

# CardioMEMS™ HF System Clinical Checklist

This form is intended to be used by the treating physician as a guide to help gather information in connection to a heart failure diagnosis. The documentation referenced below may be submitted to the insurance company prior to procedure being performed.

## HEALTHCARE PROVIDER INSTRUCTIONS

**DO NOT INCLUDE THIS INSTRUCTION PAGE IN YOUR SUBMISSION**

1. The Physician completes the CardioMEMS HF System Clinical Checklist, sign and date.
2. The documents below may be submitted to the insurance company for prior authorization.
  - CardioMEMS HF System Clinical Checklist
  - 3-6 months of treatment notes
  - Echocardiogram report
  - Hospitalizations for heart failure including discharge summaries
  - Elevated BNP/NT-proBNP lab values and reports

For questions on the CardioMEMS HF System Clinical Checklist, please contact the Patient Therapy Access Team at [PTA\\_cardiac@abbott.com](mailto:PTA_cardiac@abbott.com) or the Health Economics Team at [heartfailureeconomics@abbott.com](mailto:heartfailureeconomics@abbott.com).

### Rx Only

**Brief Summary:** Prior to using these devices, please review the Instructions for Use for a complete listing of indications, contraindications, warnings, precautions, potential adverse events and directions for use.

**CardioMEMS™ HF System Indications and Usage:** The CardioMEMS HF System is indicated for wirelessly measuring and monitoring pulmonary artery pressure and heart rate in NYHA Class II or III heart failure patients who either have been hospitalized for heart failure in the previous year and/or have elevated natriuretic peptides. The hemodynamic data are used by physicians for heart failure management with the goal of controlling pulmonary artery pressures and reducing heart failure hospitalizations.

**CardioMEMS™ HF System Contraindications:** The CardioMEMS™ HF System is contraindicated for patients with an inability to take dual antiplatelet or anticoagulants for one month post implant.

**CardioMEMS™ HF System Potential Adverse Events:** Potential adverse events associated with the implantation procedure include, but are not limited to, the following: air embolism, allergic reaction, infection, delayed wound healing, arrhythmias, bleeding, hemoptysis, hematoma, nausea, cerebrovascular accident, thrombus, cardiovascular injury, myocardial infarction, death, embolization, thermal burn, cardiac perforation, pneumothorax, thoracic duct injury and hemothorax.

### Reference

2022 AHA/ACC/HFSA Guideline for the Management of Heart Failure: A Report of the American College of Cardiology/American Heart Association Joint Committee on Clinical Practice Guidelines. <https://www.ahajournals.org/doi/full/10.1161/CIR.0000000000001063>

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Patient Name: \_\_\_\_\_

Date of Birth: \_\_\_\_\_

## Guideline Directed Medical Therapy

On GDMT with a goal of being on maximally tolerated doses (with optimal device therapy if applicable) per the 2022 ACC/AHA/HFSA/CHF guidelines based on the patient's ejection fraction.

### CURRENT MEDICATION(S):

MEDICATION	CONTRAINDICATION
<input type="checkbox"/> ACE/ARB/ARNI	
<input type="checkbox"/> SGLT2i	
<input type="checkbox"/> MRA	
<input type="checkbox"/> Beta Blocker	
<input type="checkbox"/> Diuretic	
<input type="checkbox"/> Other	

### Guideline Recommendations per the 2022 ACC/AHA/HFSA CHF Guidelines based on patient's ejection fraction

For EF  $\leq$  40%: On beta blockers for 3 months, on diuretics, SGLT2i, MRA, ACEi/ARB/ARNi

For EF 41-49%: On diuretic, SGLT2i, MRA, ACEi/ARB/ARNi, beta blockers

For EF greater than or equal to 50% on diuretic, SGLT2i, MRA, ACEi/ARG/ARNi

### PATIENT HISTORY

**NYHA CLASS**     II     III

Any major cardiovascular event within the last 3 months     YES     NO

BNP/NT-proBNP lab value (result within the last 30 days): \_\_\_\_\_

Ejection Fraction: \_\_\_\_\_

BMI  $\leq$  35     YES     NO

If no, Chest circumference measured at axillary level: \_\_\_\_\_

History of HF hospitalization within the past 12 months:     YES     NO

Dates of admission(s) \_\_\_\_\_

### DEVICE THERAPY

Evaluated for, and received if appropriate, an ICD/CRT-P/CRT-D     YES     NO

If yes, was the device implanted  $\geq$ 3 months ago?     YES     NO

### CONNECTIVITY

Has access to reliable connectivity at home     YES     NO

**DIAGNOSIS**

- |  |   |  |
|--|---|--|
| <input type="checkbox"/> I50.1 Left ventricular failure                              | <input type="checkbox"/> I50.30 Unspecified diastolic (congestive) heart failure                                    | <input type="checkbox"/> I50.42 Chronic combined systolic (congestive) and diastolic (congestive) heart failure          |
| <input type="checkbox"/> I50.20 Unspecified systolic (congestive) heart failure      | <input type="checkbox"/> I50.32 Chronic diastolic (congestive) heart failure  | <input type="checkbox"/> I50.43 Acute on chronic combined systolic (congestive) and diastolic (congestive) heart failure |
| <input type="checkbox"/> I50.22 Chronic systolic (congestive) heart failure          | <input type="checkbox"/> I50.40 Unspecified combined systolic (congestive) and diastolic (congestive) heart failure | <input type="checkbox"/> I50.9 Heart failure, unspecified  |
| <input type="checkbox"/> I50.23 Acute on chronic systolic (congestive) heart failure |   |  |
| <input type="checkbox"/> Other: _____  |   |  |

Healthcare Provider Signature: \_\_\_\_\_

Healthcare Provider Name (Printed): \_\_\_\_\_

Date: \_\_\_\_\_