

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

Fax to 1-844-403-1029

Mon-Sat: 7am to 7pm Central

## Hyrimoz® Prior Authorization Request Form (Page 1 of 2) DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

DO NOT COPT FOR FOTORE USE. FORMS ARE UPDATED FREQUENTET AND MAT 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:		
☐ Check if requesting <b>brand</b>			Directions for Use:	rections for Use:			
☐ Check if request is for <b>continuation of therapy</b>							
Clinical Information (required)							
Select the diagnosis below:							
□ Active ankylosing spondylitis							
□ Active psoriatic arthritis (PsA)							
☐ Moderate to severe chronic plaque psoriasis							
Moderate to severely active Crohn's disease							
·							
☐ Moderately to severely active polyarticular juvenile idiopathic arthritis (JIA)							
☐ Moderately to severely active rheumatoid arthritis (RA)							
☐ Moderately to severely active ulcerative colitis							
☐ Moderately to severely active ulcerative colitis							
☐ Hidradenitis Suppurativa							
□ Other diagnosis: ICD-10 C							
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							
Justification for the ulf non-preferred agent alternative:	use of a non-preferred is medically necessary	product (Hymiroz) ove or required, provide a bi	er a preferred product rief summary for use of t	( <b>Humira):</b> he non-pref	erred agent o	over a preferred	
For active ankylosing spondylitis (AS), also answer the following:							
Has the patient had ar (NSAIDs)? ☐ Yes ☐		to, intolerance to, or con	traindication to one or m	ore non-ste	roidal anti-in	ıflammatory drugs	
For active psoriatic a	arthritis (PsA), also ans	swer the following:					
Has the patient had ar	า inadequate response t	o, intolerance to, or con	traindication to methotre	xate? 🛚 Ye	s 🗆 No		
For moderate to severe chronic plaque psoriasis (PsO), also answer the following:							
Has the patient had an inadequate response to, intolerance to, or contraindication to conventional therapy with at least one of the following: phototherapy or one or more oral systemic treatments (i.e., methotrexate, cyclosporine, acitretin, sulfasalazine, calcipotriene, tazarotene, corticosteroid)?   Yes  No List							



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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 immunosuppressive agents (e.g., azathioprine, mercaptopurine, methotrexate)?   Yes 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 moderate to severe hidradenitis suppurativa, also answer the following:
Has the patient had an inadequate response to, intolerance to, or contraindication to one or more of the following: oral or topical antibiotic therapy OR oral retinoid therapy, dapsone, or acitretin? <b>□</b> Yes <b>□</b> 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?   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:  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.