

Survey Monkey Results

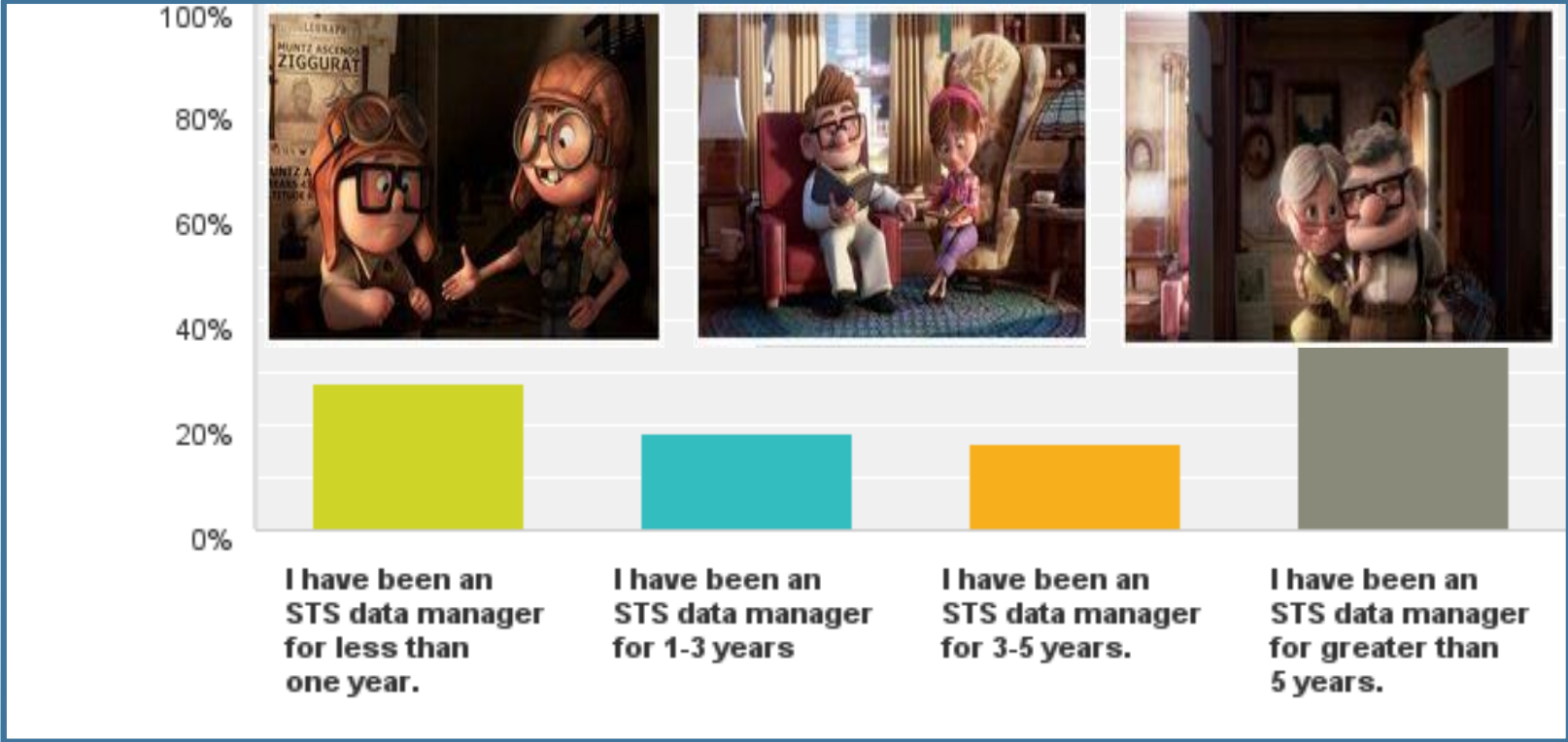
MICHIGAN STS DATA MANAGERS MEETING

November 18, 2016
St. Mary's Hospital
Saginaw, MI

Thank You to those that responded!!

- Once again, rather than multiple scenarios, all questions are based on a single scenario to mimic chart abstraction.
- This scenario was designed from the cases that you submitted for clarification. The questions were designed for the benefit of our less experienced managers.
- Please provide feedback regarding this format and future topics in your evaluations!

Experience Level



Answer Choices	Responses
I have been an STS data manager for less than one year .	27.91% 12
I have been an STS data manager for 1-3 years	18.60% 8
I have been an STS data manager for 3-5 years.	16.28% 7
I have been an STS data manager for greater than 5 years.	37.21% 16
Total	43

Patient Scenario

An 82 year old female presents to the ER on Tuesday evening, complaining of severe chest pain and SOB after watching the election results. ECG and Lab results lead to a diagnosis of NSTEMI, and the patient is taken to the Cath Lab for intervention. The cath shows 75% Left Main disease and 80% stenosis in the mid-RCA. Attempts to stent the RCA were unsuccessful, and the patient was transferred to a "Sister" hospital for further treatment. After arrival, a TEE demonstrates an EF of "30% w/ global hypokinesis", and mild mitral insufficiency. Surgery is consulted, and CABG surgery is scheduled for Friday, pending treatment and improvement of orthopnea, O2 requirements, and pulmonary edema.

