Movantik (naloxegol) Prior Authorization Request Form

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

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.

			URGENT	
MEMBER INFORMATION				
LAST NAME:		FIRST NAME:		
PHONE NUMBER:		DATE OF BIRTH:		
STREET ADDRESS:				
CITY:		STATE: ZIP CODE:		
PATIENT INSURANCE ID NUM	MBER:			
IF YOU ARE NOT THE PATIENT OR THE PRESCR FOLLOWING LINK: PRIMETHERAPEUTICS.COM	/NOPP	OSURE AUTHORIZATION FORM WITH TH	HIS REQUEST WHICH CAN BE FOUND AT THE	
PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE):				
PRESCRIBER INFORMATION				
LAST NAME:		FIRST NAME:		
PRESCRIBER SPECIALTY:		EMAIL ADDRESS:		
NPI NUMBER:		DEA NUMBER:		
PHONE NUMBER:		FAX NUMBER:		
STREET ADDRESS:				
CITY:		STATE: ZIP CODE:		
REQUESTOR (if different than prescriber):		OFFICE CONTACT PERSON:		
MEDICATION OR MEDICAL I	DISPENSING INFORMATION			
MEDICATION NAME:				
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:	
☐ NEW THERAPY	RENEWAL	IF RENEWAL: DATE THE	RAPY INITIATED:	
DURATION OF THERAPY (SPE	CIFIC DATES):			

Continued on next page.



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MEMBER'S LAST NAME:	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:	
□ Opioid-induced constipation□ Other diagnosis:	ICD-10:		
3. REQUIRED CLINICAL INFORMATION PRIOR AUTHORIZATION.	I: PLEASE PROVIDE ALL RELEVANT CLINIC.	AL INFORMATION TO SUPPORT A	
Clinical Information:			
Initial Request: Has the patient been on opioid therap	py for greater than one month (4 weeks))? □ Yes □ No	
Has the patient had a trial and failure	of laxatives? □ Yes □ No Please prov	vide documentation	
Did the patient have fewer than 3 spo ☐ Yes ☐ No	ontaneous bowel movements per week v	when off Movantik therapy?	
Will Movantik (naloxegol) be used in	combination with Relistor (methylnaltre	exone bromide)? 🗆 Yes 🗆 No	
Will Movantik (naloxegol) be used in	combination with Amitiza (lubiprostone)? □ Yes □ No	
Renewal Request: Does patient have opioid-induced cor	ostination? □ Ves. □ No		
•	re clinical response? Yes No Plea	se provide documenation.	
Are there any other comments, diagn physician feels is important to this rev	oses, symptoms, medications tried or fa view?	iled, and/or any other information the	
Please note: Not all drugs/diagnosis an information is received.	re covered on all plans. This request may	be denied unless all required	
ATTESTATION: I attest the informatio the Health Plan, insurer, Medical Grou	n provided is true and accurate to the be p or its designees may perform a routine curacy of the information reported on th	audit and request the medical	
Prescriber Signature or Electronic I.D.	Verification:	Date:	
you are not the intended recipient, you are her	companying this transmission contain confidential reby notified that any disclosure, copying, distribute have received this information in error, please no	tion, or action taken in reliance on the contents	



and arrange for the return or destruction of these documents.

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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

