 24 hours for a standard decision could seriously harm life, health, or ability to regain maximum function, you can ask for an expedited decision.
Signature: _____	

Clinical Information

General Authorization Criteria

Is member on another Cytokine or Cell Adhesion Molecule (CAM) Antagonist?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is request for Anti-Tumor Necrosis Factor?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Does member have NYHA class III OR IV CHF? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Is request for Anti-Tumor Necrosis Factors such as Stelara, Xeljanz, Xeljanz XR, Kineret, Actemra, Ilaris OR Orencia?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
Was a screen completed for Hepatitis B?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Does member have active OR chronic Hepatitis B? <input type="checkbox"/> Yes <input type="checkbox"/> No
If member has active OR chronic Hepatitis B, is member receiving appropriate antiviral treatment?	<input type="checkbox"/> Yes	<input type="checkbox"/> No <input type="checkbox"/> N/A
Was member evaluated AND given appropriate vaccinations, as recommended per CDC, for risk factors?	<input type="checkbox"/> Yes	<input type="checkbox"/> No

Was member screened for TB?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	If screening was positive for latent TB, was treatment received for latent TB?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Is request for Entyvio OR Tysabri?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is use Monotherapy AND not in combination with antineoplastic, immunosuppressive OR immunomodulating agents (AZA, 6-MP, cyclosporine, MTX, TNF inhibitors)	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Additional Criteria Based on Indication:						
<input type="checkbox"/> Rheumatoid Arthritis						
Was there inadequate response to 3-month trial of MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there intolerance OR contraindication to MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Were SSZ, LEF or HCQ used due to intolerance OR contraindication to MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Will requested medication be used concurrently with MTX or another non-biologic DMARD such as SSZ, LEF or HCQ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
<input type="checkbox"/> Systemic Juvenile Idiopathic Arthritis						
Does member have ACTIVE SYSTEMIC FEATURES such as fever, evanescent rash, lymphadenopathy, hepatomegaly, splenomegaly OR serositis? (circle one):			Is synovitis in <u>ONE OR MORE JOINTS</u> despite 3 months treatment with <u>MTX OR LEF</u> ? (circle one):			
Yes No			Yes No			
Check if ONE of the following apply:	<input type="checkbox"/> There are ACTIVE SYSTEMIC FEATURES (fever, evanescent rash, lymphadenopathy, hepatomegaly, splenomegaly OR serositis) AND synovitis is in at least <u>ONE JOINT</u> .					
	<input type="checkbox"/> There are NO ACTIVE SYSTEMIC FEATURES (fever, evanescent rash, lymphadenopathy, hepatomegaly, splenomegaly OR serositis) AND synovitis is in <u>ONE OR MORE JOINTS</u> despite 3 months treatment with <u>MTX OR LEF</u> .					
There are ACTIVE SYSTEMIC FEATURES (fever, evanescent rash, lymphadenopathy, hepatomegaly, splenomegaly, or serositis) (circle one):			Synovitis is in ONE OR MORE JOINTS despite ONE-month treatment with Kineret OR Actemra AND MTX OR LEF (circle one):			
Yes No			Yes No			
<input type="checkbox"/> Polyarticular Juvenile Idiopathic Arthritis						
Was there inadequate response to 3-months trial with MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there an intolerance OR contraindication to MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Was there trial with SSZ OR LEF for 3 months?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A			
<input type="checkbox"/> Oligoarticular Juvenile Idiopathic Arthritis						
Is disease duration > 6 months?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there documented inadequate response OR intolerable side effect with 2 NSAIDs?	<input type="checkbox"/> Yes, indicate drug:		<input type="checkbox"/> No
Was there contraindication to NSAIDs?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A	Was there inadequate response OR intolerable side effect to 3-month trial with MTX?		<input type="checkbox"/> Yes <input type="checkbox"/> No
Was there documented trial of LEF OR SSZ for 3 months?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A			
<input type="checkbox"/> Cryopyrin-Associated Periodic Syndromes						
Indicate if ONE of the following subtypes is present:	<input type="checkbox"/> Familial Cold Auto Inflammatory Syndrome		<input type="checkbox"/> Muckle-Wells syndrome	<input type="checkbox"/> Neonatal onset multi-system inflammatory disease		
Was there 3-months trial with Kineret?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A			
<input type="checkbox"/> Familial Mediterranean Fever						
Was there inadequate response, intolerance OR contraindication to colchicine at MAX indicated dose?	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
<input type="checkbox"/> Giant Cell Arteritis						
Was there inadequate response with glucocorticoids (prednisone, methylprednisolone)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there intolerance OR contraindication to glucocorticoids?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
If member had intolerance OR contraindication to glucocorticoids, was there TRIAL with MTX OR cyclophosphamide?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A	Will medication be used in combination with tapering course of glucocorticoids	<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> Ankylosing Spondylitis						
Was there inadequate response to ONE-month trial of TWO NSAIDs?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is there contraindication OR intolerance to oral NSAIDs?		<input type="checkbox"/> Yes	<input type="checkbox"/> No <input type="checkbox"/> N/A
<input type="checkbox"/> Psoriatic Arthritis						
Does member have ACTIVE Psoriatic Arthritis?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there inadequate response to 3-months trial with MTX?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Was there intolerance OR contraindication to MTX?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A	Was there 3-month trial of SSZ OR LEF?		<input type="checkbox"/> Yes <input type="checkbox"/> No
Is disease predominantly AXIAL OR ACTIVE ENTHESITIS / DACTYLITIS?				<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Was there inadequate response to ONE-month trial of 2 NSAIDs?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there contraindication OR intolerance to oral NSAIDs?		<input type="checkbox"/> Yes	<input type="checkbox"/> No <input type="checkbox"/> N/A
<input type="checkbox"/> Plaque Psoriasis						
Was there inadequate response to MTX OR cyclosporine for ≥3 months?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there intolerance OR contraindication to MTX OR cyclosporine for ≥3 months?		<input type="checkbox"/> Yes	<input type="checkbox"/> No <input type="checkbox"/> N/A
Is >10% BSA affected?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is <10% BSA affected BUT involves sensitive areas such as hands, feet, face OR genitals?		<input type="checkbox"/> Yes	<input type="checkbox"/> No