After induction, a TEE examination now reveals "moderate to severe" mitral insufficiency with an EF of 25%. The surgeon then changes the operative plan to include inspection and possible repair of the mitral valve. CABG x 3 is performed, and inspection of the mitral valve reveals significant prolapse of the posterior leaflet, requiring triangular resection and annuloplasty. Given the patient's age, the left atrial appendage is oversewn from within the left atrium. After several failed attempts, the patient is eventually weaned from CPB after insertion of an IABP. The post repair TEE now shows mild MR.

Postoperatively, the patient has the IABP removed in the afternoon of POD #1, with extubation shortly thereafter. Continued complaints of SOB and DOE lead to the discovery of a "moderate" pleural effusion on POD #2, requiring a pleurocentesis. That evening, the patient develops A-Fib resulting in recurrent dyspnea and hypotension. TEE reveals only mild mitral insufficiency, but a "significant" pericardial effusion. Further hypotension and instability results in a return trip to the OR for relief of tamponade on POD #3. The patient remained intubated until POD #5 due to "wet lungs" and recurrent A-Fib requiring Amiodarone and aggressive diuresis. Neurology was consulted for post-extubation confusion, while Nephrology monitored a transient rise in creatinine. After a prolonged recovery period, the patient is discharged to an ECF on POD #20.

On the morning of POD #27, the ECF phones to report that the patient has been found unresponsive, and is in route to the hospital while undergoing CPR. Upon arrival, the patient is found to be in refractory V-Fib. Unresponsive to all resuscitative efforts, the patient expires within 10 minutes of arrival.

Question 2 Selections:

- Previous PCI = Yes; Within this episode of Care = Yes, at this facility;
Indication for Surgery = Failure w/o Clinical Deterioration.
- Previous PCI = Yes; Within this episode of Care = Yes, at some other facility;
Indication for Surgery = Failure w/o Clinical Deterioration.
- Previous PCI = Yes; Within this episode of Care = No.
- Previous PCI = Yes; Within this episode of Care = Yes, at some other facility;
Indication for Surgery = PCI/Surgery Staged (not STEMI).

Data Points to Consider

- Did the patient undergo Previous Cardiac Intervention?
- Within this Episode of Care?
- Where did the intervention take place?
- What was the Indication for Surgery?

E. Previous Cardiac Interventions

Previous Cardiac Interventions: **PrCVInt (665)**  Yes No Unknown

(If Yes →) Previous coronary artery bypass (CAB): **PrCAB (670)** Yes No

Previous valve procedure: **PrValve (675)** Yes No If PrValve Yes, Enter at least one previous valve procedure and up to 5 ↓

Seq. #: 665

Long Name: Prev Cardiac Intervent; **Short Name:** PrCVInt

Definition: Indicate whether the patient has undergone *any previous cardiovascular intervention, either surgical or non-surgical, which may include those done during the current admission.*

Previous PCI: **POCPCI (775)**  Yes No

(If Yes →) PCI Performed Within This Episode Of Care: Yes, at this facility Yes, at some other acute care facility No

POCPCIWhen (780)

Indication for Surgery:

POCPCIndSurg (785)

PCI Complication

PCI Failure with Clinical Deterioration

PCI for STEMI, multivessel disease

PCI Failure without Clinical Deterioration

PCI/Surgery Staged (not STEMI)

Other

PCI Stent: Yes No

POCPCISt (790)

PCI Interval:

POCPCIIn (800)

(If Yes →) Stent Type: Bare metal Drug-eluting Bioresorbable Multiple Unknown




POCPCIStTy (795)

≤ 6 Hours > 6 Hours

Seq. #: 775

Long Name: Previous PCI; **Short Name:** POCPCI

Definition: Indicate whether a previous Percutaneous Coronary Intervention (PCI) was *performed any time prior to this surgical procedure.*

Previous PCI: **POCPCI (775)**  Yes No
 (If Yes →) PCI Performed Within This Episode Of Care: Yes, at this facility  Yes, at some other acute care facility No
POCPCIWhen (780)
 Indication for Surgery: PCI Complication  PCI Failure without Clinical Deterioration
POCPCIndSurg (785) PCI Failure with Clinical Deterioration PCI/Surgery Staged (not STEMI)
 PCI for STEMI, multivessel disease Other
 PCI Stent: Yes No (If Yes →) Stent Type: Bare metal Drug-eluting Bioresorbable Multiple Unknown
POCPCISt (790) **POCPCISTy (795)**
 PCI Interval: ≤ 6 Hours > 6 Hours
POCPCIIn (800)

Seq. #: 780
Long Name: Previous PCI-Within This Episode of Care; **Short Name:** POCPCIWhen
Definition: Indicate whether the previous Percutaneous Cardiac Intervention (PCI) was performed within this episode of care. *Episode of care is defined as continuous inpatient hospitalization which includes transfer from one acute care hospital to another.*
Intent/Clarification:
 This field is intended to capture PCIs done during the same episode of care prior to the surgical procedure.
Include patients who were transferred for surgery from another facility following PCI.

