



Please complete ALL information below and fax your request to 1-888-671-5285

## Enbrel® 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 <b>generic substitution</b> is acceptable			Directions for Use:		
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>					
Clinical Information (required)					
<b>Select the diagnosis below:</b>					
<input type="checkbox"/> Ankylosing spondylitis (for example: M45.0, M45.9)					
<input type="checkbox"/> Plaque psoriasis (for example: psoriasis vulgaris, psoriasis; L40.0, L40.8, L40.9)					
<input type="checkbox"/> Polyarticular juvenile idiopathic arthritis (for example: M08.09, M08.40)					
<input type="checkbox"/> Psoriatic arthritis (for example: arthropathic psoriasis; L40.5, L40.59)					
<input type="checkbox"/> Rheumatoid arthritis (for example: M05.79, M06.9)					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<b>Clinical Information:</b>					
Is this request for continuation of prior Enbrel therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if Enbrel is prescribed by or in consultation with one of the following specialists:					
<input type="checkbox"/> Dermatologist		<input type="checkbox"/> Rheumatologist			
<b>For ankylosing spondylitis, also answer the following:</b>					
Does the patient have active disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient had trial and failure, contraindication, or intolerance to <b>TWO</b> non-steroidal anti-inflammatory drugs (NSAIDs)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if the patient has had trial and failure, contraindication, or intolerance to the following, or attestation demonstrating a trial may be inappropriate:					
<input type="checkbox"/> Cimzia (certolizumab pegol)		<input type="checkbox"/> Humira (adalimumab)		<input type="checkbox"/> Simponi (golimumab) or Simponi Aria (golimumab IV)	
<b>For plaque psoriasis, also answer the following:</b>					
Does the patient have moderate to severe chronic disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Select if the patient has had trial and failure, contraindication, or intolerance to the following:					
<input type="checkbox"/> Cimzia (certolizumab pegol)		<input type="checkbox"/> Humira (adalimumab)		<input type="checkbox"/> Tremfya (guselkumab)	
<input type="checkbox"/> Cosentyx (secukinumab)		<input type="checkbox"/> Skyrizi (risankizumab)		<input type="checkbox"/> Stelara (ustekinumab)	
<b>For polyarticular juvenile idiopathic arthritis, also answer the following:</b>					
Does the patient have moderately to severely active disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient had trial and failure, contraindication, or intolerance to <b>ONE</b> of the following non-biologic modifying anti-rheumatic drugs (DMARDs): Arava (lefunomide) <b>OR</b> Rheumatrex/Trexall (methotrexate)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient had trial and failure, contraindication, or intolerance to Humira (adalimumab)? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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**Enbrel® Prior Authorization Request Form (Page 2 of 2)**  
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**For psoriatic arthritis, also answer the following:**

Does the patient have active disease?  Yes  No

Select if the patient has had trial and failure, contraindication, or intolerance to the following:

- |  |   |   |
|--|---|---|
| <input type="checkbox"/> Cimzia (certolizumab pegol) | <input type="checkbox"/> Humira (adalimumab)                                | <input type="checkbox"/> Stelara (ustekinumab)                                |
| <input type="checkbox"/> Cosentyx (secukinumab)      | <input type="checkbox"/> Simponi (golimumab) or Simponi Aria (golimumab IV) | <input type="checkbox"/> Xeljanz (tofacitinib) or Xeljanz XR (tofacitinib ER) |

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

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

Has the patient had trial and failure, contraindication, or intolerance to **ONE** non-biologic disease modifying anti-rheumatic drug (DMARD) (e.g., Rheumatrex/Trexall [methotrexate], Arava [leflunomide], Azulfidine [sulfasalazine])?  Yes  No

Select if the patient has had trial and failure, contraindication, or intolerance to the following, or attestation demonstrating a trial may be Inappropriate:

- |  |  |   |
|--|--|---|
| <input type="checkbox"/> Cimzia (certolizumab pegol) | <input type="checkbox"/> Humira (adalimumab) | <input type="checkbox"/> Simponi (golimumab) or Simponi Aria (golimumab IV) |
|--|--|---|

Select if the patient has had trial and failure, contraindication, or intolerance to the following:

- |  |   |
|--|---|
| <input type="checkbox"/> Actemra (tocilizumab) | <input type="checkbox"/> Xeljanz (tofacitinib) or Xeljanz XR (tofacitinib ER) |
|--|---|

**Reauthorization:**

**If this is a reauthorization request, answer the following:**

Is there documentation the patient has had a positive clinical response to Enbrel therapy?  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?**

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Please note: This request may be denied unless all required information is received.