Wakix (pitolisant) **Prior Authorization Request Form**

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

MEMBER'S LAST NAME: _____ MEMBER'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:	

MEDICATION	OD MEDICAL DI		
MEDICATION	OR MEDICAL DI	SPENSING IN	FORMATION

MEDICATION NAME: DOSE/STRENGTH: FREQUENCY: LENGTH OF QUANTITY: THERAPY/REFILLS: NEW THERAPY **IF RENEWAL:** DATE THERAPY INITIATED: DURATION OF THERAPY (SPECIFIC DATES): Continued on next page

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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:					
1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?						
YES (if yes, complete below) MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	NO DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:				
2. LIST DIAGNOSES:		ICD-10:				
Excessive Daytime Sleepiness (EDS) in <u>F</u>	Pediatric Patients with Narcolepsy ICD-10 Code(s):					
3. REQUIRED CLINICAL INFORMATO SUPPORT A PRIOR AUTHORIZ	ATION: PLEASE PROVIDE ALL REL ZATION.	EVANT CLINICAL INFORMATION				
Is patient going to be using drug in combination with a clinical trial? Yes No Does the patient have a diagnosis of narcolepsy? Yes No Please submit documentation.						
Does the patient have narcolepsy with	associated cataplexy? □ Yes □ No Ple	ease submit documentation.				
Has the patient had a sleep study to af	firm a diagnosis of narcolespy? D Yes	□ No Please submit documentation.				
Does the patient have an Epworth Sleepiness Scale (ESS) score of 14 or greater? Yes No Please submit documentation.						
Does the patient have another primary cause of excessive daytime sleepiness (such as a sleep Apnea Index \ge 10 per hour and/or an Apnea/Hypopnea Index \ge 15 per hour, periodic limbs movement (PLM) disorders as defined by a PLM arousal index (PLMAI) \ge 10 per hour, shift work, chronic sleep deprivation, circadian sleep wake rhythm disorder)? \Box Yes \Box No						
Will the patient take Wakix while also taking modafinil, armodafinil or any amphetamine product? 🛛 Yes 🗅 No						
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 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 verify the accuracy of the information reported on this form.						
Prescriber Signature or Electronic	c I.D. Verification:	Date:				

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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 **Phone**: 877-228-7909