NOTE THAT SEQUENCE NUMBER 785 IS A CHILD TO SEQUENCE NUMBER 780.

at least one previous other cardiac procedure and up to 7 ↓)

Intent/Clarification: Indicate whether surgery was required due to:

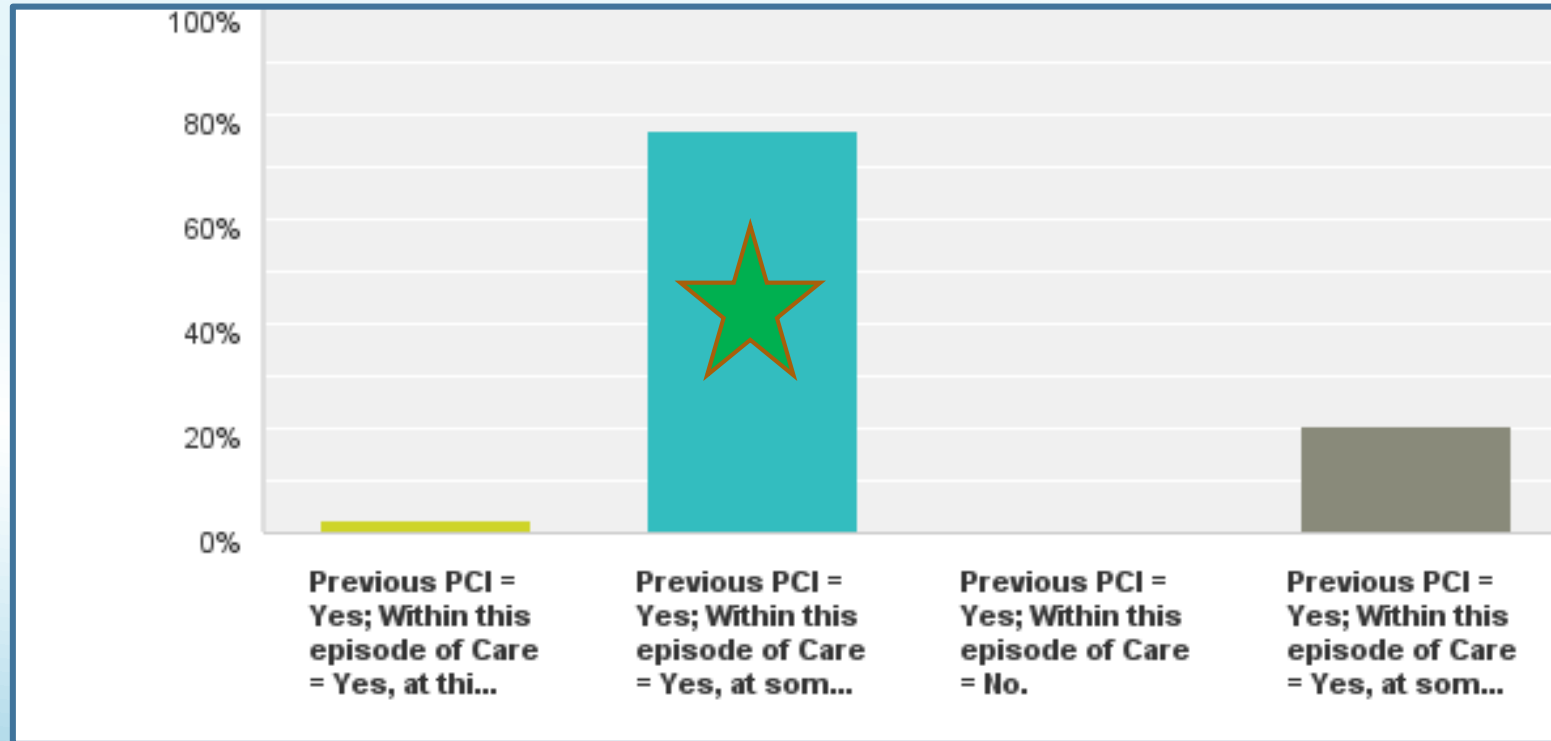
- **PCI complication** - complication during PCI necessitating surgical intervention such as dissection or acute occlusion
- **PCI failure with clinical deterioration** - PCI failed to yield expected and/or desired results, patient condition deteriorated, includes attempts to cross with the wire but unsuccessful
- **PCI for STEMI, multivessel disease** - STEMI with primary PCI (of culprit lesion) and multivessel disease requiring CABG
- **PCI failure without clinical deterioration** - PCI failed to yield expected and/or desired results, patient condition did not deteriorate, includes attempts to cross with the wire but unsuccessful
- **PCI/Surgery staged procedure (not STEMI)** - PCI and surgical procedures performed in a staged fashion in a patient not experiencing STEMI.
- **Other** - other indication for surgery not described above

Question 2 Selections:

- Previous PCI = Yes; Within this episode of Care = Yes, at this facility; Indication for Surgery = Failure w/o Clinical Deterioration.
- Previous PCI = Yes; Within this episode of Care = Yes, at some other facility; Indication for Surgery = Failure w/o Clinical Deterioration.
- Previous PCI = Yes; Within this episode of Care = No.
- Previous PCI = Yes; Within this episode of Care = Yes, at some other facility; Indication for Surgery = PCI/Surgery Staged (not STEMI).



Question 2 Results



Answer Choices	Responses
Previous PCI = Yes; Within this episode of Care = Yes, at this facility; Indication for Surgery = Failure w/o Clinical Deterioration.	2.56% 1
Previous PCI = Yes; Within this episode of Care = Yes, at some other facility; Indication for Surgery = Failure w/o Clinical Deterioration.	76.92% 30
Previous PCI = Yes; Within this episode of Care = No.	0.00% 0
Previous PCI = Yes; Within this episode of Care = Yes, at some other facility; Indication for Surgery = PCI/Surgery Staged (not STEMI).	20.51% 8
Total	39

Question 3 Selections:

- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = NSTEMI; Prior HF = No; HF w/in 2 wks = Yes (Class IV).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Prior HF = Unknown; HF w/in 2 wks = Yes (Class III).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Anginal Class = CCS III; HF w/in 2 wks = Yes (Class III).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Other; Anginal Class w/in 2 wks. = CCS IV; HF w/in 2 wks = Yes (Class IV).

