



**HEALTH PARTNERS PLANS
PRIOR AUTHORIZATION REQUEST FORM**

Camzyos - Non-PDL

Phone: 215-991-4300

Fax back to: 866-240-3712

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL DELAY the review process.

Member Name:	Prescriber Name:	
HPP Member Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Member Primary Phone:	NPI:	PA PROMISE ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Line of Business: <input type="checkbox"/> Medicaid <input type="checkbox"/> CHIP	Specialty Pharmacy (if applicable):	
Drug Name:	Strength:	
Quantity:	Refills:	
Directions:		
Diagnosis Code:	Diagnosis:	
<i>HPP's maximum approval time is 12 months but may be less depending on the drug.</i>		

Please attach any pertinent medical history including labs and information for this member that may support approval.

Please answer the following questions and sign.

Q1. Is this a renewal request? If YES, go to 12. If NO, go to 2.

Yes No

Q2. Is Camzyos (mavacamten) being prescribed by or in consultation with a cardiologist?

Yes No

Q3. Is the patient greater than or equal to 18 years of age?

Yes No

Q4. Are clinical notes attached that document a confirmed diagnosis of symptomatic New York Heart Association (NYHA) class II or class III obstructive hypertrophic cardiomyopathy (oHCM), noting associated symptoms? [for example: NYHA Functional Classification - class / symptoms: Class II - Slight limitation of physical activity. Comfortable at rest. Ordinary physical activity results in fatigue, palpitation, shortness of breath or chest pain AND Class III - Marked limitation of physical activity. Comfortable at rest. Less than ordinary activity causes fatigue, palpitation, shortness of breath or chest pain.]

Yes No

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Member Name:	Prescriber Name:
<p>Q5. Does a recent echocardiogram assessment show left ventricular ejection fraction (LVEF) greater than or equal (>/=) to 55%?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q6. Does the patient have a peak left ventricular outflow tract (LVOT) gradient greater than or equal (>/=) to 50 mmHg at rest or with provocation.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q7. Is there a documented history attached supporting trial and failure, contraindication or intolerance to a non-vasodilating beta blocker (such as metoprolol succinate, nadolol) titrated to effectiveness or maximally tolerated doses.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q8. Is there a documented history attached supporting trial and failure to at least one of the agent types below OR an intolerance to both agents below OR contraindication to both agents below:</p> <p><input type="checkbox"/> a) Non-dihydropyridine calcium channel blockers (such as verapamil er, diltiazem er) <input type="checkbox"/> Disopyramide OR,</p>	
<p>Q9. Have other disorders that cause cardiac hypertrophy been ruled out (such as, cardiac amyloidosis, Fabry disease, Noonan syndrome)?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q10. Is the drug being prescribed at an FDA-approved dose?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q11. Will the patient be taking Camzyos (mavacamten) with strong CYP2C19 inhibitors or moderate to strong CYP2C19 inducers or moderate to strong CYP3A4 inducers?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q12. Is there attached documentation showing clinical benefit from baseline as evidenced by improvement in symptoms associated with New York Heart Association (NYHA) class II or class</p>	



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<p>III obstructive hypertrophic cardiomyopathy (oHCM) OR that the NYHA is not worsening from baseline?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q13. Does the patient have a left ventricular ejection fraction (LVEF) greater than or equal to (>/=) to 50%?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q14. Is the Camzyos (mavacamten) being prescribed at an FDA-approved dose?</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q15. Additional Information:</p>	

Prescriber Signature

Date

v2026