Hydralazine:* □ YES □ NO □ UNK
Calcium channel blockers:* □ YES □ NO □ UNK
Nesiritide:* □ YES □ NO □ UNK
Angiotensin receptor blocker drug:* □ YES □ NO □ UNK
Amiodarone:* □ YES □ NO □ UNK
ACE inhibitors:* □ YES □ NO □ UNK
Beta-blockers:* □ YES □ NO □ UNK
Aldosterone antagonist:* □ YES □ NO □ UNK
Lovenox:* □ YES □ NO □ UNK
Warfarin (coumadin):* □ YES □ NO □ UNK
Antiplatelet therapy drug:* □ YES □ NO □ UNK
  Select drug(s)*

□ Aspirin
□ DEXTran
□ Dipyridamole
□ Clopidogrel
□ Ticlopidine
□ Unknown
□ Other, specify

Nitric oxide:* □ YES □ NO □ UNK
Phosphodiesterase inhibitor:* □ YES □ NO □ UNK
Digoxin:* □ YES □ NO □ UNK
Loop Diuretics:* □ YES □ NO □ UNK
Antibiotics/Anitfungals:* □ YES □ NO □ UNK
  List:

□ Amikacin
□ Amphotericin
□ Anidulafungin
□ Aztreonam
□ Caspofungin
□ Cefazolin
□ Cefuroxime
□ Ceftazidime
□ Ceftriaxone
□ Cefepime
□ Ceftaroline
□ Ciprofloxacin
## Medical Condition

**NYHA class:**
- ○ Class I: No limitation of physical activity; physical activity does not cause fatigue, palpitation or shortness of breath.
- ○ Class II: Slight limitation of physical activity; comfortable at rest, but ordinary physical activity results in fatigue, palpitations or shortness of breath.
- ○ Class III: Marked limitation of physical activity; comfortable at rest, but less than ordinary activity causes fatigue, palpitation or shortness of breath.
- ○ Class IV: Unable to carry on minimal physical activity without discomfort; symptoms may be present at rest.
- ○ Unknown

**Has patient been rehospitalized since implant hospitalization?**
- ○ YES
- ○ NO
- ○ UNK

**If yes:**

## Intervention

**Intervention since implant:**

## Registry Status

**Transferred care to another hospital (patient followed exclusively at another hospital):**
- ○ YES
- ○ NO

**Date transferred care:**

**Patient withdraws consent and therefore no more clinical data is to be collected:**
- ○ YES
- ○ NO

**Date withdrawn:**
Adverse Events

Did the patient have one or more of the following adverse events occur during this follow-up time period? Please make sure you have entered all events that occurred during this follow-up time period.

- Major Bleeding
- Major Infection
- Neurological Dysfunction
- Device Malfunction (if suspected device thrombosis, then enter as Device Malfunction)
- Death
- Transplant
- Explant due to Transplant
- Explant due to Recovery
- Explant due to Exchange
- Respiratory Failure
- Arterial Non-CNS Thromboembolism

Note: Go to section 2.10 for the definition of each Adverse Event.