



A. PARTICIPANT ADMINISTRATION:

Participant ID<sup>1000</sup>/Name<sup>1010</sup>: Medicare Provider #<sup>1015</sup>: Participant NPI<sup>1016</sup>:

B. DEMOGRAPHICS:

Last Name<sup>2000</sup>: First Name<sup>2010</sup>: Middle Name<sup>2020</sup>:
SSN<sup>2030</sup>: Unique Patient Id<sup>2040</sup>: (automatic) Other ID<sup>2045</sup>:
Date of Birth<sup>2050</sup>: Gender<sup>2060</sup>: Male; Female
Race<sup>2070</sup>: White; Black/African American; Asian; American Indian/Alaska Native; Native Hawaiian; Other
Hispanic Ethnicity<sup>2075</sup>: No; Yes
Auxiliary 1<sup>2080</sup>: Auxiliary 2<sup>2090</sup>:

C. ADMISSION:

Admission Date<sup>3000</sup>: Date of Implant<sup>3010</sup>:
Insurance Payor-Primary<sup>3020</sup>: Government; Commercial; HMO; Non-U.S. Insurance; None/Self Pay
Insurance Payor-Secondary<sup>3027</sup>: Government; Commercial; HMO; Non-U.S. Insurance; None/Self Pay
Reason for Admission<sup>3030</sup>: Admitted for this Procedure; Cardiac-CHF; Cardiac-Other; Non-Cardiac
Auxiliary 3<sup>3040</sup>: Auxiliary 4<sup>3050</sup>:

D. HISTORY AND RISK FACTORS:

Syncope<sup>3060</sup>: No; Yes Family Hx Sudden Death<sup>3070</sup>: No; Yes
CHF<sup>3080</sup>: No; Yes
CHF Duration<sup>3090</sup>: Within the past 3 months; 3 to 9 months; Greater than 9 months
Prior CHF Hospitalization<sup>3095</sup>: Not Hospitalized; Yes-Within 6 months; Yes-Greater than 6 months
NYHA Functional Class (Current Status)<sup>3100</sup>: Class I; Class II; Class III; Class IV
Cardiac Arrest<sup>3110</sup>: No Arrest; Brady Arrest; Tachy Arrest
Brady Arrest Reason<sup>3111</sup>: Acute MI; Severe Electrolyte Disturbance; Drug Induced Arrhythmia; Sinus Node Dysfunction/AV Block; Unknown Etiology
Tachy Arrest Reason<sup>3112</sup>: Acute MI; Severe Electrolyte Disturbance; Drug Induced Arrhythmia; Primary VT/VF; Unknown Etiology
Atrial Fibrillation or Flutter<sup>3120</sup>: No; Yes
Ventricular Tachycardia<sup>3130</sup>: No; Yes-VT, Non-Sustained; Yes-Monomorphic Sustained VT; Yes-Polymorphic Sustained VT
Sinus Node Function<sup>3140</sup>: Normal; Abnormal
Cardiac Transplant<sup>3150</sup>: No; Yes
Non-Ischemic Dilated Cardiomyopathy<sup>3160</sup>: No; Yes-Within the past 3 months; Yes-3 to 9 months; Yes-Greater than 9 months
Ischemic Heart Disease<sup>3180</sup>: No; Yes-At Least One Epicardial Artery > 70%; Yes-Other Diagnostic Tests
Previous MI<sup>3190</sup>: No; Yes-Within 40 days; Yes-Greater than 40 days; Yes-Both Within 40 days/Greater than 40 days
Previous CABG<sup>3200</sup>: No; Yes -> if Yes, Date<sup>3210</sup>:
Previous PCI<sup>3220</sup>: No; Yes-Within the past 3 months; Yes-Greater than 3 months
Previous Valvular Surgery<sup>3230</sup>: No; Yes
Permanent Pacemaker<sup>3240</sup>: No; Yes-Atrial Chamber; Yes-Ventricular Chamber; Yes-Dual Chamber; Yes-Biventricular
Previous ICD<sup>3250</sup>: No; Yes-Single Chamber; Yes-Dual Chamber; Yes-Biventricular
Previous ICD Reason<sup>3280</sup>: Spontaneous Monomorphic Sustained VT; Spontaneous Polymorphic Sustained VT; Ventricular Fibrillation; Cardiac Arrest/Arrhythmia-Etiology Unknown; Syncope and High Risk Characteristics; AFib; Primary Prevention; Syncope with Inducible VT
Previous ICD Implant Site<sup>3290</sup>: Pectoral; Abdominal
Cerebrovascular Disease<sup>3310</sup>: No; Yes Chronic Lung Disease<sup>3320</sup>: No; Yes
Diabetes<sup>3330</sup>: No; Yes Hypertension<sup>3340</sup>: No; Yes
Renal Failure Dialysis<sup>3350</sup>: No; Yes

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number. The valid OMB control number for this information collection is 0938-0967. The time required to complete this information collection is estimated to average fifteen (15) minutes per response, including the time to review instructions, search existing data resources, gather the data needed, and complete and review the information collection. If you have comments concerning the accuracy of the time estimate(s) or suggestions for improving this form, please write to: CMS, 7500 Security Boulevard, Attn: PRA Reports Clearance Officer, Mail Stop C4-26-05, Baltimore, Maryland 21244-1850.



