



OptumRx has partnered with CoverMyMeds to receive prior authorization requests, saving you time and often delivering real-time determinations.

Visit go.covermymeds.com/OptumRx to begin using this free service.

Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

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

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

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)							
<p>Continuation of therapy: Is this a continuation of prior therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No If "yes" to the above question, please submit documentation (e.g., medical records, chart notes, pharmacy claims) or provide the dates, duration, and previous regimen used below: (REQUIRED)</p> <hr/>							
<p>Select the diagnosis below:</p> <table style="width: 100%; border: none;"> <tr> <td style="width: 50%; vertical-align: top;"> <input type="checkbox"/> Active ankylosing spondylitis <input type="checkbox"/> Active psoriatic arthritis <input type="checkbox"/> Moderate to severe chronic plaque psoriasis <input type="checkbox"/> Moderate to severe hidradenitis suppurativa <input type="checkbox"/> Moderately to severely active Crohn's disease <input type="checkbox"/> Other diagnosis: _____ </td> <td style="width: 50%; vertical-align: top;"> <input type="checkbox"/> Moderately to severely active juvenile idiopathic arthritis <input type="checkbox"/> Moderately to severely active rheumatoid arthritis <input type="checkbox"/> Moderately to severely active ulcerative colitis <input type="checkbox"/> Non-infectious intermediate, posterior and panuveitis </td> </tr> </table> <p>ICD-10 Code(s): _____</p>						<input type="checkbox"/> Active ankylosing spondylitis <input type="checkbox"/> Active psoriatic arthritis <input type="checkbox"/> Moderate to severe chronic plaque psoriasis <input type="checkbox"/> Moderate to severe hidradenitis suppurativa <input type="checkbox"/> Moderately to severely active Crohn's disease <input type="checkbox"/> Other diagnosis: _____	<input type="checkbox"/> Moderately to severely active juvenile idiopathic arthritis <input type="checkbox"/> Moderately to severely active rheumatoid arthritis <input type="checkbox"/> Moderately to severely active ulcerative colitis <input type="checkbox"/> Non-infectious intermediate, posterior and panuveitis
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<p>Prescriber's Specialty: Is Humira prescribed by a rheumatologist, dermatologist, or gastroenterologist? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>							
<p>Clinical Information: Select if the prescriber will submit a statement or chart notes documenting the patient has experienced the following: <input type="checkbox"/> Failure on methotrexate AND one other disease-modifying antirheumatic drug (DMARD) <input type="checkbox"/> Failure of topical steroids AND PUVA <input type="checkbox"/> Failure of systemic steroids AND a disease-modifying antirheumatic drug (DMARD) (mesalamine may be considered a DMARD) Is there presence of contraindications or absolute drug interactions with existing therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>							
<p>Reauthorization: If this is a reauthorization request, answer the following question: Does the patient show symptom reduction from baseline? <input type="checkbox"/> Yes <input type="checkbox"/> No</p>							



Humira® Prior Authorization Request Form (Page 2 of 2)
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Quantity Limit Requests:

What is the quantity requested per DAY? _____

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: _____

For continuation of existing therapy, also answer the following:

Would sudden discontinuation of the dose trigger withdrawal symptoms? Yes No

Would discontinuation of the dose be unsafe for the patient and their condition may worsen or exacerbate? Yes No

Is the prescribing provider attempting to taper or reduce the dose necessary? 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.
For urgent or expedited requests please call 1-855-297-2870.
This form may be used for non-urgent requests and faxed to 1-844-403-1029.