Zeposia (ozanimod) Prior Authorization Request Form Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: ME	EMBER'S FIRST NAME:				
Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.					
MEMBER INFORMATION					
LAST NAME:	FIRST 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:					
If you are not the patient or the prescriber, you will need to submit a PHI Disclosure Authorization form with this request which can be found at the following link: primetherapeutics.com/NOPP					
PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE):					
AUTHORIZED REPRESENTATIVE'S PHONE NUMBER:					
PRESCRIBER INFORMATION					
LAST NAME:	FIRST NAME:				
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:				
NPI NUMBER:	DEA NUMBER:				
PHONE NUMBER:	FAX NUMBER:				
STREET ADDRESS:					
CITY:	STATE: ZIP CODE:				
REQUESTER (IF DIFFERENT THAN PRESCRIBER):	OFFICE CONTACT PERSON:				

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MEMBER'S LAST NAME: MEMBER'S FIRST NAME:						
MEDICATION OR MEDICAL DISPENSING INFORMATION						
MEDICATION NAME:						
DOSE/STRENGTH:	FREQU	ENCY:	LENGTH OF THERAPY/REFIL	LS:	QUANTITY:	
□ NEW THERAPY □ RENEWAL IF RENEWAL, DATE THERAPY INITIATED:						
DURATION OF THERAPY	<u> </u>	,				
1. HAS THE PATIENT T			ICATIONS FOR	THIS CC	NDITION?	
Medication/Therapy (Specify Drug Name And Dosage): Duration Of Therapy (Specify Dates):		Response/Reason For Failure/Allergy:				
2. LIST DIAGNOSES:				ICD-10:		
□ Clinically Isolated Syndrome (CIS) □ Relapsing Remitting Multiple Sclerosis (RRMS) □ Secondary Progressive Multiple Sclerosis (SPMS) □ Ulcerative Colitis (UC) □ Other diagnosis: ICD-10 CODE(S):						
3. REQUIRED CLINICA INFORMATION TO SUF				L RELE	VANT CLINICAL	
Clinical Information:						
Is drug going to be used in conjunction with a clinical trial? Yes No						
Initial Request for Multiple Sclerosis:						
Is the prescriber a neurologist? Yes No						
Has patient had a 3 month trial each of at least 2 of the following? Please provide documentation. dimethyl fumarate fingolimod glatiramer acetate teriflunomide						



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MEMBER'S LAST NAME: MEMBER'S FIRST NA	AME:
Initial Request for Ulcerative Colitis:	
Is the prescriber a gastroenterologist? Yes No	
Does patient have Crohn's disease or indeterminate colitis? Yes	No Please provide documentation.
Has patient tried and failed at least one of the following three therapies: and/or 6-mercaptopurine? Yes No Please submit chart documen	
Has patient tried and failed at least three months of the biosimilar Humi Yes No Please submit chart documentation.	ra-adalimumab-aacf?
Does patient have a absolute contraindication to the biosimilar for H Yes No Please submit documentation	lumira-adalimumab-aacf)? 🛘
Has the patient tried and had an inadequate response to a 4- month of the order of	
Does patient have a absolute contraindication to the biosimilar for S □ Yes □ No Please submit documentation.	Stelara- <u>Otulfi(ustekinumb-aauz</u>)?
Will patient use requested medication in combination with another bioloimmunomodulatory agent? ☐ Yes ☐ No	ogic response modifier or
Renewal Request:	
Is prescriber a neurologist? ☐ Yes ☐ No	
Is prescriber a gastroenterologist?	
Is patient continuing to have a positive response to therapy? $\ \square$ Yes $\ \square$ Please submit chart documentation.	No
Will patient use requested medication in combination with another biolommunomodulatory agent? \square Yes \square No	ogic response modifier or
Are there any other comments, diagnoses, symptoms, medications tried information the physician feels is important to this review?	d or failed, and/or any other



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MEMBER'S LAST NAME:	MEMBER'S FIRST NA	ME:		
Please note: Not all drugs/diagnosis are co required information is received.	vered on all plans. This reque	est may be denied unless all		
ATTESTATION: I attest the information prounderstand that the Health Plan, insurer, Marequest the medical information necessary	ledical Group or its designees	may perform a routine audit and		
Prescriber Signature or Electronic I.D. Verif	fication:	Date:		
CONFIDENTIALITY NOTICE: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is				

FAX THIS FORM TO: 800-424-7640

strictly prohibited. If you have received this information in error, please notify the sender immediately (via

return FAX) and arrange for the return or destruction of these documents.

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

