Medimpact

#### STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

#### **C1 ESTERASE INHIBITOR - CINRYZE**

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
C1 ESTERASE	CINRYZE	18568		GPI-10	FDB & MEDI-SPAN:
INHIBITOR				(8580202200)	BRAND = CINRYZE

#### **GUIDELINES FOR USE**

### INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

- 1. Does the patient have a diagnosis of hereditary angioedema (HAE) and meet **ALL** of the following criteria?
  - The patient is 6 years of age or older
  - Therapy is prescribed by or in consultation with an allergist, immunologist or hematologist
  - The patient's diagnosis of HAE is confirmed via documentation of complement testing
  - Cinryze is being used for prophylaxis against HAE attacks
  - Cinryze is NOT being used concurrently with alternative prophylactic agent for HAE (e.g., Takhzyro, Haegarda, danazol)

If yes, **approve for 12 months for all NDCs with a quantity limit of #40 vials per 28 days.** If no, do not approve.

# **INITIAL DENIAL TEXT:** \*Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **C1 ESTERASE INHIBITOR - CINRYZE** requires the following rule(s) be met for approval:

- A. You have hereditary angioedema (HAE: a type of gene condition with severe body swelling)
- B. You are 6 years of age or older
- C. Therapy is prescribed by or in consultation with an allergist (a type of allergy doctor), immunologist (a type of immune system doctor) or hematologist (a type of blood doctor)
- D. Your diagnosis is confirmed by documented complement testing (a type of lab test)
- E. Cinryze is being used for prevention of hereditary angioedema attacks
- F. You will not be using Cinryze concurrently (at the same time) with an alternative preventive agent for HAE (such as Takhzyro, Haegarda, danazol)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

## CONTINUED ON NEXT PAGE

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STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

## **C1 ESTERASE INHIBITOR - CINRYZE**

## **GUIDELINES FOR USE (CONTINUED)**

### **RENEWAL CRITERIA**

- 1. Does the patient have a diagnosis of hereditary angioedema (HAE) and meet **ALL** of the following criteria?
  - The patient has experienced improvement (i.e., reductions in attack frequency or attack severity) compared to baseline in HAE attacks
  - Cinryze will NOT be used concurrently with alternative prophylactic agent for HAE (e.g., Takhzyro, Haegarda, danazol)

If yes, **approve for 12 months by NDC with a quantity limit of #40 vials per 28 days.** If no, do not approve.

**RENEWAL DENIAL TEXT:** \*Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **C1 ESTERASE INHIBITOR - CINRYZE** requires the following rule(s) be met for renewal:

- A. You have hereditary angioedema (HAE: a type of gene condition with severe body swelling)
- B. You have experienced improvement (reductions in attack frequency or attack severity) compared to baseline in HAE attacks
- C. You will NOT be using Cinryze concurrently (at the same time) with alternative prophylactic (preventive) agent for HAE (such as Takhzyro, Haegarda, danazol)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

## RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Cinryze.

#### REFERENCES

• Cinryze [Prescribing Information]. Lexington, MA: Shire Viropharma Inc. December 2019.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Commercial Effective: 11/01/22 Created: 10/22 Client Approval: 10/22

P&T Approval: 04/22

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