Data Points to Consider

- What was the Cardiac Presentation at Admission?
- What was the Cardiac Presentation at Surgery?
- Was there a history of Prior Heart Failure ?
- Did the patient suffer from Heart Failure w/in last 2 weeks?
- Anginal and Heart Failure Classification

F. Preoperative Cardiac Status

Prior Myocardial Infarction: **PrevMI (885)** Yes No Unknown (If Yes ↓)

MI When: ≤6 Hrs. >6 Hrs. but <24 Hrs. 1 to 7 Days 8 to 21 Days >21 Days

MIWhen (890)

Cardiac Presentation/Symptoms: (Choose one from the list below for each column↓)

	At time of this admission: CardSympTimeOfAdm (895)	At time of surgery: CardSympTimeOfSurg (900)
No Symptoms		
Stable Angina		
Unstable Angina		
Non-ST Elevation MI (Non-STEMI)		
ST Elevation MI (STEMI)		
Angina Equivalent		
Other		

Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 895

Long Name: Cardiac Presentation/Symptoms - At Time Of This Admission; **Short Name:** CardSympTimeOfAdm

Definition: Indicate the patient's cardiac symptoms at the time of this admission. Cardiac presentation is not for angina only.

Intent/Clarification: Indicate the patient's cardiac presentation / symptoms. *Choose the worst status.*

- **No symptoms** – No angina, no acute STEMI, non-STEMI, no anginal equivalent, and no other atypical chest pain.
- **Stable angina** without a change in frequency or pattern for the 6 weeks prior. Angina is controlled by rest and/or oral or transcutaneous medications.
- **Unstable angina:** There are three principal presentations of unstable angina:
 - Rest angina (occurring at rest and prolonged, usually >20 minutes);
 - New-onset angina (within the past 2 months, of at least Canadian Cardiovascular Society Class III severity); or
 - Increasing angina (previously diagnosed angina that has become distinctly more frequent, longer in duration, or increased by 1 or more Canadian Cardiovascular Society class to at least CCS III severity).
- **Non-STEMI** The patient was hospitalized for a non-ST elevation myocardial infarction (NSTEMI) as documented in the medical record. Non-STEMIs are characterized by the presence of **both** criteria:
 - Cardiac biomarkers (creatinine kinase-myocardial band, Troponin T or I) exceed upper limit of normal according to the individual hospitals. Laboratory confirmation of myocardial necrosis; laboratory parameters with a clinical presentation consistent or suggestive of ischemia. ECG changes and/or ischemic symptoms may or may not be present
 - Absence of ECG changes diagnostic of a STEMI (see STEMI).
- **ST-Elevation MI (STEMI)** or equivalent. The patient presented with a ST elevation myocardial infarction (STEMI) or its equivalent as documented in the medical record. STEMI is characterized by the presence of both criteria:
 - ECG evidence of STEMI: New/presumed new ST-segment elevation or new left bundle branch block not documented to be resolved within 20 minutes.
 - Cardiac biomarkers (creatinine kinase-myocardial band, Troponin T or I) exceed the upper limit of normal according to the individual hospital's laboratory parameters and a clinical presentation which is consistent or suggestive of ischemia. Note: For purposes of the Registry, ST elevation in the posterior chest leads (V7 through V9), or ST depression that is maximal in V1-3, without ST-segment elevation in other leads, demonstrating posterobasal myocardial infarction, is considered a STEMI equivalent.
- **Anginal Equivalent** - An anginal equivalent is a symptom such as shortness of breath (dyspnea), diaphoresis, extreme fatigue, or belching, occurring in a patient at high cardiac risk. Anginal equivalents are considered to be symptoms of myocardial ischemia. Anginal equivalents are considered to have the same importance as angina pectoris in patients presenting with elevation of cardiac enzymes or certain EKG changes which are diagnostic of myocardial ischemia. For the patient with diabetes who presents with “silent angina”, code anginal equivalent.
- **Other** – Aortic dissections, sudden death, heart block, arrhythmia, syncope or heart failure.

F. Preoperative Cardiac Status

Prior Myocardial Infarction: **PrevMI (885)** Yes No Unknown (If Yes ↓)

MI When: ≤6 Hrs. >6 Hrs. but <24 Hrs. 1 to 7 Days 8 to 21 Days >21 Days

MIWhen (890)

Cardiac Presentation/Symptoms: (Choose one from the list below for each column↓)

	At time of this admission: CardSympTimeOfAdm (895)	At time of surgery: CardSympTimeOfSurg (900)
No Symptoms		
Stable Angina		
Unstable Angina		
Non-ST Elevation MI (Non-STEMI)		
ST Elevation MI (STEMI)		
Angina Equivalent		
Other		



Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 895

Long Name: Cardiac Presentation/Symptoms - At Time Of This Admission; **Short Name:** CardSympTimeOfAdm



Definition: Indicate the patient's cardiac symptoms at the time of this admission. Cardiac presentation is not for angina only.

