



A. DEMOGRAPHICS

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

B. EPISODE OF CARE (ADMISSION)

Arrival Date/Time, Admitting Provider's Name, NPI, Attending Provider's Name, NPI, Health Insurance, Payment Source, HIC #, Research Study, Study Name, Patient ID, Patient Restriction

C. HISTORY AND RISK FACTORS

Hypertension, Dyslipidemia, Prior MI, Prior PCI, Tobacco Use, Cardiac Arrest Out of Healthcare Facility, Cardiac Arrest at Transferring Healthcare Facility, Height, Weight, Family Hx. of Premature CAD, Cerebrovascular Disease, Peripheral Arterial Disease, Chronic Lung Disease, Prior CABG

(KNOWN OR DIAGNOSED PRIOR TO FIRST CATH LAB VISIT)

Diabetes Mellitus, Currently on Dialysis, CSHA Clinical Frailty Scale

Canadian Study Of Health And Aging Clinical Frailty Scale Is Used With Permission For The American College Of Cardiology Foundation By Dr. Kenneth Rockwood



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

**Heart Failure**<sup>4001</sup>:  No  Yes → **If Yes, NYHA Class**<sup>4011</sup>:  Class I  Class II  Class III  Class IV

→ **If Yes, Newly Diagnosed**<sup>4012</sup>:  No  Yes

→ **If Yes, HF Type**<sup>4013</sup>:  Diastolic  Systolic  Unknown<sup>4014</sup>

(DIAGNOSTIC TEST)

**Electrocardiac Assessment Method**<sup>5037</sup>:  ECG  Telemetry Monitor  Holter Monitor  Other  None

→ **If any methods, Results**<sup>5032</sup>:  Normal  Abnormal  Uninterpretable

→ **If Abnormal, New Antiarrhythmic Therapy Initiated Prior to Cath Lab**<sup>5033</sup>:  No  Yes

→ **If Abnormal, Electrocardiac Abnormality Type**<sup>5034</sup>: (Select all that apply)

- Ventricular Fibrillation (VF)
- Sustained VT
- Non Sustained VT
- Exercise Induced VT
- T wave inversions
- ST deviation >= 0.5 mm
- New Left Bundle Branch Block
- New Onset Atrial Fib
- New Onset Atrial Flutter
- PVC – Frequent
- PVC – Infrequent
- 2<sup>nd</sup> Degree AV Heart Block Type 1
- 2<sup>nd</sup> Degree AV Heart Block Type 2
- 3<sup>rd</sup> Degree AV Heart Block
- Symptomatic Bradycardia
- Other Electrocardiac Abnormality

→ **If New Onset Atrial Fib, Heart Rate**<sup>6011</sup>: \_\_\_\_\_ bpm

→ **If Non Sustained VT, Type**<sup>5036</sup>: (Select all that apply)  Symptomatic  Newly Diagnosed  Other

**Stress Test Performed**<sup>5200</sup>:  No  Yes → **If Yes, Specify Test Performed:**

Test Type Performed <sup>5201</sup>	Most Recent Date <sup>5204</sup>	Test Results <sup>5202</sup>	→ If Positive, Risk/Extent of Ischemia <sup>5203</sup>
<input type="radio"/> Stress Echocardiogram	mm / dd / yyyy	<input type="radio"/> Negative	<input type="radio"/> Low
<input type="radio"/> Exercise Stress Test (w/o imaging)		<input type="radio"/> Positive	<input type="radio"/> Intermediate
<input type="radio"/> Stress Nuclear		<input type="radio"/> Indeterminate	<input type="radio"/> High
<input type="radio"/> Stress Imaging w/CMR		<input type="radio"/> Unavailable	<input type="radio"/> Unavailable

**Cardiac CTA Performed**<sup>5220</sup>:  No  Yes → **If Yes, Most Recent Cardiac CTA Date**<sup>5226</sup>: mm / dd / yyyy

→ **If Yes, Results**<sup>5227</sup>: (Select all that apply)  Obstructive CAD  Unclear Severity  Structural Disease

Non-Obstructive CAD  No CAD  Unknown<sup>5228</sup>

**Agatston Coronary Calcium Score Assessed**<sup>5256</sup>:  No  Yes

→ **If Yes, Agatston Coronary Calcium Score**<sup>5255</sup>: \_\_\_\_\_ → **If any value, Most Recent Calcium Score Date**<sup>5257</sup>: mm / dd / yyyy

**LVEF Assessed**<sup>5111</sup>:  No  Yes → **If Yes, Most Recent LVEF**<sup>5116</sup>: \_\_\_\_\_ %

**Prior Dx Coronary Angiography Procedure**<sup>5263</sup>: (without intervention)  No  Yes

→ **If Yes, Most Recent Procedure Date**<sup>5264</sup>: mm / dd / yyyy

→ **If Yes, Results**<sup>5265</sup>: (Select all that apply)  Obstructive CAD  Unclear Severity  Structural Disease

