

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

Fax to 1-844-403-1029 Mon-Sat: 7am to 7pm Central

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:		Zip:		
Medication Information (required)							
Medication Name:		Strength: Dosage Form:					
			Directions for Use:				
☐ Check if requesting brand☐ Check if request is for continuation of therapy			Directions for osc.				
Clinical Information (required)							
Select the diagnosis below:							
□ Active ankylosing spondylitis							
□ Active psoriatic arthritis (PsA)□ Moderate to severe chronic plaque psoriasis							
Moderate to severe hidradenitis suppurativa (e.g., Hurley Stage II or III)							
Moderately to severely active Crohn's disease Mederately to severely active polyarticular invented idionathic arthritis (IIA)							
 Moderately to severely active polyarticular juvenile idiopathic arthritis (JIA) Moderately to severely active rheumatoid arthritis (RA) 							
Moderately to severely active medinatord artiflus (RA) Moderately to severely active ulcerative colitis							
□ Non-infectious uveitis							
☐ Other diagnosis:	ICD-10 Code(s):						
Clinical information:							
Select if the requested medication is prescribed by or in consultation with one of the following specialists: □ Dermatologist □ Gastroenterologist □ Ophthalmologist □ Rheumatologist □ Other							
Will the requested medication be used in combination with another biologic agent or targeted immunomodulator? Yes No							
If requesting a citrate-	free product, has the pa	tient tried citrate produc	t first? Yes No	When:			
For active ankylosin	g spondylitis (AS), als	o answer the following	:				
Has the patient had ar (NSAIDs)? ☐ Yes ☐	n inadequate response t No List	o, intolerance to, or con	traindication to one or n	nore non-ste	eroidal anti-iı	nflammatory drugs	
For active psoriatic a	arthritis (PsA), also an	swer the following:					
Has the patient had an inadequate response to, intolerance to, or contraindication to methotrexate? No							
For moderate to seve	ere chronic plaque pso	oriasis (PsO), also ans	wer the following:				
	n inadequate response t by or one or more oral sy roid)? □ Yes □ No						
	ere hidradenitis suppu		-				
Has the patient had an inadequate response to, intolerance to, or contraindication to one or more of the following: oral or topical antibiotic							



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For moderately to severely active Crohn's disease, also answer the following:
Has the patient had an inadequate response to, intolerance to, or contraindication to one or more conventional therapies: azathioprine, 6-mercaptopurine, methotrexate, corticosteroids (e.g., prednisone, methylprednisolone) Tes No List
For moderately to severely 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 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 moderately to severely active ulcerative colitis, also answer the following:
Has the patient had an inadequate response to, intolerance to, or contraindication to conventional therapy with one or more of the following: corticosteroids (i.e., prednisone, methylprednisolone), 5-ASAs (i.e., mesalamine, sulfasalazine, balsalazide, olsalazine), non-biologic DMARDs (i.e., azathioprine, methotrexate, mercaptopurine)? Yes No List
For non-infectious uveitis, also answer the following:
Has the patient had an inadequate response to, intolerance to, or contraindication to one or more of the following: methotrexate, mycophenolate, azathioprine, cyclosporine, tacrolimus, cyclophosphamide, oral/injectable steroid therapy? 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:
Other.
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.