Intent/Clarification: Indicate the patient's cardiac presentation / symptoms. *Choose the worst status.*

F. Preoperative Cardiac Status

Prior Myocardial Infarction: **PrevMI (885)** Yes No Unknown (If Yes ↓)
 MI When: ≤6 Hrs. >6 Hrs. but <24 Hrs. 1 to 7 Days 8 to 21 Days >21 Days
MIWhen (890)

Cardiac Presentation/Symptoms: (Choose one from the list below for each column↓)

	At time of this admission: CardSympTimeOfAdm (895)	At time of surgery: CardSympTimeOfSurg (900)
No Symptoms		
Stable Angina		
Unstable Angina		
Non-ST Elevation MI (Non-STEMI)		
ST Elevation MI (STEMI)		
Angina Equivalent		
Other		

Seq. #: 900

Definition: Indicate the patient's cardiac symptoms at the time of awake, entry to the operating room.

Intent/Clarification:

The *intent is to capture changes between admission and surgery; whether a patient improves or deteriorates*. Same definition as Seq. #895, although timeframes may overlap.

- If the patient did not improve or deteriorate between admission and surgery, the code will be the same.
- For elective admissions, patient symptoms (same value/answer) will be entered twice for seq. #895 and 900.
- **If the patient presents with STEMI or Non-STEMI, they should be coded as such in both sequence numbers 895 and 900 unless the patient remains longer than 7 days** and in that case presentation at the time of admission would be STEMI or Non-STEMI and at the time of surgery would be coded as unstable angina.
- Unstable angina at the time of admission would be coded unstable angina at the time of surgery.

Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 905

Long Name: Anginal Classification within 2 weeks **Short Name:** AnginalClass

Definition: Indicate the patient's anginal classification or symptom status within the past 2 weeks.

The anginal classification or symptom status is classified as *the highest grade of angina or chest pain* by the Canadian Cardiovascular Angina Classification System (CCS).

Intent/Clarification: Canadian Cardiovascular Angina Class - Indicate the patient's CCA Class:

- **CCS 0.** The patient has no angina.
- **CCA I.** Ordinary physical activity (for example, walking or climbing stairs) does not cause angina; angina occurs with strenuous or rapid or prolonged exertion at work or recreation
- **CCA II.** Slight limitation of ordinary activity (for example, angina occurs walking or stair climbing after meals, in cold, in wind, under emotional stress, or only during the few hours after awakening; walking more than 2 blocks on the level or climbing more than 1 flight of ordinary stairs at a normal pace; and in normal conditions)
- **CCA III.** Marked limitation of ordinary activity (for example, angina occurs with walking 1 or 2 blocks on the level or climbing 1 flight of stairs in normal conditions and at a normal pace)
- **CCA IV.** Inability to perform any physical activity without discomfort; *angina syndrome may be present at rest*. All other classes of pain go away with rest and/or treatment.

Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 910

Long Name: Heart Failure within 2 weeks; **Short Name:** CHF

Definition: Indicate if there is physician documentation the patient has been in a state of heart failure within the past 2 weeks.

Heart failure is *defined as physician documentation or report of any of the following clinical symptoms of heart failure described as unusual dyspnea on light exertion, recurrent dyspnea occurring in the supine position, fluid retention; or the description of rales, jugular venous distension, pulmonary edema on physical exam, or pulmonary edema on chest x-ray presumed to be cardiac dysfunction.* A low ejection fraction alone, without clinical evidence of heart failure does not qualify as heart failure. An elevated BNP without other supporting documentation should not be coded as CHF.

Intent/Clarification: **Capture the** patient's actual status in the weeks before surgery, the *new diagnosis or exacerbation of an existing heart failure condition.*

DO NOT code stable or asymptomatic compensated failure or patients whose symptoms improved after medical therapy. A low ejection fraction (EF) without clinical presentation does not qualify for history of heart failure

Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 915

Long Name: Classification-NYHA; Short Name: ClassNYH

Definition: Indicate the patient's *worst dyspnea or functional class*, coded as the New York Heart Association (NYHA) classification within the past 2 weeks.

This is to be used for heart failure only, is not intended to classify angina.

Intent/Clarification:

NYHA is for congestive heart failure (CHF).

Select the **highest level of heart failure within the two weeks leading up to episode of hospitalization or at the time of the procedure**. The intent is to capture the highest level of failure. If the NYHA class is not documented, use the guidelines below to assign a class based on documented symptoms.

- **Class I:** Patient has cardiac disease but without resulting limitations of ordinary physical activity. Ordinary physical activity (e.g., walking several blocks or climbing stairs) does not cause undue fatigue, palpitation, dyspnea, or anginal pain. Limiting symptoms may occur with marked exertion.
- **Class II:** Patient has cardiac disease resulting in slight limitation of ordinary physical activity. Patient is comfortable at rest. Ordinary physical activity such as walking more than two blocks or climbing more than one flight of stairs results in limiting symptoms (e.g., fatigue, palpitation, dyspnea, or anginal pain).
- **Class III:** Patient has cardiac disease resulting in marked limitation of physical activity. Patient is comfortable at rest. Less than ordinary physical activity (e.g., walking one to two level blocks or climbing one flight of stairs) causes fatigue, palpitation, dyspnea, or anginal pain.
- **Class IV:** Patient has dyspnea at rest that increases with any physical activity. Patient has cardiac disease resulting in inability to perform any physical activity without discomfort. Symptoms may be present even at rest. If any physical activity is undertaken, discomfort is increased.