Non-Obstructive CAD  No CAD  Unknown<sup>5266</sup>

PRE-PROCEDURE MEDICATIONS

MEDICATION <sup>6986</sup>	ADMINISTERED <sup>6991</sup>	MEDICATION <sup>6986</sup>	ADMINISTERED <sup>6991</sup>
Aspirin	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	Statin (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated
Beta Blockers (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	Non-Statin (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated
Ca Channel Blockers (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	PCSK9 Inhibitors	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated
Antiarrhythmic Agent Other	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	ACE (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated
Long Acting Nitrates (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	ARB (Any)	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated
Ranolazine	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated	Sacubitril and Valsartan	<input type="radio"/> No <input type="radio"/> Yes <input type="radio"/> Contraindicated



D. PRE-PROCEDURE INFORMATION (CONT.)

OPTIONAL SECTION: SEATTLE ANGINA QUESTIONNAIRE (SAQ)<sup>2</sup> – FOR PARTICIPANTS CAPTURING LONG TERM CARE

OVER THE PAST 4 WEEKS, AS A RESULT OF YOUR ANGINA, HOW MUCH DIFFICULTY HAVE YOU HAD IN:

Table with 7 columns: Activity, Extremely Limited, Quite a bit Limited, Moderately Limited, Slightly Limited, Not at all Limited, Limited for other reasons or did not do these activities. Rows include walking indoors, gardening, and lifting heavy objects.

OVER THE PAST 4 WEEKS, ON AVERAGE, HOW MANY TIMES HAVE YOU...

Table with 7 columns: Frequency (4 or more times per day, 1-3 times per day, 3 or more times per week but not every day, 1-2 times per week, less than once a week, none over the past 4 weeks). Rows include chest pain and nitroglycerin use.

OVER THE PAST 4 WEEKS, HOW MUCH HAS YOUR...:

Table with 6 columns: It has extremely limited my enjoyment of life, It has limited my enjoyment of life quite a bit, It has moderately limited my enjoyment of life, It has slightly limited my enjoyment of life, It has not limited my enjoyment of life at all. Row includes chest pain limiting enjoyment of life.

IF YOU HAD TO SPEND THE REST OF YOUR LIFE WITH YOUR CHEST PAIN, CHEST TIGHTNESS OR ANGINA THE WAY IT IS RIGHT NOW...

Table with 6 columns: Not satisfied at all, Mostly dissatisfied, Somewhat satisfied, Mostly satisfied, Completely satisfied. Row includes how you would feel about this.

OPTIONAL SECTION: ROSE DYSPNEA SCALE – FOR PARTICIPANTS CAPTURING LONG TERM CARE

PLEASE THINK ABOUT HOW YOU HAVE BEEN FEELING IN THE PAST 4 WEEKS, AS YOU ANSWER THESE FOUR QUESTIONS: DO YOU GET SHORT OF BREATH WHEN...

Table with 2 columns: Question, Yes/No. Rows include hurrying on level ground, walking with other people, walking at own pace, and washing/dressing.

2 SEATTLE ANGINA QUESTIONNAIRE (© COPYRIGHT JOHN SPERTUS, MD, MPH) IS USED WITH PERMISSION FOR NCDR BY WWW.CVOUTCOMES.ORG



E. PROCEDURE INFORMATION

**Procedure Start Date/Time**<sup>7000</sup>: mm/dd/yyyy / hh:mm      **Procedure End Date/Time**<sup>7005</sup>: mm/dd/yyyy / hh:mm

**Diagnostic Coronary Angiography Procedure**<sup>7045</sup>:  No  Yes  
 → If Yes, **Diagnostic Cath Operator's Name, NPI**<sup>7046,7047,7048,7049</sup>: \_\_\_\_\_

**Percutaneous Coronary Intervention (PCI)**<sup>7050</sup>:  No  Yes  
 → If Yes, **PCI Operator's Name, NPI**<sup>7051,7052,7053,7054</sup>: \_\_\_\_\_

**Diagnostic Left Heart Cath**<sup>7060</sup>:  No  Yes      → If Yes, **LVEF**<sup>7061</sup>: \_\_\_\_\_ %

**Concomitant Procedures Performed**<sup>7065</sup>:  No  Yes  
 → If Yes, **Procedure Type(s)**<sup>7066</sup>: (Select the best option(s)) \_\_\_\_\_, \_\_\_\_\_, \_\_\_\_\_

**Arterial Access Site**<sup>7320</sup>:  Femoral     Brachial     Radial     Other

**Arterial Cross Over**<sup>7325</sup>:  No  Yes

<b>Closure Method(s)</b> <sup>7330,7331,7333</sup> :	1	Reserved for future use	<input type="checkbox"/> <b>Method Not Documented</b> <sup>7332</sup>
	2	Reserved for future use	
	3	Reserved for future use	

**Venous Access**<sup>7335</sup>: (concomitant entry for Cath procedure)  No  Yes

**Systolic BP**<sup>6016</sup>: \_\_\_\_\_ mmHg

**Cardiac Arrest at this facility**<sup>7340</sup>:  No  Yes

RADIATION EXPOSURE AND CONTRAST

**CODE ALL AVAILABLE MEASUREMENTS** → **Fluoro Time**<sup>7214</sup>: \_\_\_\_\_ minutes      **Contrast Volume**<sup>7215</sup>: \_\_\_\_\_ mL