**E. DIAGNOSTIC STUDIES:**

**Ejection Fraction Assessed**<sup>3360</sup>: No; Yes → if Yes, **EF%**<sup>3370</sup>: \_\_\_\_\_ %  
 → if Yes, **EF Timeframe**<sup>3380</sup>: 0-1 month; 1-2 months; 2-3 months; 3-6 months; 6-12 months; >12 months

**Electrophysiology Study Done**<sup>3390</sup>: No; Yes  
 → if Yes, **EPS Timeframe**<sup>3400</sup>: 0-1 month; 1-2 months; 2-3 months; 3-6 months; 6-12 months; >12 months  
 → if Yes, **EPS Findings**<sup>3410</sup>: (Check all that apply. "No Arrhythmias Induced" is mutually exclusive.)  
 No Arrhythmias Induced     VT Induced     Non-sustained VT     Sustained Monomorphic  
 Sustained Polymorphic     Ventricular Flutter Induced     Ventricular Fibrillation Induced     Results Unattainable

**QRS Duration**<sup>3420</sup>: \_\_\_\_\_(msec)    **PR Interval Attainable**<sup>3429</sup> No; Yes → if Yes, **PR Interval**<sup>3430</sup>: \_\_\_\_\_(msec)

**AV Conduction**<sup>3440</sup>: Normal; Abnormal-1<sup>st</sup> Degree Heart Block Only; Abnormal-Heart Block 2<sup>nd</sup> or 3<sup>rd</sup> Degree(not paced); Paced (any)

**Intraventricular Conduction**<sup>3450</sup>:  
 Normal; Abnormal-Left Anterior Fascicular Block; Abnormal-Left Posterior Fascicular Block;  
 Abnormal-LBBB; Abnormal-RBBB; Abnormal-Intraventricular Conduction Delay, Nonspecific;  
 Paced; Abnormal-Bifascicular Block (RBBB Plus LAF); Abnormal-Bifascicular Block (RBBB Plus LPF)

**Creatinine**<sup>3460</sup>: \_\_\_\_\_ **BUN**<sup>3470</sup>: \_\_\_\_\_ **Sodium**<sup>3480</sup>: \_\_\_\_\_ **BNP Drawn**<sup>3485</sup>: No; Yes → if Yes, **BNP**<sup>3490</sup>: \_\_\_\_\_ **Systolic BP**<sup>3500</sup>: \_\_\_\_\_

**F. ICD PROCEDURE:**

**ICD Indication**<sup>3505</sup>: Primary Prevention; Secondary Prevention

**Reason(s) for Re-implantation**<sup>3506</sup>: (if Previous ICD<sup>3250</sup> is Yes) (Check all that apply)  
 End of Battery Life     Device Upgrade     Device Infection     Device Malfunction     Device Under Manufacturer Advisory/Recalled

**Multiple ICDs implanted during this admission**<sup>3507</sup>: No; Yes  
 → If Yes, **Reason(s) for device replacement during this admission**<sup>3508</sup>: (Check all that apply)  
 Device Upgrade     Device Infection     Device Malfunction     Device Under Manufacturer Advisory/Recalled

**Implant Operator's UPIN**<sup>3510</sup>: \_\_\_\_\_ **Implant Operator's NPI**<sup>3515</sup>: \_\_\_\_\_

**Implant Operator's Last Name**<sup>3530</sup>: \_\_\_\_\_ **First Name**<sup>3520</sup>: \_\_\_\_\_ **Middle Name**<sup>3525</sup>: \_\_\_\_\_

**ICD Type**<sup>3540</sup>: Single Chamber; Dual Chamber; Biventricular  
 → If Biventricular, **LV Lead Implant Method**<sup>3550</sup>: Coronary Sinus; Epicardial Lead; Other

	Manufacturer, Model Name, Model Number -or- ICD Device ID <sup>3565/3570</sup>	ICD Serial Number <sup>3566/3571</sup>
<b>Implant:</b>		
if Previous ICD <sup>3250</sup> is Yes then complete <b>Explant</b> below		
<b>Explant:</b>		

