

Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient? Yes No

Diagnosis related to use:

hereditary angioedema (HAE) other (please specify):

Clinical Information:

****This drug requires supportive documentation (chart notes, lab and test results, etc). Supportive documentation for all answers must be attached with this request****

For all products:

****Supportive documentation for all answers must be attached with this request.****

Does your patient have a confirmed monoallelic mutation in either the SERPING1 or F12 gene which is known to cause HAE?

Yes No ****documentation required**

(if no) Did your patient have a C4 level below the lower limit of normal as defined by the laboratory performing the test?

Yes No ****documentation of labs required**

Did/does your patient have a C1 inhibitor (C1INH) antigenic level below the lower limit of normal as defined by the laboratory performing the test?

Yes No ****documentation of labs required**

(if no) Did your patient have a C1INH functional level below the lower limit of normal as defined by the laboratory performing the test?

Yes No ****documentation of labs required**

If your patient was/is taking any medications known to cause these attacks (for example: ACE-I, ARB, or estrogens), have these been evaluated and discontinued when appropriate? Yes or patient was not taking any known medications No

Is this drug being prescribed by, or in consultation with, an allergist/immunologist or a physician who specializes in the treatment of HAE or related disorders? Yes No

For Cinryze, Haegarda, Takhzyro:

****Supportive documentation for all answers must be attached with this request.****

Is this drug being used to prevent angioedema attacks? Yes No

Does your patient have a history of 2 or more moderate or severe attacks per month (for example airway swelling, severe abdominal pain, facial swelling, nausea and vomiting, painful facial distortion)? Yes No

While receiving the requested drug, will your patient also be treated with any of the following? (check all that apply)

Cinryze Haegarda Takhzyro other (please specify):

For Berinert, Firazyr, Icatibant and Kalbitor:

****Supportive documentation for all answers must be attached with this request.****

Is this drug being used for the treatment of acute angioedema attacks? Yes No

Does your patient have a history of a moderate or severe attacks (for example: airway swelling, severe abdominal pain, facial swelling, nausea and vomiting, painful facial distortion)? Yes No

Is this request for a quantity to treat more than 2 attacks per month? Yes No

If yes- please provide clinical information to support:

While receiving the requested drug, will your patient also be treated with any of the following? (check all that apply)

Berinert Firazyr Icatibant Kalbitor Ruconest other (please specify):

For Firazyr only- Has your patient tried a generic formulation of icatibant? Yes No

(if yes) Did your patient have a documented intolerance to icatibant? Yes No

For Ruconest:

****Supportive documentation for all answers must be attached with this request.****

Is this drug being used for treatment of acute angioedema attacks? Yes No

Does your patient have a history of moderate or severe attacks involving the abdomen, face, or extremities? Yes No

Is this request for a quantity to treat more than 2 attacks per month? Yes No

If yes- please provide clinical information to support:

While receiving the requested drug, will your patient also be treated with any of the following? (check all that apply)

Berinert Firazyr Icatibant Kalbitor Ruconest other (please specify):

For Berinert, Kalbitor, Ruconest:

Does your patient have documented failure or inadequate response, contraindication per FDA label, intolerance or is not a candidate for one formulation of icatibant (generic Firazyr)? Yes No

Additional pertinent information: (include alternatives tried, date(s) taken and for how long, and what the documented results were of taking this drug, including any intolerances or adverse reactions your patient experienced):

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer 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: _____ **Date:** _____

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