**Cumulative Air Kerma**<sup>7210</sup>: \_\_\_\_\_  mGy  Gy

**Dose Area Product**<sup>7220</sup>: \_\_\_\_\_  Gy/cm<sup>2</sup>  dGy/cm<sup>2</sup>  cGy/cm<sup>2</sup>  mGy/cm<sup>2</sup>  μGy/M<sup>2</sup>

F. LABS

PRE-PROCEDURE (VALUES CLOSEST TO THE PROCEDURE)		POST-PROCEDURE	
<b>Troponin I</b> <sup>6090</sup> : _____ ng/mL <input type="checkbox"/> Not Drawn <sup>6091</sup>		<b>Troponin I</b> <sup>8515</sup> : _____ ng/mL <input type="checkbox"/> Not Drawn <sup>8516</sup>	
<b>Troponin T</b> <sup>6095</sup> : _____ ng/mL <input type="checkbox"/> Not Drawn <sup>6096</sup>		<b>Troponin T</b> <sup>8520</sup> : _____ ng/mL <input type="checkbox"/> Not Drawn <sup>8521</sup>	
<b>Creatinine</b> <sup>6050</sup> : _____ mg/dL <input type="checkbox"/> Not Drawn <sup>6051</sup>		<b>Creatinine</b> <sup>8510</sup> : (peak) _____ mg/dL <input type="checkbox"/> Not Drawn <sup>8511</sup>	
<b>Hemoglobin</b> <sup>6030</sup> : _____ g/dL <input type="checkbox"/> Not Drawn <sup>6031</sup>		<b>Hemoglobin</b> <sup>8505</sup> : (Lowest w/in 72 hours) _____ g/dL <input type="checkbox"/> Not Drawn <sup>8506</sup>	
<b>Total Cholesterol</b> <sup>6100</sup> : _____ mg/dL <input type="checkbox"/> Not Drawn <sup>6101</sup>			
<b>HDL</b> <sup>6105</sup> : _____ mg/dL <input type="checkbox"/> Not Drawn <sup>6106</sup>			



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

Indication(s) for Cath Lab Visit<sup>7400</sup>: (Select all that apply)

- ACS <= 24 hrs, ACS > 24 hrs, New Onset Angina <= 2 months, Worsening Angina, Resuscitated Cardiac Arrest, Stable Known CAD, Suspected CAD, Valvular Disease, Pericardial Disease, Cardiac Arrhythmia, Cardiomyopathy, LV Dysfunction, Syncope, Post Cardiac Transplant, Pre-operative evaluation, Evaluation for Exercise Clearance, Other

Chest Pain Symptom Assessment<sup>7405</sup>: O Typical Angina O Atypical Angina O Non-anginal Chest Pain O Asymptomatic

Cardiovascular Instability<sup>7410</sup>: O No O Yes

-> If Yes, Cardiovascular Instability Type<sup>7415</sup>: (Select all that apply)

- Persistent Ischemic Symptoms (chest pain, STE), Hemodynamic Instability (not cardiogenic shock), Ventricular Arrhythmias, Cardiogenic Shock, Acute Heart Failure Symptoms, Refractory Cardiogenic Shock

Ventricular Support<sup>7420</sup>: O No O Yes

-> If Yes, Pharmacologic Vasopressor Support<sup>7421</sup>: O No O Yes

-> If Yes, Mechanical Support<sup>7422</sup>: O No O Yes

-> If Yes, Device<sup>7423</sup>: \_\_\_\_\_

-> If Yes, Timing<sup>7424</sup>: O In place at start of procedure O Inserted during procedure and prior to intervention O Inserted after intervention has begun

-> IF INDICATION(S) FOR CATH LAB VISIT<sup>7400</sup> = 'VALVULAR DISEASE' (COMPLETE FOR EACH TYPE)

Table with 2 main sections: VALVULAR DISEASE STENOSIS TYPE and VALVULAR DISEASE REGURGITATION TYPE. Each section has a 2x3 grid of options for stenosis/regurgitation types and severity levels.

-> IF INDICATION(S) FOR CATH LAB VISIT<sup>7400</sup> = 'PRE-OPERATIVE EVALUATION'

Evaluation for Surgery Type<sup>7465</sup>: O Cardiac Surgery O Non-Cardiac Surgery

Functional Capacity<sup>7466</sup>: O < 4 METS O >= 4 METS without symptoms O >= 4 METS with symptoms O Unknown<sup>7467</sup>

Surgical Risk<sup>7468</sup>: O Low O Intermediate O High Risk: Vascular O High Risk: Non-Vascular

Solid Organ Transplant Surgery<sup>7469</sup>: O No O Yes

-> If Yes, Donor<sup>7470</sup>: O No O Yes

-> If Yes, Organ<sup>7471</sup>: (Select all that apply) O Heart O Kidney O Liver O Lung O Pancreas O Other Organ



H. CORONARY ANATOMY

Dominance<sup>7500</sup>:  Left  Right  Co-dominant

Native Vessel with Stenosis >= 50%<sup>7505</sup>:  No  Yes → If Yes, Specify Segment(s):

Table with 2 columns: SEGMENT NUMBER<sup>7507</sup> and MEASUREMENT (FOR EACH SELECTED). Rows include Native Stenosis<sup>7508</sup> and Adjunctive Measurements Obtained<sup>7511</sup> with sub-questions for FFR Ratio<sup>7512</sup>, iFR Ratio<sup>7513</sup>, IVUS MLA<sup>7514</sup>, and OCT MLA<sup>7515</sup>.

