



A. DEMOGRAPHICS

Last Name, First Name, Middle Name, SSN, Patient ID, Other ID, Birth Date, Sex, Patient Zip Code, Race, Hispanic or Latino Ethnicity

B. EPISODE OF CARE

Arrival Date, Insurance Payors, HIC, Fundamental Diagnosis Code, Prior Cardiac Catheterization, Research Study, Premature Birth, Birth Weight, Gestational Age, Prior Cardiac Surgery, Patient Restriction

GENETIC/CONGENITAL CONDITIONS (DIAGNOSED PRIOR TO OR DURING THIS EPISODE OF CARE)

22q11Deletion (DiGeorge Syndrome), Noonan Syndrome, Alagille Syndrome, Rubella, Congenital Diaphragmatic Hernia, Trisomy-13, Down Syndrome, Trisomy-18, Heterotaxy, Turner Syndrome, Marfan Syndrome, Williams-Beuren Syndrome

HISTORY & RISK FACTORS

Arrhythmia, Arrhythmia History (Atrial Fibrillation, AV conduction disturbance, Inappropriate sinus tachycardia, Macro re-entrant atrial tachycardia, Supraventricular tachycardia, Wolff-Parkinson-White syndrome, Atrial premature complexes, AV re-entrant tachycardia, Isolated ventricular pre-excitation, Permanent junctional reciprocating tachycardia, Sinus node dysfunction, Wide complex tachycardia, AV node re-entry, Focal atrial tachycardia, Junctional tachycardia, Premature ventricular complexes, Ventricular tachycardia)



B. EPISODE OF CARE (CONT.)

HISTORY & RISK FACTORS

Cardiomyopathy<sup>3170</sup>:  No  Yes

→If Yes, Cardiomyopathy History<sup>3175</sup>:

Arrhythmogenic right ventricular cardiomyopathy  
 Noncompaction of the ventricular myocardium

Dilated cardiomyopathy (DCM)  
 Restrictive cardiomyopathy (RCM)

Hypertrophic cardiomyopathy (HCM)  
 Tachycardia-induced cardiomyopathy

Chronic Lung Disease<sup>3200</sup>:  No  Yes

Heart Transplant<sup>3224</sup>:  No  Yes

Coagulation Disorder<sup>3205</sup>:  No  Yes

Hepatic Disease<sup>3225</sup>:  No  Yes

→If Yes, Hypercoagulable State<sup>3210</sup>:  No  Yes

Ischemic Heart Disease<sup>3226</sup>:  No  Yes

→If Yes, Hypocoagulable State<sup>3215</sup>:  No  Yes

Kawasaki Disease<sup>3227</sup>:  No  Yes

Diabetes Mellitus<sup>3220</sup>:  No  Yes

Renal Insufficiency<sup>3230</sup>:  No  Yes

Endocarditis<sup>3221</sup>:  No  Yes

Rheumatic Heart Disease<sup>3231</sup>:  No  Yes

Heart Failure<sup>3222</sup>: (w/in 1 month)  No  Yes

Seizure Disorder<sup>3235</sup>:  No  Yes

→If Yes, NYHA Class<sup>3223</sup>:  
 Class I  Class II  Class III  Class IV

Sickle Cell Anemia<sup>3240</sup>:  No  Yes

Stroke<sup>3250</sup>: (prior to arrival)  No  Yes

C. CATH LAB VISIT (COMPLETE FOR EACH CATH LAB VISIT)

CLINICAL EVALUATION LEADING TO THE PROCEDURE

Pre-Procedure Diagnosis Code(s)<sup>4000</sup>: \_\_\_\_\_, \_\_\_\_\_, \_\_\_\_\_ Height<sup>4005</sup>: \_\_\_\_\_ cm Weight<sup>4010</sup>: \_\_\_\_\_ kg

Pre-Procedure Labs: Hemoglobin<sup>4015</sup>: \_\_\_\_\_ g/dL  Not Drawn<sup>4016</sup> Creatinine<sup>4020</sup>: \_\_\_\_\_ mg/mL  Not Drawn<sup>4021</sup>  
O<sub>2</sub> Sat<sup>4025</sup>: \_\_\_\_\_ %

Pre-Procedure Conditions: Single Ventricle<sup>4026</sup>:  No  Yes  
Necrotizing Enterocolitis<sup>4030</sup>: (if < 30 days old)  No  Yes  
Sepsis<sup>4035</sup>:  No  Yes  
If Sex<sup>2060</sup> is 'Female', Pregnant<sup>4040</sup>:  No  Yes

Pre-Procedure Medications<sup>4041</sup>:  No  Yes  
→If Yes, (check all that apply)  Antiarrhythmics<sup>4045</sup>  Anticoagulants<sup>4046</sup>  Antihypertensives<sup>4047</sup>  Antiplatelets<sup>4048</sup>  Beta Blockers<sup>4049</sup>  
 Diuretics<sup>4050</sup>  Prostaglandins<sup>4051</sup>  Vasodilators<sup>4053</sup>

Pre-Procedure Rhythms: (check all that apply)  Sinus Rhythm<sup>4060</sup>  Atrial Ectopic Tachycardia (AET)<sup>4061</sup>  Supraventricular Tachycardia (SVT)<sup>4062</sup>  
 AFib/Flutter<sup>4063</sup>  Junctional Rhythm<sup>4064</sup>  Idioventricular Rhythm<sup>4065</sup>  
 Second Degree AV Block<sup>4066</sup>  Third Degree AV Block<sup>4067</sup>  Paced<sup>4068</sup>

D. PROCEDURE INFORMATION (COMPLETE FOR EACH CATH LAB VISIT)

Procedure(s) Performed: (check all that apply)  Diagnostic Cath<sup>5000</sup>  ASD Closure<sup>5001</sup>  Coarctation Procedure<sup>5002</sup>  Aortic Valvuloplasty<sup>5003</sup>  
 Pulmonary Valvuloplasty<sup>5004</sup>  PDA Closure<sup>5005</sup>  Proximal PA Stenting<sup>5006</sup>  
 Electrophysiology Cath<sup>5007</sup>  Electrophysiology Ablation Procedure<sup>5008</sup>  
 Transcatheter Pulmonary Valve Replacement (TPVR)<sup>5009</sup>

Specific Procedure(s)<sup>5010</sup>: \_\_\_\_\_, \_\_\_\_\_, \_\_\_\_\_

Hospital Status<sup>5015</sup>:  Outpatient  Admit to inpatient floor  Admit to inpatient ICU  
 23 Hour obs outpatient  Return to inpatient floor  Return to inpatient ICU

