

Mavacamten (Camzyos)

Coverage Criteria:

Initial Authorization Criteria:

Camzyos is reserved for:

- 1. Prescribed by or in consultation with a cardiologist; and,
- 2. Patient has a diagnosis of symptomatic obstructive hypertrophic cardiomyopathy (oHCM); and,
- 3. Patient has New York Heart Association (NYHA) class II-III symptoms (e.g., effort-related shortness of breath or chest pain); and,
- 4. Patient has a left ventricular ejection fraction (LVEF) ≥ 55%; and,
- 5. Patient has a left ventricular outflow track (LVOT) gradient of ≥ 50 mmHg or higher; and,
- 6. Patient has not undergone a septal reduction procedure within the last 6 months; and,
- 7. Patient has had a trial and failure of, intolerance to, or medical contraindication to both of the following:
 - a. Beta-blockers (e.g., metoprolol, carvedilol); and,
 - b. Non-dihydropyridine calcium channel blockers (e.g., verapamil, diltiazem); and,
- 8. Prescribed per FDA dosing regimen.

Quantity Limit: 1 capsule per day

Renewal Criteria:

- 1. Patient continues to meet the criteria above; and,
- 2. Patient has been seen and evaluated by prescriber in the past 12 months; and,
- 3. Patient has not undergone a septal reduction procedure within the last 6 months; and,
- 4. Patient has had a clinically meaningful response to therapy per medical chart notes (e.g., reduction of symptoms, NYHA classification improvement).

Coverage Duration:

Initial authorization will be provided for 6 months. Re-authorizations will be provided for 12 months.

P&T Date: 8/8/2022

Effective Date: 10/1/2022