Graft Vessel with Stenosis >= 50%<sup>7525</sup>: (Note 1)  No  Yes → If Yes, Specify Segment(s):

Table with 2 columns: SEGMENT NUMBER<sup>7527</sup> and MEASUREMENT (FOR EACH SELECTED). Rows include Graft Stenosis<sup>7528</sup>, Graft Vessel<sup>7529</sup> (LIMA, RIMA, SVG, Radial, Unknown<sup>7530</sup>), and Adjunctive Measurements Obtained<sup>7531</sup> with sub-questions for FFR Ratio<sup>7532</sup>, iFR Ratio<sup>7533</sup>, IVUS MLA<sup>7534</sup>, and OCT MLA<sup>7535</sup>.

NOTE 1: CABG DATE/TIME<sup>10011</sup> MUST BE LESS THAN PROCEDURE START DATE/TIME<sup>7000</sup> OR PRIOR CABG<sup>4515</sup> = 'YES' TO COMPLETE THESE ELEMENTS.

I. PCI PROCEDURE (COMPLETE FOR EACH CATH LAB VISIT IN WHICH A PCI WAS ATTEMPTED OR PERFORMED)

PCI Status<sup>7800</sup>:  Elective  Urgent  Emergency  Salvage

CARDIAC ARREST OUT OF HEALTHCARE FACILITY<sup>4630</sup> = 'YES' OR CARDIAC ARREST AT TRANSFERRING HEALTHCARE FACILITY<sup>4635</sup> = 'YES' OR CARDIAC ARREST AT THIS FACILITY<sup>7340</sup> = 'YES'. Includes Hypothermia Induced<sup>7806</sup> and Level of Consciousness<sup>7810</sup> questions.

Decision for PCI with Surgical Consult<sup>7815</sup>:  No  Yes. → If Yes, CV Treatment Decision<sup>7816</sup>:  Surgery Not Recommended,  Surgery Recommended, Patient/Family Declined,  Surgery Recommended, Patient/Family Accepted (Hybrid procedure)

PCI for Multi-vessel Disease<sup>7820</sup>:  No  Yes. → If Yes, Multi-vessel Procedure Type<sup>7821</sup>: (in this lab visit)  Initial PCI  Staged PCI

PCI Indication<sup>7825</sup>:  STEMI - Immediate PCI for Acute STEMI,  STEMI - Rescue (after unsuccessful lytics),  STEMI - Stable (<= 12 hrs from Sx),  STEMI - Stable (> 12 hrs from Sx),  STEMI - Unstable (> 12 hrs from Sx),  STEMI (after successful lytics),  NSTE - ACS,  Stable Angina,  CAD (without Ischemic Sx),  Other. Includes Symptom Date/Time<sup>7826,7827</sup>, Thrombolytics<sup>7829</sup>, and Syntax Score<sup>7831</sup> questions.



I. PCI PROCEDURE (COMPLETE FOR EACH CATH LAB VISIT IN WHICH A PCI WAS ATTEMPTED OR PERFORMED) (CONT.)

<p>→ IF PCI INDICATION<sup>7825</sup> = 'STEMI – IMMEDIATE PCI FOR ACUTE STEMI'</p>	<p><b>STEMI or STEMI Equivalent First Noted</b><sup>7835</sup>:</p>	<p><input type="radio"/> First ECG    <input type="radio"/> Subsequent ECG</p>
	<p>→ If Subsequent ECG, <b>ECG with STEMI/ STEMI Equivalent Date &amp; Time</b><sup>7836</sup>:</p>	<p>mm/dd/yyyy / hh:mm</p>
	<p>→ If Subsequent ECG, <b>ECG obtained in Emergency Department</b><sup>7840</sup>:</p>	<p><input type="radio"/> No    <input type="radio"/> Yes</p>
	<p><b>Transferred In For Immediate PCI for STEMI</b><sup>7841</sup>:</p>	<p><input type="radio"/> No    <input type="radio"/> Yes</p>
	<p>→ If Yes, <b>Date &amp; Time ED Presentation at Referring Facility</b><sup>7842</sup>:</p>	<p>mm/dd/yyyy / hh:mm</p>
	<p><b>First Device Activation Date &amp; Time</b><sup>7845</sup>:</p>	<p>mm/dd/yyyy / hh:mm</p>
<p><b>Patient Centered Reason for Delay in PCI</b><sup>7850</sup>:</p>		<p><input type="radio"/> No    <input type="radio"/> Yes</p>
<p>→ If Yes, <b>Reason</b><sup>7851</sup>:</p>		<p> <input type="radio"/> Difficult Vascular Access                      <input type="radio"/> Patient delays in providing consent for PCI  <input type="radio"/> Difficulty crossing the culprit lesion            <input type="radio"/> Emergent placement of LV support device before PCI  <input type="radio"/> Cardiac arrest and/or need for intubation before PCI    <input type="radio"/> Other </p>