Procedure Status<sup>5020</sup>:  Elective  Urgent  Emergency  Salvage



D. PROCEDURE INFORMATION (COMPLETE FOR EACH CATH LAB VISIT) (CONT.)

Operator's Name<sup>5030, 5031, 5032</sup>: \_\_\_\_\_ Operator's NPI<sup>5035</sup>: \_\_\_\_\_

Trainee participating in the Procedure<sup>5036</sup>:  No  Yes Second Attending participating in the Procedure<sup>5037</sup>:  No  Yes

Procedure Start Date/Time<sup>5047, 5048</sup>: mm / dd / yyyy HH:MM Procedure End Date/Time<sup>5057, 5058</sup>: mm / dd / yyyy HH:MM (break scrub at end of case)

Anesthesiologist Present<sup>5060</sup>: (start of case)  No  Yes
->If No, Anesthesiologist Called In<sup>5065</sup>: (due to escalation of care)  No  Yes

Sedation Method<sup>5070</sup>:  General Anesthesia  Epidural  Caudal  IV  IM  Oral/Intranasal  None

Airway Management<sup>5071</sup>:  No  Yes
->If Yes, (check all that apply)
 Laryngeal mask airway<sup>5076</sup>  Tracheostomy<sup>5077</sup>  Bag mask ventilation<sup>5078</sup>  CPAP<sup>5079</sup>
 Elective intubation<sup>5080</sup>  Previously intubated<sup>5081</sup>

Access Location<sup>5085</sup>:  Venous  Arterial  Both
->If Venous or Both, Venous Access Site<sup>5090</sup>: (check location for largest sheath used)
 Left brachial  Left femoral  Left jugular  Left subclavian  Hepatic  Umbilical
 Right brachial  Right femoral  Right jugular  Right subclavian  Transthoracic  Other
->If Venous or Both, Venous Sheath Size<sup>5095</sup>: \_\_\_\_\_ French (largest)
->If Venous or Both, Venous Closure Method(s)<sup>5100</sup>:
 Method Not Documented<sup>5105</sup>

Table with 3 columns for Venous Closure Method(s) with headers 1, 2, 3.

->If Arterial or Both, Arterial Access Site<sup>5110</sup>: (check location for largest sheath used)
 Left axillary  Left carotid  Left femoral  Left radial  Umbilical
 Right axillary  Right carotid  Right femoral  Right radial  Other
->If Arterial or Both, Arterial Sheath Size<sup>5115</sup>: \_\_\_\_\_ French (largest)
->If Arterial or Both, Arterial Closure Method(s)<sup>5120</sup>:
 Method Not Documented<sup>5125</sup>

Table with 3 columns for Arterial Closure Method(s) with headers 1, 2, 3.

Systemic Heparinization<sup>5140</sup>:  No  Yes
->If Yes, ACT Monitored<sup>5145</sup>:  No  Yes
->If Yes, ACT Peak<sup>5150</sup>: \_\_\_\_\_ secs
Inotrope<sup>5160</sup>:  No  Yes
->If Yes, Inotrope Use<sup>5165</sup>:
 On before case, on at the end  On before case, off at the end
 Started during the case, on at the end  Started during case, off at the end
 Used for measurement only

ECMO Use<sup>5170</sup>:  Not used  In place at start of procedure  Electively initiated during procedure

LVAD Use<sup>5175</sup>:  Not used  In place at start of procedure  Electively initiated during procedure

IABP Use<sup>5180</sup>:  Not used  In place at start of procedure  Electively initiated during procedure

FLUOROSCOPY

X-Ray Imaging<sup>5500</sup>:  Single Plane  Biplane Contrast Volume<sup>5135</sup>: \_\_\_\_\_ mL

CODE ALL AVAILABLE MEASUREMENTS:
Fluoro Time<sup>5130</sup>: \_\_\_\_\_ minutes
-> Cumulative Air Kerma<sup>5515, 5520</sup>: \_\_\_\_\_ O mGy O Gy
Dose Area Product<sup>5525, 5530</sup>: \_\_\_\_\_ O Gy-cm<sup>2</sup> O dGy-cm<sup>2</sup> O cGy-cm<sup>2</sup> O mGy-cm<sup>2</sup> O uGy-M<sup>2</sup>



E. HEMODYNAMICS (COMPLETE FOR EACH CATH LAB VISIT)

Systemic Arterial Saturation<sup>6000</sup>: \_\_\_\_\_ %  Not Assessed<sup>6001</sup>

Mixed Venous Saturation<sup>6005</sup>: \_\_\_\_\_ %  Not Assessed<sup>6006</sup>

Systemic Ventricular Systolic Pressure<sup>6010</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6011</sup>

Systemic Ventricular End Diastolic Pressure<sup>6015</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6016</sup>

Systemic Blood Pressure: (Systolic)<sup>6020</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6021</sup> (Mean)<sup>6030</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6031</sup>  
 (Diastolic)<sup>6025</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6026</sup>

PA Pressure: (Systolic)<sup>6035</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6036</sup> (Mean)<sup>6040</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6041</sup>

Pulmonary Ventricular Systolic Pressure<sup>6045</sup>: \_\_\_\_\_ mmHg  Not Assessed<sup>6046</sup>

Pulmonary Vascular Resistance Index<sup>6050</sup>: \_\_\_\_\_ Wood Units\*m<sup>2</sup>  Not Assessed<sup>6051</sup>

Cardiac Index<sup>6055</sup>: \_\_\_\_\_ L/min/m<sup>2</sup>  Not Assessed<sup>6056</sup> Qp/Qs ratio<sup>6060</sup>: \_\_\_\_\_  Not Assessed<sup>6061</sup>

F. ASD CLOSURE

Primary Procedure Indication<sup>7000</sup>:  
 Right ventricular volume overload  Chronic lung disease  Failure to thrive  
 Recurrent respiratory infections  Ventilator dependent  Cyanosis  
 Stroke prevention  Migraines  Pulmonary hypertension

Total Septal Length<sup>7005</sup>: \_\_\_\_\_ mm  Not Assessed<sup>7006</sup> Atrial Septal Aneurysm Present<sup>7010</sup>:  No  Yes

