

# Renvela Powder (sevelamer carbonate pwdr)

## 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](http://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 OR MEDICAL DISPENSING INFORMATION			
MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
<input type="checkbox"/> NEW THERAPY	<input type="checkbox"/> RENEWAL	IF RENEWAL: DATE THERAPY INITIATED:	
DURATION OF THERAPY (SPECIFIC DATES):			

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### 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:

### 2. LIST DIAGNOSES:

ICD-10:

Post-gastric bypass surgery

Stage 3 to 6 chronic kidney disease (CKD)

Other diagnosis: \_\_\_\_\_ ICD-10 Code(s): \_\_\_\_\_

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

Is patient going to be using drug in combination with a clinical trial?  Yes  No

Is the requested medication being prescribed by a nephrologist?  Yes  No

Has the patient had a trial and inadequate response or intolerance to calcium acetate tablets or capsules or sevelamer carbonate tablets?  Yes  No

Does the patient have one of the following lab measures?  Yes  No

Calcium and phosphorus product greater than 55mg<sup>2</sup>/dL<sup>2</sup>

Corrected serum calcium level greater than or equal to 9.5 mg/dL (or maximum per lab facility) and the patient is not being treated with vitamin D

Parathyroid hormone (PTH) less than 150 pg/ml (or less than 2 times the upper limit of normal) in a patient with corrected calcium levels of 8.4 mg/dL or greater

Serum phosphorus levels greater than 6.0 mg/dL (or maximum per lab facility)

*Please provide 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?

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**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: \_\_\_\_\_

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**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.

**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