Anginal Classification Within 2 weeks: CCS Class 0 CCS Class I CCS Class II CCS Class III CCS Class IV

AnginalClass (905)

Heart Failure Within 2 weeks : Yes No Unknown (If Yes→) Classification-NYHA: Class I Class II Class III Class IV

CHF (910)

ClassNYH (915)

Prior Heart failure: Yes No Unknown

PriorHF (920)

Seq. #: 920

Long Name: Prior Heart failure; **Short Name:** PriorHF

Definition: Indicate history of *heart failure occurring more than 2 weeks prior to current episode of care.*

A previous hospital admission with principal diagnosis of heart failure is considered evidence of heart failure history but is not essential.

Intent/Clarification:

The goal is to capture patients who have improved following medical management and do not exhibit clinical signs of failure within 2 weeks of surgery but have documented failure symptoms prior to that.

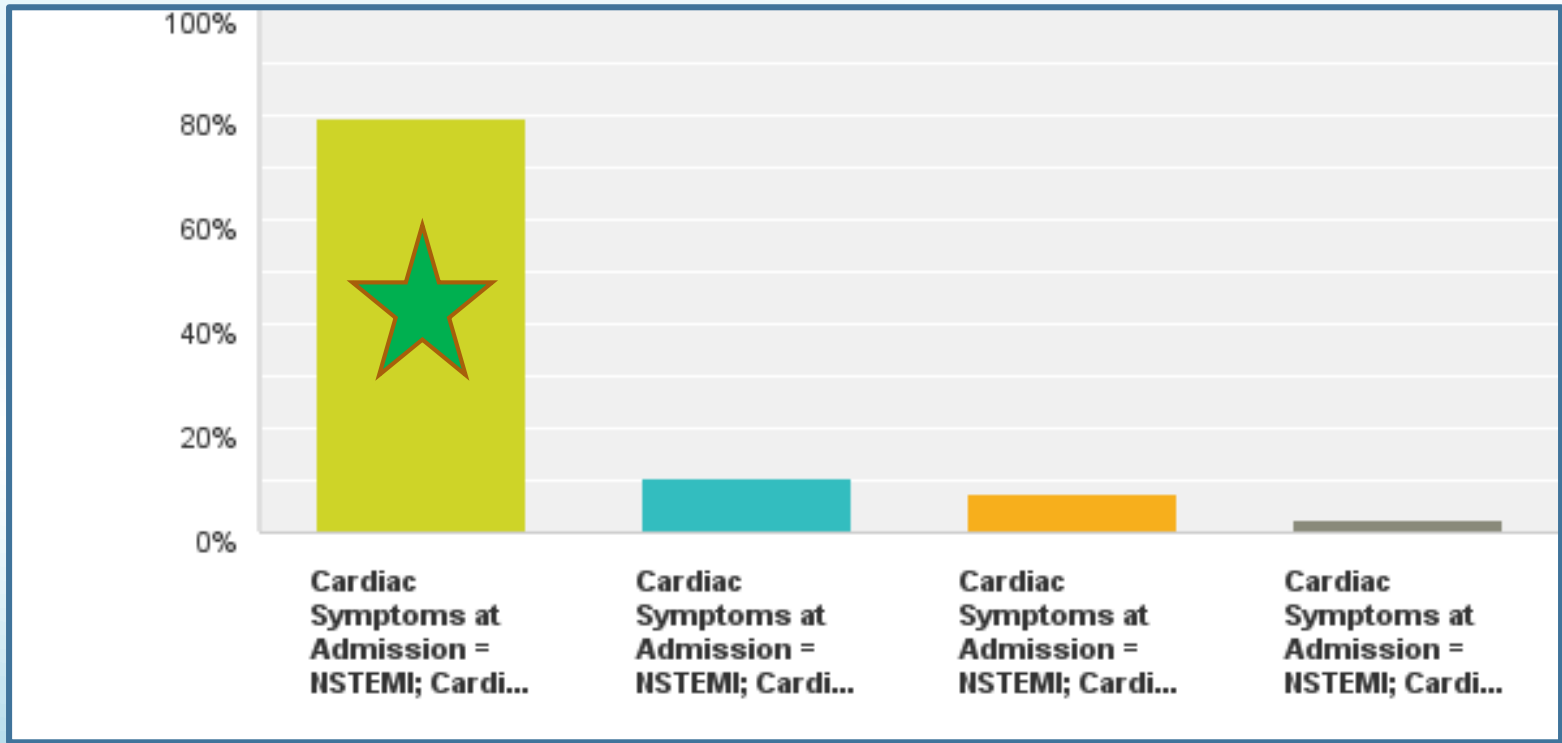
Newly diagnosed HF that is described to have onset of symptoms over past few months (but not previously known or treated) that is now worsening –code yes to both 910 and 920

Question 3 Selections:



- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = NSTEMI; Prior HF = No; HF w/in 2 wks = Yes (Class IV).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Prior HF = Unknown; HF w/in 2 wks = Yes (Class III).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Anginal Class = CCS III; HF w/in 2 wks = Yes (Class III).
- Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Other; Anginal Class w/in 2 wks. = CCS IV; HF w/in 2 wks = Yes (Class IV).

