



Actemra® Prior Authorization Request Form (Page 1 of 2)

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Member Information (required) Provider Information (required)

Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

Medication Information (required)

Medication Name:		Strength:	Dosage Form:
<input type="checkbox"/> Check if requesting brand		Directions for Use:	
<input type="checkbox"/> Check if request is for continuation of therapy			

Clinical Information (required)

Select the diagnosis below:

Moderately to severely active rheumatoid arthritis (RA)
 Active polyarticular juvenile idiopathic arthritis (pJIA)
 Active systemic juvenile idiopathic arthritis (sJIA)
 Temporal arteritis or giant cell arteritis (GCA)
 Systemic sclerosis-associated interstitial lung disease
 Other diagnosis: _____ ICD-10 Code(s): _____

Clinical information:
 Select if Actemra is prescribed by or in consultation with one of the following specialists:
 Allergist/Immunologist Pulmonologist Rheumatologist Other _____
 Will Actemra be used in combination with another biologic agent or targeted immunomodulator? Yes No

For moderately to severely active rheumatoid arthritis (RA), also answer the following:
 Has the patient had an inadequate response to, intolerance to, or contraindication to one or more non-biologic disease modifying anti-rheumatic drugs (DMARDs)? Yes No List _____

For active polyarticular juvenile idiopathic arthritis (pJIA), also answer the following:
 Has the patient had an inadequate response to, intolerance to, or contraindication to one or more non-biologic disease modifying anti-rheumatic drugs (DMARDs)? Yes No List _____

For active systemic juvenile idiopathic arthritis (sJIA), also answer the following:
 Has the patient had an inadequate response or intolerance to at least one oral systemic agent [i.e., non-steroidal anti-inflammatory drugs (NSAIDs), corticosteroid]? Yes No List _____

For temporal arteritis or giant cell arteritis (GCA), also answer the following:
 Has the patient had an inadequate response to, intolerance to, or contraindication to oral or parenteral corticosteroid? Yes No List _____

Quantity limit requests:
 What is the quantity requested per TREATMENT? _____ syringe every _____ weeks

What is the reason for exceeding the plan limitations?

Titration or loading dose purposes
 Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)
 Requested strength/dose is not commercially available
 Other: _____

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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-855-401-4262.
This form may be used for non-urgent requests and faxed to 1-844-403-1029.