

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

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

AbriladaTM Prior Authorization Request Form (Page 1 of 2)

	Provider Information (required)						
Member Information (required) Member Name:			Provider Name:				
Insurance ID#:			NPI#:	PI#:		Specialty:	
Date of Birth:			Office Phone:				
Street Address:			Office Fax:				
City: State: Zip:			Office Street Address:				
Phone:		City: State: Zip:					
				'			
		Medication Info	ormation (required)				
Medication Name:			Strength:	Dosage Form:			
☐ Check if requesting brand			Directions for Use:				
☐ Check if request is	for continuation of the	rapy					
		Clinical Inform	mation (required)				
□ Moderately to sevee □ Moderately to sevee □ Moderately to sevee □ Moderately to sevee □ Hidradenitis Support □ Uveitis □ Other diagnosis: Clinical information: Select if the requested □ Dermatologist Will the requested me	spondylitis chritis (PsA) e chronic plaque psorias erely active Crohn's dise erely active polyarticular erely active rheumatoid a erely active ulcerative co urativa d medication is prescribe G Gastroenterologist edication be used in com	iase juvenile idiopathic arthranthritis (RA) plitis ed by or in consultation of the consultation with another bid	with one of the following t □ Rheumatologist ologic agent or targeted	☐ Othe immunomod	er	Yes □ No	
Justification for the	use of a non-preferred	product (Abrilada) ov	er a preferred product	(Humira):			
	is medically necessary		rief summary for use of	the non-pref	erred agent	over a preferred	
	g spondylitis (AS), also n inadequate response t I No List			nore non-ste	eroidal anti-i	nflammatory drugs	
-	arthritis (PsA), also an	•					
· · · · · · · · · · · · · · · · · · ·	n inadequate response t			exate? u Y	es u no		
Has the patient had an following: phototherap	ere chronic plaque pson inadequate response to by or one or more oral syroid)?	to, intolerance to, or convstemic treatments (i.e.,	traindication to conventi				



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

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? □ 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? 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.					