

**Kerendia (finerenone)**  
**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: |
| REQUESTOR (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):        |                                  |                                     |           |

*Continued on next page.*

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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:**

|  |  |
|--|--|
| <input type="checkbox"/> Type II Diabetes(T2D)<br><br><input type="checkbox"/> Other diagnosis: _____ ICD-10 _____ |  |
|--|--|

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

**Clinical Information:**

Is the drug going to be used in conjunction with a clinical trial?  Yes  No

Does patient have a diagnosis of chronic kidney disease(CKD) ?  Yes  No

Does patient have persistent high albuminuria(UACR  $\geq 30$  to  $< 300$ mg/g) ?  Yes  No *Please submit chart documentation.*

Does patient have an estimated glomerular filtration rate(eGFR)  $\geq 25$  but  $< 60$ mL/min/1.73m<sup>2</sup> (CKD EPI)? *Please submit chart documentation.*

Does patient have presence of diabetic retinopathy?  Yes  No *Please submit chart documentation.*

Does patient have persistent high albuminuria(UACR  $\geq 300$ mg/g ?  Yes  No *Please submit chart documentation.*

Does patient have an estimated glomerular filtration rate(eGFR)  $\geq 25$  but  $< 75$ mL/min/1.73m<sup>2</sup> (CKD EPI)?  Yes  No *Please submit chart documentation.*

Has patient been treated with the maximally tolerated dose of an ACE or an ARB at least 4 weeks or more prior to starting Kerendia(finerenone)?  Yes  No *Please submit chart documentation.*

Does patient have known non-diabetic renal disease?  Yes  No

Does patient have renal stenosis?  Yes  No

Does patient have a mean sitting systolic blood pressure(SBP)  $\geq 160$ mmHg?  Yes  No

Does patient have a mean sitting diastolic blood pressure(DBP)  $\geq 100$ mmHg?  Yes  No

Does patient have a mean SBP  $< 90$ mmHg?  Yes  No

Does patient have a HbA1c  $> 12\%$ ?  Yes  No

Does patient have a clinical diagnosis of heart failure with reduced ejection fracture(EF)  $\leq 40\%$  ?  Yes  No *Please submit chart documentation.*

Does patient have NYHA class II-IV?  Yes  No

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Has patient had a stroke, transient ischemic attack, acute coronary syndrome, or hospitalization for worsening heart failure in the last 30 days prior to starting treatment with Kerendia(finerenone)?  Yes  No

Has patient had dialysis for acute renal failure within 12 weeks of starting Kerendia(finerenone)?  Yes  No

Has patient had a renal allograft in place or scheduled withing the next 12 months of starting Kerendia(finerenone)?  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?

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**\*Please note:** Not all drugs/diagnoses 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