DEFECT COUNTER <sup>7020</sup>	1	2	3
ASD Multi-Fenestrated <sup>7022</sup> : →If No, ASD Size <sup>7025</sup> : _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm
Balloon Sizing Performed <sup>7030</sup> : →If Yes, Stretched Diameter Performed <sup>7035</sup> : →If Yes, Size <sup>7040</sup> : _____ mm →If Yes, Stop Flow Technique Performed <sup>7045</sup> : →If Yes, Size <sup>7050</sup> : _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm <input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm
Rim Measurement Performed <sup>7055</sup> : →If Yes, IVC Rim Length <sup>7060</sup> : _____ mm →If Yes, Minimum Aortic Rim Length <sup>7065</sup> : _____ mm →If Yes, Posterior Rim Length <sup>7066</sup> : _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm _____ mm _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm _____ mm _____ mm	<input type="checkbox"/> No <input type="checkbox"/> Yes _____ mm _____ mm _____ mm
Residual Shunt Size <sup>7080</sup> : (immed after device placement)	<input type="checkbox"/> None to trivial (<3 mm) <input type="checkbox"/> Significant (>=3 mm)	<input type="checkbox"/> None to trivial (<3 mm) <input type="checkbox"/> Significant (>=3 mm)	<input type="checkbox"/> None to trivial (<3 mm) <input type="checkbox"/> Significant (>=3 mm)

	Device(s) <sup>7085</sup>	Associated Defect(s) <sup>7089</sup>	Outcome of Device <sup>7090</sup>
1	_____ , _____ , _____	_____ , _____ , _____	<input type="checkbox"/> Implanted, not released <input type="checkbox"/> Implanted, released <input type="checkbox"/> Implanted, released and retrieved
2	_____ , _____ , _____	_____ , _____ , _____	<input type="checkbox"/> Implanted, not released <input type="checkbox"/> Implanted, released <input type="checkbox"/> Implanted, released and retrieved
3	_____ , _____ , _____	_____ , _____ , _____	<input type="checkbox"/> Implanted, not released <input type="checkbox"/> Implanted, released <input type="checkbox"/> Implanted, released and retrieved



G. COARCTATION PROCEDURE

Primary Procedure Indication 7100:
O Abnormal ventricular function
O High resting gradient
O Congestive heart failure
O Angiographic appearance
O Exercise hypertension
O Pseudoaneurysm
O Systemic hypertension

Nature of simple discrete coarctation (One site of intervention) 7101:
O Native
O Post Treatment
->If Post Treatment, Most Recent Prior Treatment 7102:
O Surgical Repair
O Catheter-based Intervention

Pre-Procedure Minimal Diameter 7107: \_\_\_\_\_ mm
[ ] Not Assessed 7108

Pre-Procedure Peak Systolic Gradient 7110: \_\_\_\_\_ mmHg
[ ] Not Assessed 7111

Post-Procedure Minimal Diameter 7120: \_\_\_\_\_ mm
[ ] Not Assessed 7121

Post-Procedure Peak Systolic Gradient 7125: \_\_\_\_\_ mmHg
[ ] Not Assessed 7124

Coarctation with additional associated aortic obstruction 7126:
O No
O Yes
->If Yes, Additional intervention on aortic arch 7127:
O No
O Yes
->If Yes, Pre-Procedure Total ascending to descending Aortic Systolic Gradient 7128: \_\_\_\_\_ mmHg
->If Yes, Post-Procedure Total ascending to descending Aortic Systolic Gradient 7129: \_\_\_\_\_ mmHg

DEVICE COUNTER 7130

Table with 4 columns: Device Counter, 1, 2, 3. Rows include Device ID 7135, Device Type 7140, and various procedural details like Purpose 7145, Max Inflation Pressure 7150, Outcome 7155, and In Stent Minimal Diameter Assessed 7164.



H. AORTIC VALVULOPLASTY

Primary Procedure Indication 7200: O Aortic stenosis gradient O Abnormal stress test/EKG
O LV dysfunction O Symptoms

Valve Morphology 7205: O Unicuspid O Bicuspid O Tricuspid O Quadracuspid O Uncertain

Pre-Procedure Aortic Valve Regurgitation 7210: O None O 1+ (mild) O 2+ (moderate) O 3+ (moderately severe) O 4+ (severe)

Aortic Valve Diameter 7215: (used to select balloon) \_\_\_\_\_ mm Pre-Procedure Peak Systolic Gradient 7220: \_\_\_\_\_ mmHg

Table with 4 columns: BALLOON COUNTER 7231, 1, 2, 3. Rows include Balloon Technique 7236, Balloon Stabilization 7243, Max Inflation Pressure 7244, Balloon Outcome 7256, Post Dilatation Systolic Gradient 7257, and Post Dilatation Regurgitation 7258.

I. PULMONARY VALVULOPLASTY

Primary Procedure Indication 7400: O High resting gradient O R to L shunting O RV dysfunction O Symptoms

Valve Morphology 7405: O Typical O Dysplastic/Complex Subpulmonary Stenosis Present 7410: O No O Yes

Pulmonary Valve Diameter 7415: (used to select balloon) \_\_\_\_\_ mm

Pre-Procedure Peak Systolic Gradient 7420: \_\_\_\_\_ mmHg O Not Assessed 7421

Balloon Technique (Final Balloon) 7520: O Single O Double

->If Single or Double, Device ID Balloon 1 7525: \_\_\_\_\_

->If Double, Device ID Balloon 2 7530: \_\_\_\_\_

Balloon Stabilization 7535: O No O Yes

Max Inflation Pressure 7540: \_\_\_\_\_ atm(s)

Balloon Outcome 7545: O Inflated with rupture O Inflated without rupture

Post-Procedure Peak Systolic Gradient 7550: \_\_\_\_\_ mmHg O Not Assessed 7551

J. PDA CLOSURE

Primary Procedure Indication 7600: O SBE prevention O Left ventricular volume overload O Pulmonary hypertension

PDA Diameter Aortic Side 7605: \_\_\_\_\_ mm PDA Minimum Luminal Diameter 7610: \_\_\_\_\_ mm PDA Length 7615: \_\_\_\_\_ mm

PDA Classification 7620: O Type A (conical) O Type B (window) O Type C (tubular) O Type D (complex) O Type E (elongated)

PA Obstruction 7630: (caused by implant) O No O Yes O Not Assessed

Aortic Obstruction 7635: (caused by implant) O No O Yes O Not Assessed

Residual Shunt 7640: (immed after device placement) O None to trivial O Significant

Table with 4 columns: Device(s) 7645, Outcome of Device 7650, and three columns for device status: O Implanted, not released; O Implanted, released; O Implanted, released and retrieved.



K. PROXIMAL PULMONARY ARTERY STENTING PROCEDURE

Primary Procedure Indication 7700: O PA gradient O PA flow discrepancy O RV hypertension/dysfunction O Pulmonary regurgitation O PA angiographic narrowing

Table with 4 columns: Defect Counter 7705, Defect Location 7710, Distal Obstruction Present 7720, Sidebranch Jailing 7725, and associated flow/pressure indicators.

