Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME:		MEMBER'S FIRST NAME:		
that is important for the re		lab data, to support the	y. Attach any additional documene authorization request). Inform	
			□ U	RGENT
MEMBER INFORMATION	ON			
LAST NAME:	AST NAME:			
PHONE NUMBER:		DATE OF BIRTH:		
STREET ADDRESS:		-		
CITY:		STATE:	ZIP CODE:	
PATIENT INSURANCE	ID NUMBER:			
☐ MALE ☐ FEMALE	HEIGHT (IN/CM):	_ WEIGHT (LB/KG)	ALLERGIES:	
FOLLOWING LINK: PRI		HIS REQUEST WHICH	H CAN BE FOUND AT THE	
AUTHORIZED REPRESI	ENTATIVE'S PHONE NU	MBER:		
PRESCRIBER INFORM	ATION			
LAST NAME:		FIRST NAME:		
PRESCRIBER SPECIA	LTY:	EMAIL ADDRESS:		
NPI NUMBER:		DEA NUMBER:		
PHONE NUMBER:		FAX NUMBER:		
STREET ADDRESS:		I		
CITY:		STATE:	ZIP CODE:	
REQUESTER (if different than prescriber):		OFFICE CONTACT PERSON:		
MEDICATION OR MED	ICAL DISPENSING INFO	RMATION		
MEDICATION NAME:				
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REF	QUANTITY:	
☐ NEW THERAPY	RENEWAL IF		HERAPY INITIATED:	
DURATION OF THERA	PY (SPECIFIC DATES):			
Continued on next page				

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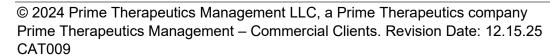
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MEMBER'S LAST NAME:	MEMBER'S FIRST N	IAME:				
1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?						
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:				
2. LIST DIAGNOSES:		ICD-10:				
Rheumatoid arthritis(RA) Polyarticular Juvenile idiopath Moderate to severe Atopic Derma Psoriatic Arthritis (PsA) Ulcerative Colitis(UC) Crohn's Disease(CD) Ankylosing Spondylitis Non-radiographic Axial Spondyla Atopic Dermatitis Giant Cell Arteritis Other diagnosis:	titis (AD)					
-	• •					
3. REQUIRED CLINICAL INFORMATO SUPPORT A PRIOR AUTHORIA	ATION: PLEASE PROVIDE ALL REL ZATION.	LEVANT CLINICAL INFORMATION				
	in combination with a clinical trial	? 🗌 Yes 🔲 No				
Is the prescriber a rheumatologis Is the prescriber a gastroenterologist Is the prescriber a dermatologist Will the patient use drug in comb	gist? □ Yes □ No	onse modifier or				
immunomodulatory agent? Ye						
•	nadequate response to a three mo s □ No Please submit documenta					
Does patient have a absolute con ☐ No Please submit documentation		Humira- adalimumab-aacf? □ Yes				
	nadequate response to a trial of th s □ No Please submit documenta					
Does patient have a absolute con	traindication to the biosimilar for A	Actemra, Tyenne(tocilizumab-				

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MEMBER'S LAST NAME: MEMBER'S FIRST NAME:
Has the patient tried and had an inadequate response to a 4- month trial of the biosimilar for Stelara-
Otulfi(ustekinumb-aauz)? Yes No Please submit documentation.
Does patient have a absolute contraindication to the biosimilar for Stelara-Otulfi(ustekinumb-aauz)? □ Yes □ No Please submit documentation.
For diagnosis of Rheumatoid Arthritis only:
Does the patient have a diagnosis of moderately to severely active rheumatoid arthritis? \hdots Yes \hdots No
Has the patient had a trial of methotrexate or another oral non-biologic disease modifying anti-rheumatic agent (DMARD) such as Imuran, Ridaura, Plaquenil, sulfasalazine or Arava? No Please submit documentation with dates of service.
Does patient have chronic alcohol abuse/alcoholism, chronic liver disease such as chronic hepatitis, fatty liver, nonalcoholic steatohepatitis/NASH, elevated liver enzymes)? □ Yes □ No Please submit documentation.
For diagnosis of Polyarticular Juvenile idiopathic arthritis(pJIA):
Has patient tried and failed previous therapy with oral disease modifying anti-rheumatic agents (DMARDs) [e.g. for JIA: methotrexate or sulfasalazine or leflunamide]? \[\text{ Yes } \text{ No } Please submit documentation.} \]
For renewal only: Does the patient continue to have a positive clinical response and remission of disease maintained with continued use of the medication? Yes No Please submit chart documentation.
Will the patient use drug in combination with another biologic or immunomodulatory agent? $\ \square$ Yes $\ \square$ No
Is the prescriber a rheumatologist? □ Yes □ No Is the prescriber a gastroenterologist? □ Yes □ No
For diagnosis of <u>Atopic Dermatitis</u> only:
Has the patient had the diagnosis of atopic dermatitis for at least 12 months? □ Yes □ No *Please submit documentation.
Does the patient have atopic dermatitis on at least 10% or more of their body surface area? — Yes — No *Please submit documentation.