PCI PROCEDURE MEDICATIONS (ADMINISTERED WITHIN 24 HOURS PRIOR TO AND DURING THE PCI PROCEDURE)

MEDICATION <sup>7990</sup>		ADMINISTERED <sup>7995</sup>	MEDICATION <sup>7990</sup>		ADMINISTERED <sup>7995</sup>
ANTICOAGULANT	Argatroban	<input type="radio"/> No <input type="radio"/> Yes	GLYCOPROTEIN (GP) IIb/IIIa INHIBITORS	GP IIb/IIIa Inhibitors (Any)	<input type="radio"/> No <input type="radio"/> Yes
	Bivalirudin	<input type="radio"/> No <input type="radio"/> Yes			
	Fondaparinux	<input type="radio"/> No <input type="radio"/> Yes	NON-VITAMIN K DEPENDENT ORAL ANTICOAGULANT	Apixaban	<input type="radio"/> No <input type="radio"/> Yes
	Low Molecular Wt Heparin	<input type="radio"/> No <input type="radio"/> Yes		Dabigatran	<input type="radio"/> No <input type="radio"/> Yes
	Unfractionated Heparin	<input type="radio"/> No <input type="radio"/> Yes		Edoxaban	<input type="radio"/> No <input type="radio"/> Yes
	Warfarin	<input type="radio"/> No <input type="radio"/> Yes		Rivaroxaban	<input type="radio"/> No <input type="radio"/> Yes
ANTIPLATELET	Vorapaxar	<input type="radio"/> No <input type="radio"/> Yes	P2Y12 INHIBITORS	Cangrelor	<input type="radio"/> No <input type="radio"/> Yes
				Clopidogrel	<input type="radio"/> No <input type="radio"/> Yes
				Prasugrel	<input type="radio"/> No <input type="radio"/> Yes
				Ticagrelor	<input type="radio"/> No <input type="radio"/> Yes



J. LESIONS AND DEVICES (COMPLETE FOR EACH PCI ATTEMPTED OR PERFORMED)

<b>Lesion Counter</b> <sup>8000</sup> :	<b>1</b>		<b>2</b>		
<b>Segment Number(s)</b> <sup>8001</sup> :	____, _____, _____, _____, _____		____, _____, _____, _____, _____		
<b>If PCI Indication</b> <sup>7825</sup> is STEMI or NSTEMI-ACS, <b>Culprit Stenosis</b> <sup>8002</sup> :	<input type="radio"/> No <input type="radio"/> Yes <input type="checkbox"/> Unknown <sup>8003</sup>		<input type="radio"/> No <input type="radio"/> Yes <input type="checkbox"/> Unknown <sup>8003</sup>		
<b>Stenosis Immediately Prior to Rx</b> <sup>8004</sup> :	_____%		_____%		
→ <b>If 100%, Chronic Total Occlusion</b> <sup>8005</sup> :	<input type="radio"/> No <input type="radio"/> Yes <input type="checkbox"/> Unknown <sup>8006</sup>		<input type="radio"/> No <input type="radio"/> Yes <input type="checkbox"/> Unknown <sup>8006</sup>		
<b>TIMI Flow (Pre-Intervention)</b> <sup>8007</sup> :	<input type="radio"/> TIMI-0 <input type="radio"/> TIMI-1 <input type="radio"/> TIMI-2 <input type="radio"/> TIMI-3		<input type="radio"/> TIMI-0 <input type="radio"/> TIMI-1 <input type="radio"/> TIMI-2 <input type="radio"/> TIMI-3		
<b>Previously Treated Lesion</b> <sup>8008</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, Date</b> <sup>8009</sup> :	mm / dd / yyyy		mm / dd / yyyy		
→ <b>If Yes, Treated with Stent</b> <sup>8010</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, In-Stent Restenosis</b> <sup>8011</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, In-Stent Thrombosis</b> <sup>8012</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, Stent Type</b> <sup>8013</sup> :	<input type="radio"/> DES <input type="radio"/> BMS <input type="checkbox"/> Unknown <sup>8014</sup> <input type="radio"/> Bioabsorbable		<input type="radio"/> DES <input type="radio"/> BMS <input type="checkbox"/> Unknown <sup>8014</sup> <input type="radio"/> Bioabsorbable		
<b>Lesion in Graft</b> <sup>8015</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, Type of CABG Graft</b> <sup>8016</sup> :	<input type="radio"/> LIMA <input type="radio"/> Vein <input type="radio"/> Other Artery		<input type="radio"/> LIMA <input type="radio"/> Vein <input type="radio"/> Other Artery		
→ <b>If Yes, Location in Graft</b> <sup>8017</sup> :	<input type="radio"/> Aortic <input type="radio"/> Body <input type="radio"/> Distal		<input type="radio"/> Aortic <input type="radio"/> Body <input type="radio"/> Distal		
<b>Navigate through Graft to Native Lesion</b> <sup>8018</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
<b>Lesion Complexity</b> <sup>8019</sup> :	<input type="radio"/> Non-High/Non-C <input type="radio"/> High/C		<input type="radio"/> Non-High/Non-C <input type="radio"/> High/C		
<b>Lesion Length</b> <sup>8020</sup> :	____mm		____mm		
<b>Severe Calcification</b> <sup>8021</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
<b>Bifurcation Lesion</b> <sup>8022</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
<b>Guidewire Across Lesion</b> <sup>8023</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, Device(s) Deployed</b> <sup>8024</sup> :	<input type="radio"/> No <input type="radio"/> Yes		<input type="radio"/> No <input type="radio"/> Yes		
→ <b>If Yes, Stenosis (Post-Intervention)</b> <sup>8025</sup> :	_____%		_____%		
→ <b>If Yes, TIMI Flow (Post-Intervention)</b> <sup>8026</sup> :	<input type="radio"/> TIMI-0 <input type="radio"/> TIMI-1 <input type="radio"/> TIMI-2 <input type="radio"/> TIMI-3		<input type="radio"/> TIMI-0 <input type="radio"/> TIMI-1 <input type="radio"/> TIMI-2 <input type="radio"/> TIMI-3		
<b>Intracoronary Device(s) Used</b> <sup>8027,8028</sup>	<b>Unique Device Identifier (UDI)</b> <sup>8029</sup>	<b>Associated Lesion(s)</b> <sup>8030</sup>	<b>Diameter</b> <sup>8031</sup>	<b>Length</b> <sup>8032</sup>	
1	Reserved for future use	____, _____, _____	____ mm	____ mm	
2	Reserved for future use	____, _____, _____	____ mm	____ mm	