PRE-PROCEDURE MEASUREMENTS (TO PA DEFECT)

Table for pre-procedure measurements including Proximal Systolic Pressure, Distal Systolic Pressure, Proximal Mean Pressure, Distal Mean Pressure, Proximal Diameter, Distal Diameter, and PA Vessel Diameter Minimum.

POST-PROCEDURE MEASUREMENTS (TO PA DEFECT)

Table for post-procedure measurements including Proximal Systolic Pressure, Distal Systolic Pressure, Proximal Mean Pressure, Distal Mean Pressure, Proximal Diameter, Distal Diameter, and PA Vessel Diameter in Stent Minimum.

Table with 3 columns: Device(s) 7820, Associated Defect(s) 7824, and Outcome of Device 7825, containing rows 1, 2, and 3.



L. ELECTROPHYSIOLOGY PROCEDURE

Primary Procedure Indication 10000: O Evaluation of specific arrhythmia O Evaluation of event or symptoms suggesting arrhythmia
O Evaluation of prior antiarrhythmic treatment O Evaluation of risk for ventricular tachyarrhythmia
O Preoperative evaluation

PRIOR ELECTROPHYSIOLOGY HISTORY

History of Congenital Heart Disease 10005: O No structural heart disease or trivial, unoperated congenital heart disease
O Repaired functionally two-ventricle congenital heart disease
O Repaired tetralogy of Fallot and tetralogy-like variants
O Transposition of the great arteries following atrial-level (Mustard or Senning) palliation
O Fontan palliation of functionally univentricular heart
O Pre-Fontan palliation of functionally univentricular heart
O Unoperated acyanotic congenital heart disease
O Unoperated cyanotic congenital heart disease

Previous EP Therapy Attempted 10010: O No O Yes
->If Yes, EP therapy(ies) attempted: O Catheter Ablation 10011 O Pharmacologic Therapy 10012 O Chemical cardioversion 10013
(check all that apply) O DC cardioversion 10014 O Pacemaker insertion 10015 O ICD insertion 10016
O Arrhythmia surgery 10017
->If Catheter Ablation, Number of prior ablation procedures 10018: \_\_\_\_\_

PRE-PROCEDURE SYMPTOM SEVERITY SURVEY (SSS)

SSSQ1: In the past 6 months, what symptom or feeling that comes from the patient's heart rhythm problem bothers her/him most often? 10020

O Palpitations O Chest pain O Shortness of breath O Dizziness O Fatigue O Fainting O No symptoms

->If any symptoms present, SSSQ2: In the past 6 months how often has patient had this feeling? 10021

O Every day O At least once per week O At least once per month O At least once in the last 6 months

SSSQ3: In the past 6 months, what symptom or feeling that comes from the patient's heart rhythm problem was the worst (most severe or unpleasant)? 10022

O Palpitations O Chest pain O Shortness of breath O Dizziness O Fatigue O Fainting O No symptoms

SSSQ4: In the past 6 months, for any heart rhythm episodes the patient has had, what is the most intense (severe) treatment that the patient has endured to treat the rhythm problem? 10023

O No rhythm problems during this time O Rhythm is always present and no effort was made to try and relieve it O Self-Resolving
O Vagal Maneuvers O ER visit, symptoms self-resolved or with vagal maneuvers
O ER-Treated with medication O Admitted for >= 1 day, treated with medication O Hospital/ER Cardioversion

SSSQ5: In the past 6 months, has the patient taken any of the following medications? 10024 (check all that apply)

O Amiodarone O Beta Blocker O Digoxin O Diltiazem O Dofetilide O Dronedaron O Flecainide O Mexiletine
O Propafenone O Sotalol O Verapamil O None

SSSQ6: In the past 6 months, does the patient feel that their rhythm problem has interfered with how well they are able to work, go to school or play? 10025 O No O Yes

SSSQ7: Indicate who completed the Symptom Severity Survey (SSS)? 10026 O Caregiver O Parent O Patient

Tachyarrhythmias Observed during EP Study 10040: (check all that apply)

O Atrial Fibrillation O Atrial Flutter - CTI-dependent O Atrial Flutter - Non-CTI-dependent
O Atrial premature complexes O AV node re-entry Typical (slow/fast) O AV node re-entry Atypical (fast/slow)
O AV node re-entry Atypical (slow/slow) O AV node re-entry Atypical (unknown) O AVRT - antidromic
O AVRT - orthodromic O Ectopic atrial tachycardia O Inappropriate sinus tachycardia
O Isolated ventricular pre-excitation O Junctional tachycardia O Premature ventricular complexes
O Ventricular fibrillation O Ventricular tachycardia, monomorphic O Ventricular tachycardia, monomorphic, non-sustained
O Ventricular tachycardia, polymorphic O Ventricular tachycardia, polymorphic, non-sustained
O No tachyarrhythmias or ectopy observed

Sedation Medication 10065: O Cisatracurium O Desflurane O Dexmedetomidine O Fentanyl O Isoflurane O Ketamine
(check all that apply) O Midazolam O Morphine O Nitrous oxide O Propofol O Remifentanyl O Rocuronium
O Sevoflurane O Succinylcholine O Vecuronium



L. ELECTROPHYSIOLOGY PROCEDURE (CONT.)

Imaging System(s) Used (check all that apply):
[ ] CARTO 3 [ ] CARTO XP [ ] CARTO Sound [ ] ICE [ ] Ensite NavX
[ ] Velocity NavX [ ] EnSite Balloon Array [ ] Velocity Balloon Array [ ] TEE [ ] None

ABLATION PROCEDURE

Table with 3 columns: Ablation Target Counter, Indications for Ablation, Approach to Ablation Target, Targeted ablation substrate, Ablation Target Location ID, Methods to localize ablation target, Ablation Attempted. Each row contains checkboxes and text for clinical data entry.