Question 3 Results



Answer Choices	Responses
Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = NSTEMI; Prior HF = No; HF w/in 2 wks = Yes (Class IV).	79.49% 31
Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Prior HF = Unknown; HF w/in 2 wks = Yes (Class III).	10.26% 4
Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Unstable Angina; Anginal Class = CCS III; HF w/in 2 wks = Yes (Class III).	7.69% 3
Cardiac Symptoms at Admission = NSTEMI; Cardiac Symptoms at Surgery = Other; Anginal Class w/in 2 wks. = CCS IV; HF w/in 2 wks = Yes (Class IV).	2.56% 1
Total	39

Question 4 Selections:

- Number of Diseased Vessels = 3; EF = 25; Mitral Insufficiency = Severe; Aortic, Tricuspid, and Pulmonic Insufficiency = None
- Number of Diseased Vessels = 3; EF = 30; Mitral Insufficiency = Severe; Aortic Insufficiency = Not Documented.
- Number of Diseased Vessels = 2; EF = 30; Mitral Insufficiency = Moderate; Aortic Insufficiency = Not Documented.
- Number of Diseased Vessels = 2; EF = 25; Mitral Insufficiency = Moderate; Aortic Insufficiency = None.

Data Points to Consider

- What were the number of diseased Vessels?
- Determination of Ejection Fraction.
- What degree of insufficiency was present preoperatively in the native valves?

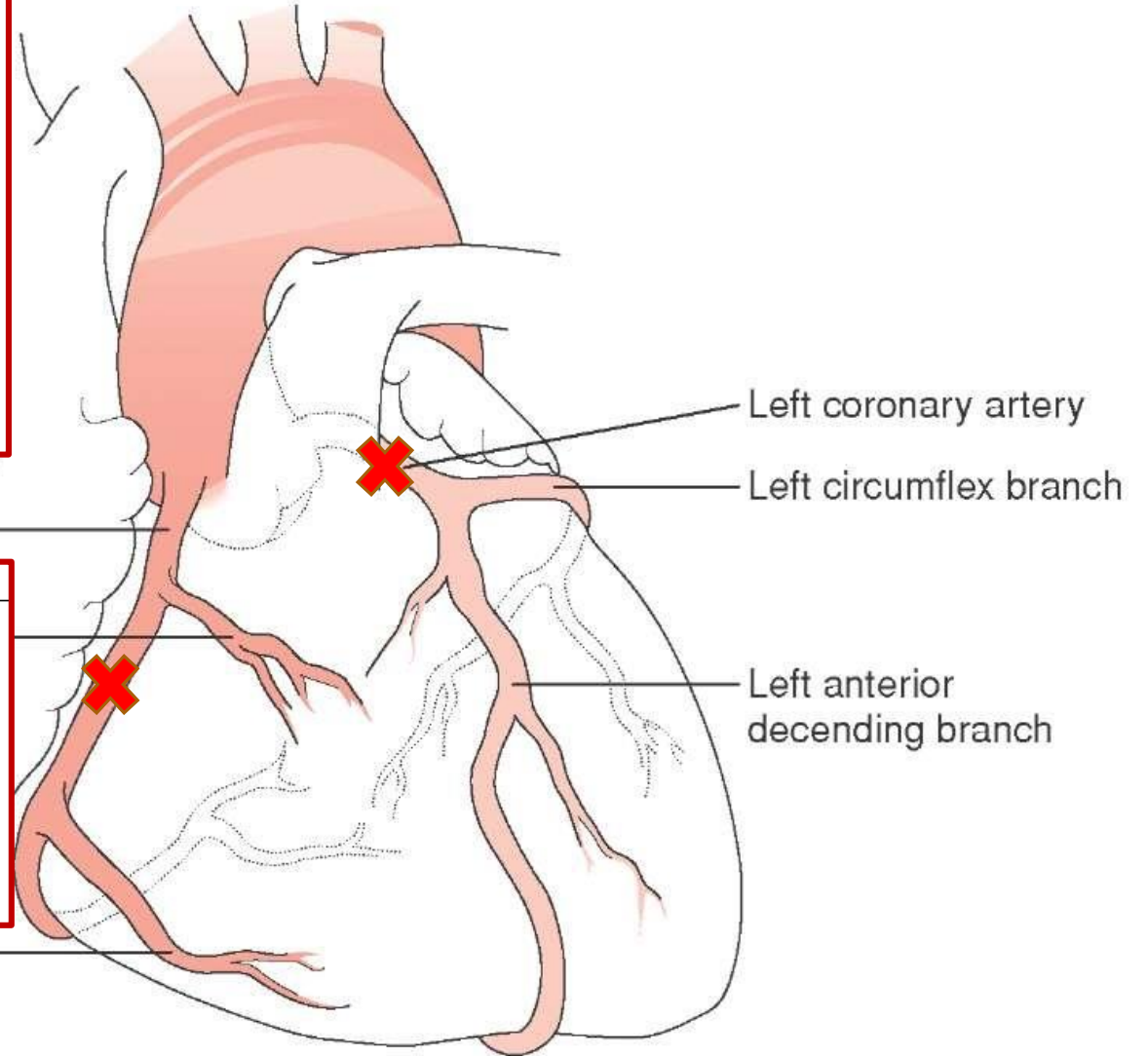
Coronary Anatomy

Seq. #: 1170

Long Name: Num Dis Vessels; Short Name: NumDisV

Definition: Indicate the number of **diseased major native coronary vessel systems**: LAD system, Circumflex system, and/or Right system with $\geq 50\%$ narrowing of any vessel preoperatively.

