

## **Request for Prior Authorization**

### FAX Completed Form To I (800) 574-2515

**Provider Help Desk** 1 (877) 776-1567

# **MAVACAMTEN (CAMZYOS)**

(PLEASE PRINT – ACCURACY IS IMPORTANT)

	(			
IA Medicaid Member ID #	Patient name		DOB	
Patient address				
Provider NPI	Prescriber name		Phone	
Prescriber address			Fax	
Pharmacy name	Address		Phone	
Prescriber must complete all informa	tion above. It must be legible, con	rect, and complete or fo	rm will be returned.	
Pharmacy NPI	Pharmacy fax	NDC		
considered when documented evid contraindicated. Payment will be contrained the following when the following I) Request adheres to all FDA appropriate contraindications, warnings and	onsidered for an FDA approved conditions are met: proved labeling for requested d	d or compendia indica	ted diagnosis for the cluding age, dosing,	
2) Patient has a diagnosis of obstructive hypertrophic cardiomyopathy (HCM); and				
3) Patient exhibits symptoms of New York Heart Association (NYHA) class II or III symptoms; and				
4) Is prescribed by or consultation with a cardiologist; and				
5) Patient has a left ventricular ejo	ection fraction (LVEF) $\geq$ 55%; a	nd		
6) Patient has a peak left ventricular outflow tract (LVOT) gradient ≥ 50 mmHg at rest or with provocation; and				
7) Documentation of a previous to	rial and therapy failure, at a ma	ximally tolerated dos	e, with all of the following:	
b. Non-dihydropyridine calciu	ker (atenolol, metoprolol, biso m channel blocker (verapamil, disopyramide plus beta-blocke	diltiazem); and		
The required trials may be override be medically contraindicated.	den when documented evidenc	e is provided that the	use of these agents would	
Request for continuation of therapy evidenced by improvement in obst		nentation of a positive	e response to therapy as	
Non-Preferred				
☐ Camzyos				
Strength	Dosage Instructions	Quantity D	ays Supply	
Diagnosis:				
Prescriber Specialty:   Cardi	ologist Other (specify):			

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If other, note consultation with cardiologist: Consultation date:

Physician name, specialty & phone:				
Does patient exhibit symptoms of NYHA class II or	III symptoms?			
<b>Does patient have LVEF</b> ≥ <b>55%?</b>				
Does patient have LVOT gradient ≥ 50 mmHg at re	est or with provocation? No Yes			
Document trials, at a maximally tolerated dose, with	th all of the following:			
Non-vasodilating beta-blocker trial (atenolol, metolorug Name & Dose:  Failure reason:	Trial dates:			
Non-dihydropyridine calcium channel blocker trial	(verapamil, diltiazem): Trial dates:			
Combination therapy with disopyramide plus beta-blocker:				
Failure reason:	Trial dates:			
Non-vasodilating beta-blocker trial (atenolol, metoprolol, b Drug Name & Dose:	Trial dates:			
OR				
Non-dihydropyridine calcium channel blocker trial (verapar Drug Name & Dose:  Failure reason:	Trial dates:			
Renewal Requests:  Document positive response to therapy as evidence	ed by improvement in HCM symptoms:			
Attach lab results and other documentation as necessary.				
Prescriber signature (Must match prescriber listed above.)	Date of submission			
<b>IMPORTANT NOTE:</b> In evaluating requests for prior authorization the co	nsultant will consider the treatment from the standpoint of medical necessity only.			

IMPORTANT NOTE: In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only if approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member continues to be eligible for Medicaid.

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