L. ELECTROPHYSIOLOGY PROCEDURE (CONT.)

Ablation Target Counter <sup>10075</sup>	1	2
<b>EP – Outcome of Ablation<sup>10110</sup></b> (Select from below)		
→If Accessory pathway - concealed, <b>Outcome:</b>	<input type="radio"/> Elimination of retrograde AP conduction <input type="radio"/> Persistence of retrograde conduction, without SVT <input type="radio"/> Persistence of retrograde conduction, with SVT <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of retrograde AP conduction <input type="radio"/> Persistence of retrograde conduction, without SVT <input type="radio"/> Persistence of retrograde conduction, with SVT <input type="radio"/> Unknown or Ambiguous
→If Accessory pathway - manifest (bidirectional WPW), <b>Outcome:</b>	<input type="radio"/> Elimination of antegrade conduction (Elimination of retrograde conduction) <input type="radio"/> Elimination of antegrade conduction (Persistence of retrograde conduction, with SVT) <input type="radio"/> Elimination of antegrade conduction (Persistence of retrograde conduction, without SVT) <input type="radio"/> Persistence of antegrade conduction (Elimination of retrograde conduction, with SVT) <input type="radio"/> Persistence of antegrade conduction (Elimination of retrograde conduction, without SVT) <input type="radio"/> Persistence of antegrade conduction (Persistence of retrograde conduction, with SVT) <input type="radio"/> Persistence of antegrade conduction (Persistence of retrograde conduction, without SVT) <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of antegrade conduction (Elimination of retrograde conduction) <input type="radio"/> Elimination of antegrade conduction (Persistence of retrograde conduction, with SVT) <input type="radio"/> Elimination of antegrade conduction (Persistence of retrograde conduction, without SVT) <input type="radio"/> Persistence of antegrade conduction (Elimination of retrograde conduction, with SVT) <input type="radio"/> Persistence of antegrade conduction (Elimination of retrograde conduction, without SVT) <input type="radio"/> Persistence of antegrade conduction (Persistence of retrograde conduction, with SVT) <input type="radio"/> Persistence of antegrade conduction (Persistence of retrograde conduction, without SVT) <input type="radio"/> Unknown or Ambiguous
→If Accessory pathway manifest (antegrade only WPW), <b>Outcome:</b>	<input type="radio"/> Elimination of antegrade AP conduction <input type="radio"/> Persistence of antegrade conduction, without SVT <input type="radio"/> Persistence of antegrade conduction, with SVT <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of antegrade AP conduction <input type="radio"/> Persistence of antegrade conduction, without SVT <input type="radio"/> Persistence of antegrade conduction, with SVT <input type="radio"/> Unknown or Ambiguous
→If Accessory pathway - manifest (unidirectional anterograde decremental pathway - Mahaim), <b>Outcome:</b>	<input type="radio"/> Elimination of antegrade AP conduction <input type="radio"/> Persistence of antegrade conduction, without SVT <input type="radio"/> Persistence of antegrade conduction, with SVT <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of antegrade AP conduction <input type="radio"/> Persistence of antegrade conduction, without SVT <input type="radio"/> Persistence of antegrade conduction, with SVT <input type="radio"/> Unknown or Ambiguous
→If AV node, <b>Outcome:</b>	<input type="radio"/> Ablation ineffective <input type="radio"/> Attenuation of AV conduction <input type="radio"/> Elimination of AV conduction <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Ablation ineffective <input type="radio"/> Attenuation of AV conduction <input type="radio"/> Elimination of AV conduction <input type="radio"/> Unknown or Ambiguous
→If AV node (fast pathway), <b>Outcome:</b>	<input type="radio"/> Elimination of fast pathway conduction (Attenuation of AV conduction) <input type="radio"/> Elimination of fast pathway conduction (Elimination of AV conduction) <input type="radio"/> Elimination of fast pathway conduction (No change in AV conduction) <input type="radio"/> Persistence of fast pathway conduction (Attenuation of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (Elimination of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (No change in AV conduction) <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of fast pathway conduction (Attenuation of AV conduction) <input type="radio"/> Elimination of fast pathway conduction (Elimination of AV conduction) <input type="radio"/> Elimination of fast pathway conduction (No change in AV conduction) <input type="radio"/> Persistence of fast pathway conduction (Attenuation of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (Elimination of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (No change in AV conduction) <input type="radio"/> Unknown or Ambiguous
→If AV node (slow pathway), <b>Outcome:</b>	<input type="radio"/> Elimination of slow pathway conduction <input type="radio"/> Persistence of slow pathway conduction (Persistence of spontaneous or inducible SVT) <input type="radio"/> Persistence of slow pathway conduction (with single echoes but no SVT) <input type="radio"/> Persistence of slow pathway conduction (without echoes) <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of slow pathway conduction <input type="radio"/> Persistence of slow pathway conduction (Persistence of spontaneous or inducible SVT) <input type="radio"/> Persistence of slow pathway conduction (with single echoes but no SVT) <input type="radio"/> Persistence of slow pathway conduction (without echoes) <input type="radio"/> Unknown or Ambiguous



L. ELECTROPHYSIOLOGY PROCEDURE (CONT.)

Ablation Target Counter <sup>10075</sup>		1	2	
EP – Outcome of Ablation <sup>10110</sup> (Select from below)				
→If His bundle, Outcome:	<input type="radio"/> Elimination of ectopic focus/tachycardia (Attenuation of AV conduction) <input type="radio"/> Elimination of ectopic focus/tachycardia (Elimination of AV conduction) <input type="radio"/> Elimination of ectopic focus/tachycardia (No change in AV conduction) <input type="radio"/> Persistence of ectopic focus/tachycardia (Attenuation of AV conduction) <input type="radio"/> Persistence of ectopic focus/tachycardia (Elimination of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (No change in AV conduction) <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of ectopic focus/tachycardia (Attenuation of AV conduction) <input type="radio"/> Elimination of ectopic focus/tachycardia (Elimination of AV conduction) <input type="radio"/> Elimination of ectopic focus/tachycardia (No change in AV conduction) <input type="radio"/> Persistence of ectopic focus/tachycardia (Attenuation of AV conduction) <input type="radio"/> Persistence of ectopic focus/tachycardia (Elimination of AV conduction) <input type="radio"/> Persistence of fast pathway conduction (No change in AV conduction) <input type="radio"/> Unknown or Ambiguous		
→If Myocardium - atrial, Outcome:	<input type="radio"/> Substrate eliminated <input type="radio"/> Substrate attenuated <input type="radio"/> Ablation ineffective <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Substrate eliminated <input type="radio"/> Substrate attenuated <input type="radio"/> Ablation ineffective <input type="radio"/> Unknown or Ambiguous		
→If Myocardium - coronary sinus, Outcome:	<input type="radio"/> Substrate eliminated <input type="radio"/> Substrate attenuated <input type="radio"/> Ablation ineffective <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Substrate eliminated <input type="radio"/> Substrate attenuated <input type="radio"/> Ablation ineffective <input type="radio"/> Unknown or Ambiguous		
→If Myocardium-ventricular, Outcome:	<input type="radio"/> Elimination of spontaneous/inducible VT <input type="radio"/> Persistence of spontaneous/inducible VT (with non-sustained VT) <input type="radio"/> Persistence of spontaneous/inducible VT (with sustained VT) <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Elimination of spontaneous/inducible VT <input type="radio"/> Persistence of spontaneous/inducible VT (with non-sustained VT) <input type="radio"/> Persistence of spontaneous/inducible VT (with sustained VT) <input type="radio"/> Unknown or Ambiguous		
→If Sinus node, Outcome:	<input type="radio"/> Normalization of sinus node function <input type="radio"/> Persistent inappropriate sinus tachycardia <input type="radio"/> Sinus bradycardia or arrest <input type="radio"/> Unknown or Ambiguous	<input type="radio"/> Normalization of sinus node function <input type="radio"/> Persistent inappropriate sinus tachycardia <input type="radio"/> Sinus bradycardia or arrest <input type="radio"/> Unknown or Ambiguous		
	Ablation Catheter(s) <sup>10120</sup>	Associated Ablation Targets(s) <sup>10125</sup>	Seconds on Ablation Target <sup>10130</sup>	Number of Ablation Activations <sup>10135</sup>
1		_____ , _____ , _____		
2		_____ , _____ , _____		
3		_____ , _____ , _____		