**G. ADVERSE EVENTS:** (During or after the implant procedure until discharge.)

**Adverse Events Exist**<sup>3580</sup>: No; Yes → if Yes, then complete **Adverse Events** below.

Adverse Event <sup>3581</sup>		Date <sup>3583</sup>	Adverse Event <sup>3581</sup>		Date <sup>3583</sup>
<b>Cardiac Arrest</b> <sup>ae001</sup> :	<input type="checkbox"/>	___/___/___	<b>Phlebitis - Deep</b> <sup>ae014</sup> :	<input type="checkbox"/>	___/___/___
<b>Drug Reaction</b> <sup>ae002</sup> :	<input type="checkbox"/>	___/___/___	<b>TIA</b> <sup>ae015</sup> :	<input type="checkbox"/>	___/___/___
<b>Cardiac Perforation</b> <sup>ae003</sup> :	<input type="checkbox"/>	___/___/___	<b>CVA/Stroke</b> <sup>ae016</sup> :	<input type="checkbox"/>	___/___/___
<b>Cardiac Valve Injury</b> <sup>ae004</sup> :	<input type="checkbox"/>	___/___/___	<b>MI</b> <sup>ae0017</sup> :	<input type="checkbox"/>	___/___/___
<b>Conduction Block</b> <sup>ae005</sup> :	<input type="checkbox"/>	___/___/___	<b>Pericardial Tamponade</b> <sup>ae018</sup> :	<input type="checkbox"/>	___/___/___
<b>Coronary Venous Dissect</b> <sup>ae006</sup> :	<input type="checkbox"/>	___/___/___	<b>AV Fistula</b> <sup>ae019</sup> :	<input type="checkbox"/>	___/___/___
<b>Hematoma</b> <sup>ae007</sup> :	<input type="checkbox"/>	___/___/___	<b>Infection Related to Device</b> <sup>ae020</sup> :	<input type="checkbox"/>	___/___/___
<b>Lead Dislodgement</b> <sup>ae008</sup> :	<input type="checkbox"/>	___/___/___		<input type="checkbox"/>	___/___/___
<b>Hemothorax</b> <sup>ae009</sup> :	<input type="checkbox"/>	___/___/___		<input type="checkbox"/>	___/___/___
<b>Pneumothorax</b> <sup>ae010</sup> :	<input type="checkbox"/>	___/___/___			
<b>Peripheral Nerve Injury</b> <sup>ae011</sup> :	<input type="checkbox"/>	___/___/___			
<b>Peripheral Embolus</b> <sup>ae012</sup> :	<input type="checkbox"/>	___/___/___			
<b>Phlebitis - Superficial</b> <sup>ae013</sup> :	<input type="checkbox"/>	___/___/___			

**H. DISCHARGE:** (Complete this section at discharge)CABG During this Admission<sup>3590</sup>: No; Yes → if Yes, Date<sup>3600</sup>: \_\_\_\_/\_\_\_\_/\_\_\_\_PCI During this Admission<sup>3610</sup>: No; Yes → if Yes, Date<sup>3620</sup>: \_\_\_\_/\_\_\_\_/\_\_\_\_Vital Status<sup>3630</sup>: Alive; Deceased-Cardiac Death; Deceased-Non-Cardiac Death→ if Deceased, Date<sup>3640</sup>: \_\_\_\_/\_\_\_\_/\_\_\_\_ → if Deceased, Death in Lab<sup>3645</sup>: No; YesDischarge Date<sup>3650</sup>: \_\_\_\_/\_\_\_\_/\_\_\_\_**I. DISCHARGE MEDICATIONS:** (Medications prescribed at discharge.)if Vital Status<sup>3630</sup> is **Alive** then complete **Discharge Medications** below.

Category	Medication Name <sup>3660</sup>	Prescribed <sup>3665</sup>				Category	Medication Name <sup>3660</sup>	Prescribed <sup>3665</sup>				
		No	Yes	Con	Blind			No	Yes	Con	Blind	
Ace Inhibitor	ACE-Inhibitor (any) <sup>m001</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Calcium Channel Blocker	Diltiazem <sup>m016</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
							Verapamil <sup>m017</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Antiarrhythmic Agent	Amiodarone <sup>m002</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Coumadin	Other CCB <sup>m018</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
	Disopyramide <sup>m003</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>		Coumadin	Coumadin <sup>m019</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	Dofetilide <sup>m004</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Digoxin						
	Flecainide <sup>m005</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>		Digoxin	Digoxin <sup>m020</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	Mexiletine <sup>m006</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Diuretic						
	Procainamide <sup>m007</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>		Diuretic	Diuretic (any) <sup>m021</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	Propafenone <sup>m008</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Nitrate						
	Quinidine <sup>m009</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>		Nitrate	Nitroglycerin SL, PRN <sup>m022</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	Sotalol <sup>m010</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>		Nitroglycerin Long Acting <sup>m023</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
	Other Anti. Arrhy. <sup>m011</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>							
Antihypertensive	Hydralazine <sup>m012</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Platelet Aggregation Inhibitor	Clopidogrel <sup>m024</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
							Ticlopidine <sup>m025</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
ARB	ARB (any) <sup>m013</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	Statin						
							Statin (any) <sup>m026</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
ASA	ASA <sup>m014</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>							
Beta Blocker	Beta-Blocker (any) <sup>m015</sup>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>							