

## Inflectra® & Remicade® Coverage Determination Request Form (Page 1 of 2)

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Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		Office Contact:
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name: Select one of the following: <input type="checkbox"/> Request is for <b>GENERIC</b> <input type="checkbox"/> Request is for <b>BRAND</b> (unable to take the generic)			Strength:		Dosage Form:
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>			Directions for Use:		
Clinical Information <small>(required)</small>					
<b>Select the Type of Coverage Determination Requested:</b>					
<input type="checkbox"/> <b>Prior Authorization</b> - Request is for a drug that requires prior authorization under the plan.					
<b>Select the diagnosis below:</b>					
<input type="checkbox"/> Ankylosing spondylitis		<input type="checkbox"/> Pulmonary sarcoidosis			
<input type="checkbox"/> Crohn's disease		<input type="checkbox"/> Pyoderma gangrenosum			
<input type="checkbox"/> Plaque psoriasis		<input type="checkbox"/> Rheumatoid arthritis			
<input type="checkbox"/> Polyarticular juvenile idiopathic arthritis		<input type="checkbox"/> Ulcerative colitis			
<input type="checkbox"/> Psoriatic arthritis		<input type="checkbox"/> Wegener's granulomatosis			
<input type="checkbox"/> Other diagnosis: _____		ICD-10 Code(s): _____			
<b>Clinical Information:</b>					
Will the patient receive concurrent therapy with other biologic disease modifying anti-rheumatic drugs (DMARDs) or other tumor necrosis factor antagonists? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is there documentation the patient has been evaluated for active or latent tuberculosis (TB)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For ankylosing spondylitis, also answer the following:</b>					
Does the patient have active, moderate to severe disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is there documentation that the patient has had an inadequate response or inability to tolerate at least one other treatment such as NSAIDs, COX-2 inhibitors, or methotrexate? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For chronic pulmonary sarcoidosis, also answer the following:</b>					
Is there documentation that the patient has had history of trial and failure, or intolerance to a 3-month trial of corticosteroids and immunosuppressive agents? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For Crohn's disease, also answer the following:</b>					
Is there documentation that the patient has had an inadequate response or inability to tolerate at least one conventional treatment (e.g., corticosteroids, aminosalicylates, immunomodulators)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>For plaque psoriasis, also answer the following:</b>					
Does the patient have chronic, severe (i.e., extensive and/or disabling) disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is the patient a candidate for systemic therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is there documentation that the patient has had an inadequate response or is unable to tolerate other systemic therapies? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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Office use only: Inflectra-Remicade\_FSPartD\_2018Jul-W



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**For polyarticular juvenile idiopathic arthritis, also answer the following:**

Does the patient have evidence of active disease?  Yes  No

Is there documentation the patient has had failure or intolerance to a 3-month trial of an FDA-approved biologic disease modifying anti-rheumatic drug (DMARD)?  Yes  No

**For psoriatic arthritis, also answer the following:**

Does the patient have active disease?  Yes  No

Is there documentation that the patient has an inability to tolerate or inadequate response to at least one DMARD?  Yes  No

**For rheumatoid arthritis, also answer the following:**

Will the requested medication be used in combination with methotrexate?  Yes  No

If "no" to the above question, will the requested medication be used as monotherapy due to an intolerance or contraindication to methotrexate?  Yes  No

Is there documentation that the patient has had an inadequate response to any DMARD (e.g. sulfasalazine, azathioprine, cyclophosphamide, cyclosporine, methotrexate, and/or other anti-tumor necrosis factor agents)?  Yes  No

**For ulcerative colitis, also answer the following:**

Does the patient have moderate to severe disease?  Yes  No

Is there documentation that the patient has had an inadequate response or inability to tolerate at least one conventional treatment (e.g., corticosteroids, aminosalicylates, immunomodulators)?  Yes  No

**For pediatric patients, also answer the following:**

Does the patient have moderately to severely active disease?  Yes  No

Does the patient have documentation of an inadequate response to conventional therapies?  Yes  No

**For Wegener's granulomatosis, also answer the following:**

Does the patient have evidence of severe active disease?  Yes  No

Is there documentation that the patient has had an inadequate response or inability to tolerate corticosteroids and immunosuppressant agents?  Yes  No

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.