

Ubrelvy (ubrogepant)
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.

URGENT

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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MEDICATION OR MEDICAL DISPENSING INFORMATION

MEDICATION NAME:

DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
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NEW THERAPY RENEWAL IF RENEWAL, DATE THERAPY INITIATED:

DURATION OF THERAPY (SPECIFIC DATES):

1. HAS THE PATIENT TRIED ANY OTHER 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:
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2. LIST DIAGNOSES: ICD-10:

Acute Migraines

Other diagnosis: _____

ICD-10 CODE(S): _____

3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.

Clinical Information:

Is taking Ubrelvy(ubrogepant) going to be part of a clinical trial? Yes No

Will patient use in combination with another CGRP product for prevention and/or acute migraines?

Yes No --PATIENT IS NOT ALLOWED COMBINATION USE OF 2 CGRP'S OF ANY KIND—

If patient is using another CGRP product for acute or prevention of migraines and prescriber would like to use Ubrelvy(ubrogepant) for prevention or acute migraines, is prescriber ok with terming use of the other CGRP for acute or prevention of migraines in lieu of using NurtecODT(rimegepant) for prevention or acute migraines? Yes No *Any current active PA for an acute or prevention CGRP will be termed.

Has patient had acute migraines for at least 1 year? Yes No

Has patient received at least two different triptans and failed to have relief of their acute migraine episodes? Yes No *Please submit documentation.*

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Does the patient have an absolute contraindication to triptans: such as, ischemic heart disease, ischemic bowel disease, cerebrovascular disease, peripheral vascular disease, cardiac conduction pathway disorder, hemiplegic migraines, basilar migraines, or severe hepatic impairment? Yes No Please submit documentation.

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 I.D. Verification: _____ 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 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.

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