Is Psoriasis Area and Severity Index score >10?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was phototherapy PUVA, UVB ineffective?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
For Siliq ONLY:	Is there history of a prior suicide attempt, bipolar disorder OR depressive disorder?		<input type="checkbox"/> Yes	<input type="checkbox"/> No	
	Was a mental health evaluation completed by prescriber OR psychiatrist?		<input type="checkbox"/> Yes	<input type="checkbox"/> No	
<input type="checkbox"/> Ulcerative Colitis					
<input type="checkbox"/> STEROID DEPENDENT	A relapse occurred within 3-months of stopping glucocorticoids (circle one):		There is inability to taper steroids to acceptable dose after 3 months W/O having symptom recurrence:(circle one):		
	Yes No		Yes No		
<input type="checkbox"/> STEROID REFRACTORY	Inadequate response OR intolerable side effect to IV glucocorticoids after 7-10 days (circle one):		Inadequate response OR intolerable side effect to oral prednisone ≥40mg per day after 30 days (circle one):		
	Yes No		Yes No		
<input type="checkbox"/> Crohn's Disease					
<input type="checkbox"/> STEROID DEPENDENT	A relapse occurred within 3-months of stopping glucocorticoids (circle one):		There was inadequate response OR intolerable side effect, with 3-month trial of 6-MP OR AZA OR injectable MTX (circle one):		
	Yes No		Yes No		
<input type="checkbox"/> STEROID REFRACTORY	There is inability to taper steroids to acceptable dose after 3 months W/O having symptom recurrence (circle one):		There was contraindication to 6-MP, AZA, AND injectable MTX (circle one):		
	Yes No		Yes No		
<input type="checkbox"/> STEROID REFRACTORY	There was inadequate response OR intolerable side effect to IV glucocorticoids after 7-10 days (circle one):		There was inadequate response OR intolerable side effect to oral prednisone ≥40mg per day after 30 days (circle one):		
	Yes No		Yes No		
<input type="checkbox"/> Hidradenitis Suppurativa (Acne Inversa)					
Does member have moderate to severe disease (Hurley stage II-III)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there trial AND failure of 90-day treatment with oral antibiotics (doxycycline, minocycline OR clindamycin with rifampin)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> Behçet's Disease					
Does member have ACTIVE RECURRENT oral ulcers?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there trial AND failure with ONE Non-Biologic DMARD (MTX, LEF, SSZ OR HCQ)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> Uveitis					
Was intermediate, posterior OR pan uveitis caused by infection?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	There was inadequate response OR intolerable side effect with following:	<input type="checkbox"/> cyclosporine	<input type="checkbox"/> tacrolimus
				<input type="checkbox"/> MTX	<input type="checkbox"/> AZA
Are medications such as corticosteroids, MTX, AZA, MMF, cyclosporine, AND tacrolimus are NOT appropriate?				<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> Cytokine Release Syndrome					
Is diagnosis grade 3 OR 4, severe OR life-threatening due to chimeric antigen receptor-T cell therapy?				<input type="checkbox"/> Yes	<input type="checkbox"/> No
Additional information the prescribing provider feels is important to this review. Please specify below or submit medical records					

Signature affirms that information given on this form is true and accurate and reflects office notes.

Prescribing Provider's Signature: _____ **Date:** _____

Please note: Incomplete forms or forms without the chart notes will be returned

Office notes, labs, and medical testing relevant to the request that show medical justification are required.
Standard turnaround time is 24 hours. You can call 800-624-3879 to check the status of a request.