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

INTRA PCI ONLY	PERCUTANEOUS CORONARY INTERVENTION (PCI) <sup>7050</sup> = 'YES'	Coronary Artery Perforation <sup>9145</sup> :	<input type="radio"/> No	<input type="radio"/> Yes
		Significant Coronary Artery Dissection <sup>9146</sup> :	<input type="radio"/> No	<input type="radio"/> Yes

INTRA AND POST-PROCEDURE EVENTS (NOTE 1: RECORD EACH EVENT SEPARATELY INDICATING THE DATE AND TIME)

EVENT(S) <sup>9001</sup>	EVENT(S) OCCURRED <sup>9002</sup>	→ IF YES, EVENT DATE/TIME(S) <sup>9003</sup>
Bleeding – Access Site	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Bleeding – Gastrointestinal	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Bleeding – Genitourinary	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Bleeding – Hematoma at Access Site	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Bleeding – Other	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Bleeding – Retroperitoneal	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Cardiac Arrest	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Cardiac Tamponade	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Cardiogenic Shock	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Heart Failure	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Myocardial Infarction	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
New Requirement for Dialysis	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Other Vascular Complications Req Tx	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Stroke – Hemorrhagic	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Stroke – Ischemic	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm
Stroke – Undetermined	<input type="radio"/> No <input type="radio"/> Yes	mm/dd/yyyy / hh:mm

RBC Transfusion<sup>9275</sup>:  No  Yes

→ If Yes, Number of Units Transfused<sup>9276</sup>: \_\_\_\_\_

→ If Yes, Transfusion PCI<sup>9277</sup>: (within 72 hours)  No  Yes

→ If Yes, Transfusion Surgical<sup>9278</sup>: (within 72 hours)  No  Yes



L. DISCHARGE

Intervention(s) this Hospitalization<sup>10030</sup>: (not during same lab visit as Cath or PCI)  No  Yes

→ If Yes, Type<sup>10031</sup>: (Select all that apply)  CABG  Cardiac Surgery (non CABG)  Surgery (non Cardiac)  Valvular Intervention  Structural Heart Intervention (non-valvular)  EP Study  Other

→ IF CABG = 'YES' CABG Status<sup>10035</sup>:  Elective  Urgent  Emergency  Salvage CABG Indication<sup>10036</sup>:  PCI/CABG Hybrid Procedure  Recommendation from Dx Cath (instead of PCI)  PCI Failure  PCI Complication CABG Date/Time<sup>10011</sup>: mm/dd/yyyy / hh:mm

Creatinine<sup>10060</sup>: (at D/C) \_\_\_\_\_ mg/dL  Not Drawn<sup>10061</sup> Hemoglobin<sup>10065</sup>: (at D/C) \_\_\_\_\_ g/dL  Not Drawn<sup>10066</sup>

