



Kalbitor

HMSACOM - Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-237-5512.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-808-254-4414**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Patient's Phone Number: _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Additional Demographic Information:

Patient Weight: _____ *kg*
Patient Height: _____ *ft* _____ *inches*

Indicate where the drug is being dispensed:

- Office Outpatient Hospital Ambulatory Surgical Inpatient Hospital
- Off Campus Outpatient Hospital Urgent Care Emergency Room Birthing Center
- Military Facility Skilled Nursing Facility Nursing Facility Hospice
- Inpatient Psychiatric Psychiatric Residential Treatment End Stage Renal Facility
- Psychiatric Facility Pharmacy Other

Indicate where the drug is being administered:

- Ambulatory surgical Home Inpatient Hospital
- Office Outpatient Hospital Pharmacy

What is the ICD-10 code? _____

Send completed form to: CVS Caremark Specialty Programs. Fax: 1-866-237-5512

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Exception Criteria Questions:

- A. Is the product being requested for the treatment of acute attacks of hereditary angioedema?
 Yes No *If No, skip to Criteria Questions.*
- B. The preferred products for your patient's health plan are icatibant and Ruconest.
Can the patient's treatment be switched to icatibant or Ruconest?
 Yes, *Please obtain Form for preferred product and submit for corresponding PA.*
 No
- C. Is the patient less than 12 years of age? *If Yes, skip to Criteria Questions* Yes No
- D. Is the patient 12 years of age or older but less than 18 years of age? Yes No *If No, skip to G.*
- E. Has the patient tried and experienced a documented inadequate response or intolerable adverse event to Ruconest? ***ACTION REQUIRED: If 'Yes', attach supporting chart note(s).***
If Yes, skip to Criteria Questions Yes No
- F. Does the patient have a documented contraindication to Ruconest (i.e., a known or suspected allergy to rabbits or rabbit-derived products)? ***ACTION REQUIRED: If 'Yes', attach supporting chart note(s).***
If Yes, skip to Criteria Questions Yes No
- G. Has the patient tried and experienced a documented inadequate response, intolerable adverse event or contraindication to both of the preferred products, icatibant AND Ruconest? ***ACTION REQUIRED: If 'Yes', attach supporting chart note(s).*** Yes No

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Criteria Questions:

1. What is the diagnosis?

- Hereditary angioedema (HAE) with C1 inhibitor deficiency or dysfunction confirmed by laboratory testing, *Continue to #2*
- HAE with normal C1 inhibitor confirmed by laboratory testing, *Continue to #3*
- Other, *No Further Questions*

2. Which of the following conditions does the patient have at the time of diagnosis? **ACTION REQUIRED:** *For any answer, attach laboratory test or medical record documentation confirming C1 inhibitor functional and antigenic protein levels*

- A C1 inhibitor (C1-INH) antigenic level below the lower limit of normal as defined by the laboratory performing the test, *Continue to #4*
- A normal C1-INH antigenic level and a low C1-INH functional level (functional C1-INH less than 50% or C1-INH functional level below the lower limit of normal as defined by the laboratory performing the test), *Continue to #4*
- Other, *Continue to #4*

3. Which of the following conditions does the patient have at the time of diagnosis? **ACTION REQUIRED:** *For any answer, attach laboratory test or medical record documentation confirming normal C1 inhibitor. Based on the answer provided, attach genetic test or medical record documentation confirming F12, angiotensin-converting enzyme 1 (ACE-1), plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosaminase 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation testing or chart notes confirming family history of angioedema and the angioedema was refractory to a trial of high-dose antihistamine therapy*

- F12, angiotensin-converting enzyme 1, plasminogen, kininogen-1 (KNG1), heparan sulfate-glucosaminase 3-O-sulfotransferase 6 (HS3ST6), or myoferlin (MYOF) gene mutation as confirmed by genetic testing, *Continue to #4*
- BOTH of the following: 1) Angioedema refractory to a trial of high-dose antihistamine therapy (i.e., cetirizine at 40 mg per day or the equivalent) for at least one month AND 2) Family history of angioedema, *Continue to #4*
- Other, *Continue to #4*

4. Is the requested medication being used for the treatment of acute HAE attacks?

- Yes, *Continue to #5*
- No, *Continue to #5*

5. Will the requested medication be used in combination with any other medication used for the treatment of acute HAE attacks (e.g., Berinert, Firazyr, or Ruconest)?

- Yes, *Continue to #6*
- No, *Continue to #6*

6. Will the requested medication be prescribed by or in consultation with a prescriber who specializes in the management of hereditary angioedema (HAE)?

- Yes, *Continue to #7*
- No, *Continue to #7*

7. Have other causes of angioedema been ruled out (e.g., angiotensin-converting enzyme inhibitor [ACE-I] induced angioedema, angioedema related to an estrogen-containing drug, allergic angioedema)?

- Yes, *Continue to #8*
- No, *Continue to #8*

8. Is this request for continuation of therapy with the requested drug?

- Yes, *Continue to #9*
- No, *No Further Questions*

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9. Has the patient experienced a reduction in severity and/or duration of attacks? **Action Required:** If 'Yes', attach supporting chart note(s) demonstrating a reduction in severity and/or duration of attacks

- Yes, *Continue to #10*
- No, *Continue to #10*

10. Does the patient's attack frequency, attack severity, comorbid conditions and patient's quality of life warrant prophylactic therapy?

- Yes, *Continue to #11*
- No, *No Further Questions*

11. Has prophylactic treatment been considered?

- Yes, *No Further Questions*
- No, *Continue to #12*

12. Please provide a brief rationale as to why prophylactic treatment has not been considered.

_____, *No Further Questions*

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____
Prescriber or Authorized Signature **Date (mm/dd/yy)**

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