MEMBER'S LAST NAME: MEMBER'S FIRST NAME:
Has the patient tried at least two different topical steroids? □ Yes □ No *Please submit documentation.
If patient has not had at least 2 different topical steroids, has the patient tried at least one topical steroid AND one topical calcineurin inhibitor (tacrolimus or pimecrolimus)? — Yes — No *Please submit documentation.
If patient has not had at least 2 different topical steroids, has the patient tried at least one topical steroid AND Eucrisa(crisaborole)? □ Yes □ No *Please submit documentation.
If patient has not had at least 2 different topical steroids, has the patient tried at least one topical steroid AND Zoryve(roflumilast)? Yes No *Please submit documentation.
If patient has not had at least 2 different topical steroids, has the patient tried at least one topical steroid AND Vtama(tapinarof)? □ Yes □ No *Please submit documentation.
Has patient tried and failed a 3-month trial of Dupixent(dupilumab)? □ Yes □ No *Please submit documentation.
Has patient tried and failed a 3-month trial of Adbry(tralokinumab-ldrm)? Yes No *Please submit documentation.
Has patient tried and failed a 3-month trial of Cibinqo(abrocitinib)? □ Yes □ No *Please submit documentation.
Will RinvoqER(upadacitinib) be used in combination with Cibinqo(abrocitinib), Olumiant(baracitinib), Opzelura(ruxolitinib), Dupixent(dupilumab), Adbry(tralokinumab), Xolair(omalizumab), Nucala(mepolizumab) or Fasenra(benralizumab? Yes No
For renewal only: Does the patient continue to have a positive clinical response and remission of disease maintained with continued use of the medication? Yes No Please submit chart documentation.
Is the patient currently being treated with another biologic or immunomodulatory agent? $\ \square$ Yes $\ \square$ No
Is RinvoqER(upadacitinib) being used in combination with Cibinqo(abrocitinib), Olumiant(baracitinib), Opzelura(ruxolitinib), Dupixent(dupilumab), Adbry(tralokinumab), Xolair(omalizumab), Nucala(mepolizumab) or Fasenra(benralizumab? ☐ Yes ☐ No
Is the prescriber a dermatologist or allergist? □ Yes □ No
For diagnosis of <u>Psoriatic Arthritis</u> only:



MEMBER'S LAST NAME: MEM	BER'S FIRST NAME:
Does the patient have documented moderately to sessibility documentation	verely active disease? Yes No Please
Has the patient had a trial and failed previous therapy agents (DMARDs, e.g., methotrexate, sulfasalazine (A cyclosporine)? ☐ Yes ☐ No Please submit documentation with dates of service.	
For renewal only: Does the patient continue to have a positive clinical with continued use of the medication? Yes No	
Is the patient currently being treated with another bid	ologic or immunomodulatory agent? □ Yes □
Is the prescriber a rheumatologist or dermatologist?	□ Yes □ No
For diagnosis of <u>Ulcerative Colitis and Crohn's Diseased</u> Is prescriber a gastroenterologist? Yes No Has patient tried and failed at least one of the following azathioprine, and/or 6-mercaptopurine? Yes No	ng three therapies: corticosteroids,
For renewal only: Does the patient continue to have a positive clinical with continued use of the medication? Yes No	
Is the patient currently being treated with another bid	ologic or immunomodulatory agent? □ Yes □
Is the prescriber a rheumatologist or gastroenterolog	gist? □ Yes □ No
For diagnosis of <u>Ankylosing Spondylitis</u> only:	
Is the prescriber a rheumatologist? ☐ Yes ☐ No	
Does the patient have documented active disease?	Yes No Please submit documentation
Has the patient had a trial and failed previous therap inflammatory agents (NSAIDS), unless use is contrai Please submit documentation with dates of service.	
For renewal only: Does the patient continue to have a positive clinical with continued use of the medication? Yes No	



MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:
Is the patient currently being treated with an No	other biologic or immunomodulatory agent? □ Yes □
Is the prescriber a rheumatologist? □ Yes	□ No
	endyloarthritis only: ammation by presence of sacroiliitis on MRI imaging vel? □ Yes □ No Please submit imaging and/or lab
Has patient had an inadequate response to a submit documentation.	it least two different NSAIDs? □ Yes □ No Please
	dication or intolerance to a 3-month trial with at least ibitor or an IL-17 inhibitor? Yes No Please submit
	clinical response and remission of disease maintained No <i>Please submit chart documentation</i> .
Is the patient currently being treated with an No	other biologic or immunomodulatory agent? □ Yes □
Is the prescriber a rheumatologist? □ Yes	□ No
For Giant Cell Arteritis: Does patient have a diagnosis of Giant Cell Adocumentation	Arteritis? □ Yes □ No <i>Please submit chart</i>
Is patient currently on a tapering dose of cor Will Rinvoq be used as monotherapy? Yes	
• • • • • • • • • • • • • • • • • • •	clinical response and remission of disease maintained No <i>Please submit chart documentation</i> .
Is the patient currently being treated with an No	other biologic or immunomodulatory agent? □ Yes □
Is the prescriber a rheumatologist? □ Yes	□ No
Are there any other comments, diagnoses, s information the physician feels is important	ymptoms, medications tried or failed, and/or any other to this review?





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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:	
Please note: Not all drugs/diagnosis are covered	on all plans. This request may be denied unless all	
required information is received.		
ATTESTATION: I attest the information provided	is true and accurate to the best of my knowledge. I	
understand that the Health Plan, insurer, Medical	Group or its designees may perform a routine audit and	
request the medical information necessary to verif	y the accuracy of the information reported on this form.	
	,	
Prescriber Signature or Electronic I.D. Verificat	tion: Date:	
U		
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MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program

Attn: CP-4201 P.O. Box 64811 St. Paul, MN 55164-0811 **Phone**: 877-228-7909

