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

MEMBER'S LAST NAME:		_ MEMBER'S FIRST NAME: _		
	g., chart notes or lab data, to	tely and legibly. Attach any addi support the authorization reque		
			☐ URGENT	
MEMBER INFORMATION				
LAST NAME:		FIRST NAME:		
PHONE NUMBER:		DATE OF BIRTH:		
STREET ADDRESS:				
CITY:		STATE: ZIP CODE	::	
PATIENT INSURANCE ID NU	UMBER:	I		
FOLLOWING LINK: PRIMETHERAPEUTICS.CC PATIENT'S AUTHORIZED REI	PRESENTATIVE (IF APPLICABI	SCLOSURE AUTHORIZATION FORM WITH THIS RE		
PRESCRIBER INFORMATIO	N			
LAST NAME:		FIRST NAME:	FIRST NAME:	
PRESCRIBER SPECIALTY:		EMAIL ADDRESS:	EMAIL ADDRESS:	
NPI NUMBER:		DEA NUMBER:	DEA NUMBER:	
PHONE NUMBER:		FAX NUMBER:	FAX NUMBER:	
STREET ADDRESS:				
CITY:	CITY:		E:	
REQUESTOR (if different than prescriber):		OFFICE CONTACT PERSON:		
MEDICATION OR MEDICA	L DISPENSING INFORMATION			
MEDICATION NAME:				
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:	
■ NEW THERAPY	RENEWAL	IF RENEWAL: DATE THERAP	Y INITIATED:	
DURATION OF THERAPY (SE	PECIFIC DATES):			

Continued on next page.



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MEMBER'S LAST NAME:	R'S LAST NAME: MEMBER'S FIRST NAME:			
1. HAS THE PATIENT TRIED ANY OTHE	R MEDICATIONS FOR THIS CONDITION?	YES (if yes, complete below) NO		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:		
2. LIST DIAGNOSES:		ICD-10:		
 □ Plaque psoriasis □ Psoriatic arthritis □ Ankylosing spondylitis □ Nonradiographic Axial Spondyloarthritis: 	ICD-10 Code(s):			
3. REQUIRED CLINICAL INFORMATION PRIOR AUTHORIZATION.	: PLEASE PROVIDE ALL RELEVANT CLINIC	AL INFORMATION TO SUPPORT A		
Clinical Information: Initial Request: Is drug being used as part of a clinical	trial? □ Yes □ No			
Is the patient on concurrent treatment Yes $\ \square$ No	t with another biologic response modifi	ier or immunomodulatory agent? $\ \square$		
Has the patient tried and had an inade adalimumab-aacf?* *Must provide documentation, included.*	equate response to a three month trial of trial of trial dates.	of the biosimilar for Humira-		
Is Taltz prescribed by a dermatologist Is Taltz prescribed by a rheumatologis				
For Request Plaque Psoriasis, also an	swer the following: ing greater than or equal to 10% of thei	r hody surface area (BSA)? □ Yes □ No		
	ng less than 10% of BSA with involveme			
Has the patient had an inadequate restazarotene)? ☐ Yes ☐ No	sponse to a topical therapy (e.g., cortico	osteroids, anthralin, calcipotriene		
	ase provide supporting documentation,			
Select if the patient has had an inaded Psoralens with UVA light (PUVA) UVB with coal tar	quate response to previous treatment w	vith the following phototherapies:		
Please provide supporting docume	ntation, including which agent(s) have I	been tried and trial dates:		



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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:	
□ Acitretin□ Methotrexate□ Cyclosporine	ad an inadequate response to the following oral systemic therapies: mentation, including which agent(s) have been tried and trial dates:	
AcitretinMethotrexateCyclosporine	ication to ALL of the follow ing oral systemic therapies:* of the contraindications to all three drugs.	
For Request of Ankylosing Spondyling Has patient had a trial of at least two Yes No	tis, also answer the following: vo (2) NSAIDs OR has patient had a trial of one NSAID AND methotrexate?	
For Request of Nonradiographic Axi Did the patient's back pain begin be	ial Spondyloarthritis, also answer the following: efore age 45 years? Yes No	
Does the patient have objective sign ☐ Yes ☐ No Please submit MRI repo	ns of inflammation by presence of sacroiliitis on MRI? ort.	
Does the patient have objective sign ☐ Yes ☐ No Please submit lab report	ns of inflammation by presence of an elevated C-reactive protein level?	
Has the patient had an inadequate □ Yes □ No Please submit documen	response to at least two different NSAIDs for at least 4 weeks? tation.	
Is the patient intolerant of NSAIDs?	□ Yes □ No Please submit documentation.	
Does the patient have radiographic imaging (x-ray) report.	sacroiliitis (per 1984 modified New York criteria)? □ Yes □ No <i>Please submit</i>	
Renewal Requests: Is Taltz prescribed by a dermatologi	st? 🗆 Yes 🗆 No	
Is Taltz prescribed by a rheumatolog	gist? □ Yes □ No	
Is patient continuing to respond to	therapy? Yes No Please submit documentation.	
Will patient use requested medication in combination with another biologic response modifier or immunomodulatory agent? $\ \square$ Yes $\ \square$ No		



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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: Not all drugs/diagnosis are cover information is received.	red on all plans. This request may be denied unless all required	
·	ded is true and accurate to the best of my knowledge. I understand that designees may perform a routine audit and request the medical of the information reported on this form.	
Prescriber Signature or Electronic I.D. Verifica	ation: Date:	
you are not the intended recipient, you are hereby notifi	ng this transmission contain confidential health information that is legally privileged. If fied that any disclosure, copying, distribution, or action taken in reliance on the contents served this information in error, please notify the sender immediately (via return FAX) ments.	

FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program
Attn: CP - 4201
P.O. Box 64811
St. Paul, MN 55164-0811

St. Paul, IVIN 55164-0811