Discharge Date/Time<sup>10101</sup>: mm/dd/yyyy / hh:mm Discharge Provider's Name, NPI<sup>10070,10071,10072,10073</sup>: \_\_\_\_\_

Comfort Measures Only<sup>10075</sup>:  No  Yes

Discharge Status<sup>10105</sup>:  Alive  Deceased

→ If Alive, Discharge Location<sup>10110</sup>:  Home  Skilled Nursing facility  Extended care/TCU/rehab  Other  Other acute care hospital  Left against medical advice (AMA)

→ If Other acute care hospital, Transferred for CABG<sup>10111</sup>:  No  Yes

→ If Not Left against medical advice (AMA) OR Other acute care hospital, CABG Planned after Discharge<sup>10112</sup>:  No  Yes

→ If Alive, Hospice Care<sup>10115</sup>:  No  Yes

→ If Alive, Cardiac Rehabilitation Referral<sup>10116</sup>:  No – Reason Not Documented  No – Health Care System Reason Documented  No – Medical Reason Documented  Yes

→ If Deceased AND any (CARDIAC ARREST OUT OF HEALTHCARE FACILITY<sup>4630</sup> = 'YES' OR CARDIAC ARREST AT TRANSFERRING HEALTHCARE FACILITY<sup>4635</sup> = 'YES' OR CARDIAC ARREST AT THIS FACILITY<sup>7340</sup> = 'YES'), Level of Consciousness<sup>10117</sup>: (highest s/p cardiac arrest)

(A) Alert  (V) Verbal  (P) Pain  (U) Unresponsive  Unable to assess

→ If Deceased, Death During the Procedure<sup>10120</sup>:  No  Yes

→ If Deceased, Cause of Death<sup>10125</sup>:

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

DISCHARGE MEDICATIONS (PRESCRIBED AT DISCHARGE - COMPLETE FOR EACH EPISODE OF CARE IN WHICH A PCI WAS ATTEMPTED OR PERFORMED)

Medications prescribed at discharge are not required for patients who expired, discharged to "Other acute care Hospital", "AMA", or are receiving Hospice Care.

MEDICATION <sup>10200</sup>	PRESCRIBED <sup>10205</sup>				→ IF YES, DOSE <sup>10207</sup>			→ IF NO - PT. REASON, PATIENT RATIONALE <sup>10206</sup> (Select all that apply)			
	YES	No - NO REASON	No - MEDICAL REASON	No - PT. REASON	LOW	MODERATE	HIGH				
ACE INHIBITORS (ANGIOTENSIN CONVERTING ENZYME)	ACE Inhibitors (Any)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="checkbox"/> Cost <input type="checkbox"/> Alternative Therapy Preferred <input type="checkbox"/> Negative Side Effect					
ANTICOAGULANT	Warfarin	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>						
ANTIPLATELET	Aspirin	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>						
	Vorapaxar	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>						
									<input type="checkbox"/> Cost <input type="checkbox"/> Alternative Therapy Preferred <input type="checkbox"/> Negative Side Effect		
									<input type="checkbox"/> Cost <input type="checkbox"/> Alternative Therapy Preferred <input type="checkbox"/> Negative Side Effect		



L. DISCHARGE (CONT.)

DISCHARGE MEDICATIONS (PRESCRIBED AT DISCHARGE - COMPLETE FOR EACH EPISODE OF CARE IN WHICH A PCI WAS ATTEMPTED OR PERFORMED)

Medications prescribed at discharge are not required for patients who expired, discharged to "Other acute care Hospital", "AMA", or are receiving Hospice Care.

Table with columns: MEDICATION, PRESCRIBED (Yes, No - No Reason, No - Medical Reason, No - Pt. Reason), IF YES, DOSE (Low, Moderate, High), and IF NO - PT. REASON, PATIENT RATIONALE. Rows include ARB, Beta Blockers, Non-Vitamin K Dependent Oral Anticoagulant, P2Y12 Inhibitors, Statin, Non-Statins, and PCSK9 Inhibitors.

Discharge Medication Reconciliation Completed: O No O Yes

If Yes, Reconciled Medications: (Select all that apply)

- Prescriptions: Cardiac, Over the Counter (OTC) Medications, Vitamins/Minerals, Prescriptions: Non-Cardiac, Herbal Supplements



M. FOLLOW-UP: 30 DAY (23 TO 75 DAYS POST INDEX PCI PROCEDURE): 1 YEAR (305 TO 425 DAYS POST INDEX PCI PROCEDURE)

Assessment Date<sup>11000</sup>: mm / dd / yyyy Reference Episode Arrival Date/Time<sup>11002</sup>: mm/dd/yyyy / hh:mm

Reference Procedure Start Date/Time<sup>11001</sup>: mm/dd/yyyy / hh:mm Reference Episode Discharge Date/Time<sup>11015</sup>: mm/dd/yyyy / hh:mm

Method(s) to Determine Status<sup>11003</sup>: (Select all that apply)
Office Visit, Medical Records, Letter from Medical Provider, Phone Call, Social Security Death Master File, Hospitalized, Other

