

US Family Health Plan
Prior Authorization Request Form for
mavacamten (Camzyos)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

Prior authorization approval expires after 1 year. For renewal of therapy an initial Tricare prior authorization approval is required.

Step 1 Please complete patient and physician information (please print):

Patient Name: _____	Physician Name: _____
Address: _____	Address: _____
Sponsor ID # _____	Phone #: _____
Date of Birth: _____	Secure Fax #: _____

Step 2 Please complete the clinical assessment:

1. Has the patient received this medication under the TRICARE benefit in the last 6 months? Please choose "No" if the patient did not previously have a TRICARE approved PA for Camzyos	<input type="checkbox"/> Yes (subject to verification) Proceed to question 2	<input type="checkbox"/> No Proceed to question 3
2. Has the patient responded to therapy, as evidenced by improvement in obstructive hypertrophic cardiomyopathy symptoms?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved
3. Is the patient greater than or equal to 18 years of age?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No STOP Coverage not approved
4. Is the requested medication prescribed by a cardiologist?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No STOP Coverage not approved
5. Does the patient have documented evidence of obstructive hypertrophic cardiomyopathy (HCM)?	<input type="checkbox"/> Yes Proceed to question 6	<input type="checkbox"/> No STOP Coverage not approved
6. Is the left ventricular outflow tract (LVOT) pressure greater than or equal to 50 mmHg?	<input type="checkbox"/> Yes Proceed to question 7	<input type="checkbox"/> No STOP Coverage not approved
7. Does the patient have New York Heart Association (NYHA) Class II to III obstructive hypertrophic cardiomyopathy that is symptomatic (for example; dyspnea, chest pain, light headedness, syncope, fatigue, reduced exercise capacity)?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No STOP Coverage not approved

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8. Is the left ventricular ejection fraction (LVEF) greater than or equal to 55%?	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No STOP Coverage not approved
9. Has the patient failed therapy with at least one agent from both classes: <ul style="list-style-type: none"> • beta blocker (non-vasodilating) propranolol, metoprolol AND • calcium channel blockers (non-dihydropyridine) verapamil or diltiazem? 	<input type="checkbox"/> Yes Proceed to question 10	<input type="checkbox"/> No STOP Coverage not approved
10. Is the patient on dual calcium channel blocker and beta blocker therapy concurrently?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 11
11. Is the patient receiving ranolazine (Ranexa) or disopyramide (Norpace, Rythmodan) concurrently?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 12
12. What is the patient's sex?	<input type="checkbox"/> Female Proceed to question 13	<input type="checkbox"/> Male Proceed to question 16
13. Is the patient of childbearing potential?	<input type="checkbox"/> Yes Proceed to question 14	<input type="checkbox"/> No Proceed to question 16
14. Has the patient received counseling for using effective contraception during therapy with Camzyos and for 4 months after the last dose?	<input type="checkbox"/> Yes Proceed to question 15	<input type="checkbox"/> No STOP Coverage not approved
15. Is the patient pregnant?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 16
16. Are the patient and provider aware of the risks of systolic dysfunction, as outlined in the REMS program?	<input type="checkbox"/> Yes Proceed to question 17	<input type="checkbox"/> No STOP Coverage not approved
17. Will the patient and the provider agree to comply to all requirements of the REMS program, including echocardiogram at 0, 4, 8, 12 weeks follow by every 12 weeks and drug interaction monitoring requirements?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

Step 3 I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date