M. TRANSCATHETER PULMONARY VALVE REPLACEMENT (TPVR)

Clinical Indication<sup>11000</sup> :

Symptomatic  Prevention of symptoms in asymptomatic patient  Declining ventricular function  Worsening arrhythmias  Other

Hemodynamic Indication<sup>11005</sup> :

Predominant valve/conduit Obstruction  Predominant valve/conduit Regurgitation  Mixed obstruction/regurgitation

Underlying anatomic reason for Right Ventricular Outflow Tract (RVOT) dysfunction<sup>11010</sup>:

Congenital Heart Disease repaired using RVOT valve/conduit  s/p Ross Procedure with repair using RVOT valve/conduit  
 No Congenital Heart Disease with RVOT valve/conduit  Native RVOT dysfunction secondary to surgical intervention  
 Native RVOT dysfunction secondary to transcatheter intervention (other than preparation for transcatheter valve)  
 Native RVOT dysfunction with no prior interventions

PRE-PROCEDURE TESTING

Echocardiogram<sup>11015</sup>:  No  Yes

→If Yes,

Mean gradient across valve/conduit<sup>11016</sup>: \_\_\_\_\_ mmHg

Maximum gradient across valve/conduit<sup>11017</sup>: \_\_\_\_\_ mmHg

Pulmonary Valve Regurgitation<sup>11018</sup>:  None  1+ (mild)  2+ (moderate)  3+ (moderately severe)  4+ (severe)

LVEF<sup>11019</sup>: \_\_\_\_\_ %

Tricuspid Regurgitation Severity<sup>11020</sup>:  None  1+ (mild)  2+ (moderate)  3+ (moderately severe)  4+ (severe)

MRI<sup>11030</sup>:  No  Yes

→If Yes,

RVEF<sup>11031</sup>: \_\_\_\_\_ %

LVEF<sup>11032</sup>: \_\_\_\_\_ %

PR Fraction<sup>11037</sup>: \_\_\_\_\_ %

RVEDV Index<sup>11033</sup>: \_\_\_\_\_ ml/m<sup>2</sup>

RVESV Index<sup>11034</sup>: \_\_\_\_\_ ml/m<sup>2</sup>

LVEDV Index<sup>11035</sup>: \_\_\_\_\_ ml/m<sup>2</sup>

LVESV Index<sup>11036</sup>: \_\_\_\_\_ ml/m<sup>2</sup>

RIGHT VENTRICULAR OUTFLOW TRACT (RVOT) ANATOMY AND FUNCTION

Type of RVOT valve/conduit<sup>11040</sup>:  Homograft (aortic)  Homograft (pulmonary)  Homograft (unknown)  Contegra  
 Bioprosthetic valve/conduit  Non-valved synthetic tube  Native/patched RVOT

→If Not Native/patched RVOT, Surgically implanted valve/conduit size<sup>11041</sup>: \_\_\_\_\_ mm

Existing stent within valve/conduit<sup>11045</sup>:  No  Yes

Prior TPVR (Valve-in-Valve)<sup>11050</sup>:  No  Yes

Cath Peak gradient across valve/conduit<sup>11055</sup>: \_\_\_\_\_ mmHg

Narrowest angiographic valve/conduit diameter<sup>11060</sup>: \_\_\_\_\_ mm

CORONARY ARTERY ASSESSMENT

Aortography performed<sup>11065</sup>:  No  Yes

Selective coronary angiography performed<sup>11070</sup>:  No  Yes

Coronary compression testing performed<sup>11075</sup>:  No  Yes

→If Yes, Max Balloon size<sup>11076</sup>: \_\_\_\_\_ mm

→If Yes, Coronary compression present<sup>11077</sup>:  No  Yes  Uncertain



M. TRANSCATHETER PULMONARY VALVE REPLACEMENT (CONT.)

CONDUIT PREPARATION

Pre-dilation performed<sup>11080</sup>:  No  Yes

→If Yes, First Balloon size<sup>11081</sup>: \_\_\_\_\_ mm

→If Yes, Maximum Balloon size<sup>11082</sup>: \_\_\_\_\_ mm

→If Yes, Highest pressure inflation performed<sup>11083</sup>: \_\_\_\_\_ atm(s)

New Pre-Stent implanted<sup>11085</sup>:  No  Yes

→If Yes, Number of new stents<sup>11086</sup>: \_\_\_\_\_

Access vessel for valve delivery<sup>11090</sup>:  Femoral Vein  Jugular Vein  Subclavian Vein  Per ventricular  Other

Delivery Balloon size<sup>11095</sup>: \_\_\_\_\_ mm

Transcatheter Pulmonary Valve (TPV) deployed<sup>11100</sup>:  No  Yes

→If Yes, Post-dilation of TPV<sup>11101</sup>:  No  Yes

→If Yes, Final Balloon size<sup>11102</sup>: \_\_\_\_\_ mm

→If Yes, Final Balloon Pressure<sup>11103</sup>: \_\_\_\_\_ atm(s)

→If Yes, Post-Procedure Peak RVOT gradient<sup>11105</sup>: \_\_\_\_\_ mmHg

→If Yes, Post-Procedure Pulmonary Valve Regurgitation<sup>11110</sup>:  None  1+ (mild)  2+ (moderate)  3+ (moderately severe)  4+ (severe)

→If Yes, Final minimal diameter of valve<sup>11115</sup>: \_\_\_\_\_ mm

→If No, Reason TPV not deployed<sup>11120</sup>:

- Not indicated based on invasive hemodynamics
- Coronary artery compression risk
- Pre-stent implanted, planned TPVR at a later date
- Other
- Other treatment performed instead with adequate result
- Valve could not be advanced to implant location
- Patient unstable
- Complication before deployment
- No treatable landing zone

	Device(s) <sup>11130</sup>	Outcome of Device <sup>11135</sup>
1		<input type="radio"/> Implanted in intended location <input type="radio"/> Implanted, not released <input type="radio"/> Implanted in unintended location <input type="radio"/> Implanted, released and retrieved
2		<input type="radio"/> Implanted in intended location <input type="radio"/> Implanted, not released <input type="radio"/> Implanted in unintended location <input type="radio"/> Implanted, released and retrieved
3		<input type="radio"/> Implanted in intended location <input type="radio"/> Implanted, not released <input type="radio"/> Implanted in unintended location <input type="radio"/> Implanted, released and retrieved

POST-PROCEDURE TESTING (POST PROCEDURE AND PRIOR TO DISCHARGE)

Echocardiogram<sup>11140</sup>:  No  Yes

→If Yes, Mean gradient across valve/conduit<sup>11145</sup>: \_\_\_\_\_ mmHg

→If Yes, Maximum gradient across valve/conduit<sup>11150</sup>: \_\_\_\_\_ mmHg

→If Yes, Pulmonary Valve Regurgitation<sup>11155</sup>:  None  1+ (mild)  2+ (moderate)  3+ (moderately severe)  4+ (severe)



N. INTRA AND POST-PROCEDURE EVENTS (COMPLETE FOR EACH CATH LAB VISIT)

<b>Cardiac Arrest</b> <sup>8000</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Bleeding Event</b> <sup>8090</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Arrhythmia</b> <sup>8005</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Bleeding at Access Site</b> <sup>8095</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>AV Block</b> <sup>8006</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Hematoma at Access Site</b> <sup>8100</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Spontaneously resolved</b> <sup>8007</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Retroperitoneal Bleeding</b> <sup>8110</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Antiarrhythmic Medication</b> <sup>8010</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>GI Bleed</b> <sup>8115</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Cardioversion</b> <sup>8015</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>GU Bleed</b> <sup>8120</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Temporary Pacemaker</b> <sup>8020</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Other Bleed</b> <sup>8125</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Permanent Pacemaker</b> <sup>8025</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>RBC Transfusion</b> <sup>8130</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>New Heart Valve Regurgitation</b> <sup>8030</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Drop in Hgb ≥ 3</b> <sup>8131</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Tamponade</b> <sup>8035</sup> : (req pericardial drainage)	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Anemia prior to Cath Procedure</b> <sup>8132</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Air Embolus</b> <sup>8040</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Post-operative Blood Loss</b> <sup>8133</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Embolic Stroke</b> <sup>8045</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>ECMO Blood Replacement</b> <sup>8134</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Device Malposition or Thrombus</b> <sup>8050</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Other Vascular Complications Req Rx</b> <sup>8140</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Retrieved via Catheterization</b> <sup>8051</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Other Events</b> <sup>8145</sup> : (optional)	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Retrieved via Surgery</b> <sup>8052</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, select <b>Event(s)</b> <sup>8150</sup> from list: _____, _____, _____	
<b>Device Embolization</b> <sup>8055</sup> : (req device retrieval)	<input type="radio"/> No <input type="radio"/> Yes	<b>Peripheral Nerve Injury</b> <sup>8200</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Retrieved via Catheterization</b> <sup>8060</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Phrenic Nerve Paralysis</b> <sup>8205</sup> :	<input type="radio"/> No <input type="radio"/> Yes
→If Yes, <b>Retrieved via Surgery</b> <sup>8065</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Pneumothorax</b> <sup>8210</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>New Requirement for Dialysis</b> <sup>8070</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Pulmonary Embolism</b> <sup>8215</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Coronary Artery Compression</b> <sup>8071</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Pulmonary Vein Stenosis</b> <sup>8220</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Erosion</b> <sup>8072</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Radiation Burn to Skin</b> <sup>8225</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Esophageal Fistula</b> <sup>8073</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Deep Vein Thrombosis</b> <sup>8230</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>LBBB</b> <sup>8074</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Conduit Tear</b> <sup>8235</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>RBBB</b> <sup>8076</sup> :	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Location</b> <sup>8236</sup> :	
<b>Airway Event Requiring Escalation of Care</b> <sup>8075</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<input type="radio"/> Confined or therapeutic tear without hemodynamic change	
<b>Event Requiring ECMO</b> <sup>8080</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<input type="radio"/> Rupture into pericardial or pleural space	
<b>Event Requiring LVAD</b> <sup>8085</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<input type="radio"/> Rupture into bronchus, cardiac chamber, aorta, or other vessel	
		→If Yes, <b>Treatment</b> <sup>8237</sup> : (check all that apply)	
		<input type="checkbox"/> No specific treatment	
		<input type="checkbox"/> Pericardial or pleural drain	
		<input type="checkbox"/> Covered with TPV	
		<input type="checkbox"/> Other catheter device (covered stent, occluder, coils)	
		<input type="checkbox"/> Surgery	

POST-PROCEDURE TREATMENTS

<b>Planned Cardiac Surgery</b> <sup>8155</sup> :	<input type="radio"/> No <input type="radio"/> Yes	<b>Unplanned Other Surgery</b> <sup>8170</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Unplanned Cardiac Surgery</b> <sup>8160</sup> : (due to cath complication)	<input type="radio"/> No <input type="radio"/> Yes	→If Yes, <b>Due to Cath Complication</b> <sup>8175</sup> :	<input type="radio"/> No <input type="radio"/> Yes
<b>Unplanned Vascular Surgery</b> <sup>8165</sup> : (due to cath complication)	<input type="radio"/> No <input type="radio"/> Yes	<b>Subsequent Cardiac Cath</b> <sup>8180</sup> : (due to cath complication)	<input type="radio"/> No <input type="radio"/> Yes



O. DISCHARGE (COMPLETE FOR EACH EPISODE OF CARE)

Cardiac Surgery during this admission<sup>8305</sup>:  No  Yes → If Yes, Cardiac Surgery Date/Time<sup>8310,8315</sup>: mm / dd / yyyy HH:MM

Discharge Date<sup>9000</sup>: mm / dd / yyyy

Discharge Status<sup>9005</sup>:  Alive  Deceased

→ If Deceased, Death in Lab<sup>9010</sup>:  No  Yes

→ If Deceased, Primary Cause of Death<sup>9015</sup>:

- Acute myocardial infarction, Sudden cardiac death, Heart failure, Stroke, Cardiovascular procedure, Cardiovascular hemorrhage, Other cardiovascular reason, Pulmonary, Renal, Gastrointestinal, Hepatobiliary, Pancreatic, Infection, Inflammatory/Immunologic, Hemorrhage, Non-cardiovascular procedure or surgery, Trauma, Suicide, Neurological, Malignancy, Other non-cardiovascular reason

P. FOLLOW-UP (COMPLETE AFTER DISCHARGE FROM FACILITY)

Assessment Date<sup>12000</sup>: mm / dd / yyyy

Reference Procedure Start Date/Time<sup>12001/12002</sup>: mm / dd / yyyy HH:MM

Method(s) to Determine Status:  Office Visit<sup>12005</sup>  Medical Records<sup>12006</sup>  Letter from Medical Provider<sup>12007</sup>  Phone Call<sup>12008</sup>  Social Security Death Master File<sup>12009</sup>  Hospitalized<sup>12010</sup>  Other<sup>12011</sup>

Follow-up Status<sup>12015</sup>:  Alive  Deceased  Lost to Follow-up

→ If Deceased, Date of Death<sup>12020</sup>: mm / dd / yyyy

→ If Deceased, Cause of Death<sup>12025</sup>:

- Acute myocardial infarction, Sudden cardiac death, Heart failure, Stroke, Cardiovascular procedure, Cardiovascular hemorrhage, Other cardiovascular reason, Pulmonary, Renal, Gastrointestinal, Hepatobiliary, Pancreatic, Infection, Inflammatory/Immunologic, Hemorrhage, Non-cardiovascular procedure or surgery, Trauma, Suicide, Neurological, Malignancy, Other non-cardiovascular reason

EVENTS SINCE DISCHARGE

Readmitted<sup>12030</sup>:  No  Yes

→ If Yes, Readmission Length of Stay<sup>12031</sup>: \_\_\_\_\_ days

→ If Yes, Readmission Date<sup>12032</sup>: mm / dd / yyyy

→ If Yes, Hospitalized at time of Follow-up<sup>12033</sup>:  No  Yes

ASD PROCEDURE

Erosion<sup>12040</sup>:  No  Yes

Device Embolization<sup>12045</sup>: (req device retrieval)  No  Yes

→ If Yes, Retrieved via Catheterization<sup>12046</sup>:  No  Yes

→ If Yes, Retrieved via Surgery<sup>12047</sup>:  No  Yes

Endocarditis<sup>12050</sup>:  No  Yes

→ If Yes, Date of Endocarditis Diagnosis<sup>12051</sup>: mm / dd / yyyy

→ If Yes, Predisposing Factors for Endocarditis<sup>12052</sup>:  Recent dental work or poor dentition  History of Endocarditis  Other implanted foreign bodies  Other surface injuries/infections  IV drug use

→ If Yes, Treatment<sup>12053</sup>:  Antibiotics  Surgical Explant  Transcatheter reintervention  Other

Residual Shunt Size<sup>12055</sup>:  None to trivial (<3 mm)  Significant (>=3 mm)



P. FOLLOW-UP (CONT.)

ELECTROPHYSIOLOGY ABLATION PROCEDURE

POST-PROCEDURE SYMPTOM SEVERITY SURVEY (SSS)

SSSQ1: In the past 6 months, what symptom or feeling that comes from the patient's heart rhythm problem bothers her/him most often?

- O Palpitations O Chest pain O Shortness of breath O Dizziness O Fatigue O Fainting O No symptoms

->If any symptoms present, SSSQ2: In the past 6 months how often has patient had this feeling?

- O Every day O At least once per week O At least once per month O At least once in the last 6 months

SSSQ3: In the past 6 months, what symptom or feeling that comes from the patient's heart rhythm problem was the worst (most severe or unpleasant)?

- O Palpitations O Chest pain O Shortness of breath O Dizziness O Fatigue O Fainting O No symptoms

SSSQ4: In the past 6 months, for any heart rhythm episodes the patient has had, what is the most intense (severe) treatment that the patient has endured to treat the rhythm problem?

- O No rhythm problems during this time O Rhythm is always present and no effort was made to try and relieve it O Self-Resolving O Vagal Maneuvers O ER visit, symptoms self-resolved or with vagal maneuvers O ER-Treated with medication O Admitted for >= 1 day, treated with medication O Hospital/ER Cardioversion

SSSQ5: In the past 6 months, has the patient taken any of the following medications? (check all that apply)

- O Amiodarone O Beta Blocker O Digoxin O Diltiazem O Dofetilide O Dronedaron O Flecainide O Mexiletine O Propafenone O Sotalol O Verapamil O None

SSSQ6: In the past 6 months, does the patient feel that their rhythm problem has interfered with how well they are able to work, go to school or play?

- O No O Yes

SSSQ7: Indicate fate of ablated substrates

- O No Recurrence O Confirmed No Recurrence O Possible Recurrence O Probable Recurrence O Confirmed Recurrence

TRANSCATHETER PULMONARY VALVE REPLACEMENT (TPVR) PROCEDURE

Transcatheter Pulmonary Valve (TPV) still in place

- O No O Yes

-> If No, Reason TPV is not still in place

- O Migration O Embolization O Explanted

TPV Reintervention

- O No O Yes

-> If Yes, TPV Surgical Reintervention

- O No O Yes

-> If Yes, TPV Surgical Reintervention Date

mm / dd / yyyy

-> If Yes, TPV Catheter Reintervention

- O No O Yes

-> If Yes, TPV Catheter Reintervention Date

mm / dd / yyyy

-> If Yes, Reason for TPV Reintervention

- O Stenosis O Pulmonary Regurgitation O Endocarditis O Other

Endocarditis

- O No O Yes

-> If Yes, Date of Endocarditis Diagnosis

mm / dd / yyyy

-> If Yes, Predisposing Factors for Endocarditis

- O Recent dental work or poor dentition O History of Endocarditis O Other implanted foreign bodies O Other surface injuries/infections O IV drug use

-> If Yes, Treatment

- O Antibiotics O Surgical Explant O Transcatheter reintervention O Other

TRANSCATHETER PULMONARY VALVE (TPV) FUNCTION

Mean gradient across valve/conduit

\_\_\_\_\_ mmHg

Maximum gradient across valve/conduit

\_\_\_\_\_ mmHg

Pulmonary Valve Regurgitation

- O None O 1+ (mild) O 2+ (moderate) O 3+ (moderately severe) O 4+ (severe)