



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

MAVACAMTEN

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
MAVACAMTEN	CAMZYOS	47972		GPI-10 (4019005000)	

GUIDELINES FOR USE

NITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

1. Does the patient have a diagnosis of symptomatic obstructive hypertrophic cardiomyopathy (HCM) and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has New York Heart Association (NYHA) class II-III symptoms
 - The patient has a left ventricular outflow track (LVOT) gradient of 50 mmHg or higher
 - Therapy is prescribed by or in consultation with a cardiologist
 - The patient had a trial of or contraindication to beta-blockers (e.g., metoprolol, carvedilol) AND non-dihydropyridine calcium channel blockers (e.g., verapamil, diltiazem)

If yes, **approve for 4 months by HICL or GPI-10 with a quantity limit of #1 per day.**

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 **MAVACAMTEN (Camzyos)** requires the following rule(s) be met for approval:

- A. You have symptomatic obstructive hypertrophic cardiomyopathy (HCM: a type of heart condition)
- B. You are 18 years of age or older
- C. You have New York Heart Association (NYHA) class II-III (classification system for heart failure) symptoms
- D. You have a left ventricular outflow track gradient (a predictor of heart failure and cardiovascular death) of 50 mmHg or higher
- E. Therapy is prescribed by or in consultation with a cardiologist (a type of heart doctor)
- F. You had a trial of or contraindication (harmful for) to beta-blockers (such as metoprolol, carvedilol) AND non-dihydropyridine calcium channel blockers (such as verapamil, diltiazem)

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



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

MAVACAMTEN

GUIDELINES FOR USE (CONTINUED)

RENEWAL CRITERIA

1. Does the patient have a diagnosis of symptomatic obstructive hypertrophic cardiomyopathy (HCM) **AND** meet the following criterion?

- The patient has experienced continued clinical benefit (e.g., reduction of symptoms, NYHA classification improvement)

If yes, **approve for 12 months by HICL or GPI-10 with quantity limit of #1 per day.**

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 **MAVACAMTEN (Camzyos)** requires the following rule(s) be met for renewal:

- A. You have symptomatic obstructive hypertrophic cardiomyopathy (HCM: a type of heart condition)
- B. You have experienced continued clinical benefit (such as reduction of symptoms, NYHA classification improvement)

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 Camzyos.

REFERENCES

- Camzyos [Prescribing Information]. Brisbane, CA: MyoKardia, Inc.; April 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

Commercial Effective:06/01/22

Created: 05/22

Client Approval: 05/22

P&T Approval: 04/22