

Pharmacy Request for Prior Approval – Camzyos

Beneficiary Information		
1. Beneficiary Last Name:		
3. Beneficiary ID #:	4. Beneficiary Date of Birth:	5. Beneficiary Gender:
Prescriber Information		
7. Prescriber Name:		
Mailing address:	City:	State: ZIP:
8. Requester Contact Information:		
Name:	Phone #:	Fax #:
Drug Information		
8. Drug Name:	9. Strength:	10. Quantity Per 30 Days:
	60 days90 days120 day	s180 days365 daysOther:
Clinical Information		
Initial Requests for Camzyos: (questions 1-	-10)	
1. Is the beneficiary 18 years of age or olde		
2. Does the beneficiary have a diagnosis of obstructive hypertrophic cardiomyopathy (oHCM) consistent with current guidelines (e.g., American College of Cardiology Foundation/American Heart Association, European Society of Cardiology guidelines)? YesNo		
3. Does the beneficiary have New York Hea	rt Association (NYHA) Class 2 or Class	3? YesNo
4. Will the beneficiary be monitored for LVI	EF, Valsalva left ventricular outflow tr	act (LVOT) gradient assessment, and heart failure
		atigue, swelling in the legs)? Yes No
5. Does the beneficiary have adequate echo	ocardiogram or cardiovascular magne	tic resonance imaging (CMR)? Yes No
6. Will the beneficiary avoid concomitant use with moderate to strong CYP2C19 inhibitors, strong CYP3A4 inhibitors, and moderate to strong CYP2C19 and CYP3A4 inducers (e.g., carbamazepine, cimetidine, esomeprazole, omeprazole, phenobarbital, phenytoin, rifampin, St. John's wort)? Yes No		
7. For females of childbearing potential, has a pregnancy test been performed ensuring beneficiary is not pregnant? YesNo		
8. Will Mavacamten be prescribed by or in	consultation with a cardiologist? Yes	No
9. Has the beneficiary had an adequate tria	I and failure of ≥ 1 beta-blocker? Yes	s No List:
10. Does the beneficiary have documented left ventricular ejection fraction (LVEF) ≥ 55%? (for initiation of treatment only) YesNo		
Continuation Requests for Camzyos: (ques	stions 1-9 above and 11-13)	
11. Has the beneficiary had disease improvement and/or stabilization of disease from baseline (e.g., NYHA class improvement [class 3 to class 2], ≥ 1.5 mL/kg/min in pVO2 increase or ≥ 3 mL/kg/min in pVO2 without NYHA class worsening)? Yes No		
12. Does the beneficiary have left ventricular ejection fraction (LVEF) ≥ 50%? Yes No		
13. Has the beneficiary experienced any treatment-restricting adverse effects (e.g., heart failure)? Yes No		
13. Has the beneficially experienced any treatment-restricting adverse effects (e.g., flear trailule): TesNO		
Signature of Prescriber:	Date:	

*Prescriber signature mandatory

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.