Follow-Up Status<sup>11004</sup>: O Alive O Deceased O Lost to Follow-up
If Alive, Chest Pain Symptom Assessment<sup>11005</sup>: O Typical Angina O Atypical Angina O Non-anginal Chest Pain O Asymptomatic

If Deceased, Date of Death<sup>11006</sup>: mm / dd / yyyy

If Deceased, Primary Cause of Death<sup>11007</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

Research Study<sup>11008</sup>: O No O Yes If Yes, Study Name<sup>11009</sup>, Patient ID<sup>11010</sup>: \_\_\_\_\_

EVENTS, INTERVENTIONS AND/OR SURGICAL PROCEDURES (ANY OCCURRENCE BETWEEN DISCHARGE (OR PREVIOUS FOLLOW-UP) AND THE CURRENT FOLLOW-UP ASSESSMENT) (NOTE 1: RECORD EACH EVENT SEPARATELY INDICATING THE DATE)

Table with 4 columns: EVENT(S)<sup>11011</sup>, EVENT(S) OCCURRED<sup>11012</sup>, IF YES, DEVICE(S) EVENT OCCURRED IN<sup>11013</sup>, IF YES, EVENT DATE(S)<sup>11014</sup>. Rows include Bleeding Event, CABG, Myocardial Infarction, PCI, Readmission, Stroke, and Thrombosis.



M. FOLLOW-UP (CONT.)

FOLLOW-UP MEDICATIONS		PRESCRIBED <sup>11995</sup>				→ IF YES, DOSE <sup>11996</sup>		
MEDICATION <sup>11990</sup>		YES	NO - NO REASON	NO - MEDICAL REASON	NO - PT. REASON	LOW	MODERATE	HIGH
		<b>ACE INHIBITORS (ANGIOTENSIN CONVERTING ENZYME)</b>	ACE Inhibitors (Any)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>ANTICOAGULANT</b>	Warfarin	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>ANTIPLATELET</b>	Aspirin	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Vorapaxar	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>ARB (ANGIOTENSIN RECEPTORS BLOCKERS)</b>	ARB (Any)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>NON-VITAMIN K DEPENDENT ORAL ANTICOAGULANT</b>	Apixaban	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Dabigatran	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Edoxaban	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Rivaroxaban	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>P2Y12 INHIBITORS</b>	Clopidogrel	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Prasugrel	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Ticagrelor	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Ticlopidine	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>STATIN</b>	Statin (Any)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	
<b>NON-STATIN</b>	Non-Statins (Any)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
<b>PCSK9 INHIBITORS</b>	Alirocumab	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			
	Evolocumab	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>			



M. FOLLOW-UP (CONT.)

OPTIONAL SECTION: SEATTLE ANGINA QUESTIONNAIRE (SAQ)<sup>2</sup> – FOR PARTICIPANTS CAPTURING LONG TERM CARE

OVER THE PAST 4 WEEKS, AS A RESULT OF YOUR ANGINA, HOW MUCH DIFFICULTY HAVE YOU HAD IN:

Table with 7 columns: Activity, Extremely Limited, Quite a bit limited, Moderately limited, Slightly limited, Not at all limited, Limited for other reasons or did not do these activities. Rows include walking indoors, gardening, and lifting heavy objects.

OVER THE PAST 4 WEEKS, ON AVERAGE, HOW MANY TIMES HAVE YOU...

Table with 7 columns: Frequency (4 or more times per day, 1-3 times per day, 3 or more times per week but not every day, 1-2 times per week, less than once a week, none over the past 4 weeks). Rows include chest pain and nitroglycerin use.

OVER THE PAST 4 WEEKS, HOW MUCH HAS YOUR...:

Table with 6 columns: It has extremely limited my enjoyment of life, It has limited my enjoyment of life quite a bit, It has moderately limited my enjoyment of life, It has slightly limited my enjoyment of life, It has not limited my enjoyment of life at all. Row includes chest pain limiting enjoyment of life.

IF YOU HAD TO SPEND THE REST OF YOUR LIFE WITH YOUR CHEST PAIN, CHEST TIGHTNESS OR ANGINA THE WAY IT IS RIGHT NOW...

Table with 6 columns: Not satisfied at all, Mostly dissatisfied, Somewhat satisfied, Mostly satisfied, Completely satisfied. Row includes how you would feel about this.

OPTIONAL SECTION: ROSE DYSPNEA SCALE – FOR PARTICIPANTS CAPTURING LONG TERM CARE

PLEASE THINK ABOUT HOW YOU HAVE BEEN FEELING IN THE PAST 4 WEEKS, AS YOU ANSWER THESE FOUR QUESTIONS: DO YOU GET SHORT OF BREATH WHEN...

Table with 4 rows of questions about shortness of breath during activities like hurrying, walking with others, walking at own pace, and washing/dressing, with Yes/No options.

<sup>2</sup>SEATTLE ANGINA QUESTIONNAIRE (© COPYRIGHT JOHN SPERTUS, MD, MPH) IS USED WITH PERMISSION FOR NCDR BY WWW.CVOUTCOMES.ORG