NOTE: Left main disease ($\geq 50\%$) is counted as TWO vessels (LAD and Circumflex, which may include a Ramus Intermedius). For example, left main and RCA would count as three total.



H. Hemodynamics/Cath/Echo

Cardiac Catheterization Performed: Yes No (If Yes→)

CarCathPer (1145)

Coronary Anatomy/Disease known: Yes No (If Yes↓)

CorAnatDisKnown (1155)

Dominance:

Dominance (1160)

Source(s) used to quantify stenosis: **StenSource** (1165)

Number Diseased Vessels:

NumDisV (1170)

Cardiac Catheterization Date: ___/___/___

CarCathDt (1150)

Left Right Co-dominant Not Documented

Angiogram CT IVUS Progress/OP Note Other

Multiple

None One Two Three

Marginal branch

Ejection Fraction

Seq. #: 1540

Long Name: Hemo Data-EF Done; **Short Name:** HDEFD

Definition: Indicate whether the Ejection Fraction was measured prior to the induction of anesthesia.

Intent/Clarification:

Some patients may not have had an LV Gram performed during cardiac catheterization due to existing clinical conditions. Ejection fraction (EF) and hemodynamic pressures may be obtained from other sources other than coronary angiogram, such as echo, or MUGA.

Note: Because anesthesia can alter the values to be collected, do not collect data from intra-operative transesophageal echo (TEE) after the induction of anesthesia, **unless** you have no other source to collect the information.

Time Frame: Do not use results more than 6 months prior to this operation.

Ejection Fraction Done: HDEFD (1540) Yes No (If Yes→) Ejection Fraction: HDEF (1545) 30 (%)

Seq. #: 1545

Long Name: Hemo Data-EF; **Short Name:** HDEF

Definition: Indicate the percentage of the blood emptied from the left ventricle at the end of the contraction. Use the most recent determination prior to the surgical intervention documented on a diagnostic report.

Enter a percentage in the range of 1 - 99. If a percentage range is reported, report a whole number using the "mean" (i.e., 50-55% is reported as 53%).

- Hyperdynamic: >70%
- Normal: 50%–70% (midpoint 60%)
- Mild dysfunction: 40%–49% (midpoint 45%)
- Moderate dysfunction: 30%–39% (midpoint 35%)
- Severe dysfunction: <30%

Note: If no diagnostic report is in the medical record, a value documented in the medical record is acceptable. ACCF/AHA 2013

Intent/Clarification:

Time Frame: Collect the last value closest to incision, not greater than 6 months.

Use the most recent determination prior to the induction of anesthesia documented on a diagnostic report, regardless of the diagnostic procedure to obtain it.

Note: If no diagnostic report specifying an EF is in the medical record, a value documented in the progress record is acceptable.

Note: If there is no documentation of a pre-op EF, **then** it is acceptable to code the EF from the intra-op TEE prior to incision.

Valve Insufficiency and Disease

Seq. #: 1680

Long Name: VD-Insuff-Mitral

Short Name: VDInsufM

The same rules apply to all four valves

Definition: Indicate whether there is evidence of Mitral valve insufficiency/regurgitation. Enter the level of valve function associated with highest risk (i.e., worst performance). Enter the highest level recorded in the chart. "Moderately severe" should be coded as "Severe".

Intent/Clarification:

Time Frame: Collect the last value closest to incision, not greater than 6 months.

Choose the highest level of valve dysfunction when there are differences in interpretation of the most recent study.

Capture even if patient is not scheduled for valve repair and/or replacement when available.

January 2016 FAQ: Clarify coding of valve disease from echocardiograms.

If there is a preoperative echo, use those values UNLESS the diagnostic information from the TEE changes the procedure performed. If there is no pre-op information, you may use the pre-incision intraoperative TEE.

Seq. #: 1685

Long Name: VD-Mitral; Short Name: VDMit

Definition: Indicate whether Mitral valve disease is present.

Intent/Clarification:

When insufficiency is noted in the valve, at what level should the valve be considered diseased? The valve should be coded as being diseased if there is mild, moderate or severe insufficiency.

Mitral Valve

Mitral Insufficiency: **VDInsufM (1680)** None Trivial/Trace Mild Moderate Severe Not Documented

Mitral Valve Disease: **VDMit (1685)** Yes No

(If Yes→) Mitral Stenosis: Yes No (If Yes→)

VDStenM (1690)

Hemodynamic/ Echo data available: Yes No (If Yes ↓)

MiHemoDatAvail (1695)

Smallest Valve Area: _____ cm²

VDMVA (1700)

Highest Mean Gradient: _____ mmHg

There is not an option to select “Not Documented” for Valve Disease. When an Echo is not done, or a valve is not mentioned in the study, the best answer is “No” if there is no other documentation in the medical record that there is valve disease.

None = Valves were Studied & No Insufficiency Found

Not Documented = Procedure to Study Valves Not Done (or not mentioned)

No means No

Aortic Valve

Aortic Insufficiency: None Trivial/Trace Mild Moderate Severe Not Documented

VDInsufA (1590)

Aortic Valve Disease: VDAort (1595) Yes No

(If Yes→) Aortic Stenosis: Yes No (If Yes→) Hemodynamic/Echo data available: Yes No (If Yes ↓)

VDStenA (1600) AoHemoDatAvail (1605)

Tricuspid Valve

Tricuspid Insufficiency: VDInsufT (1775) None Trivial/Trace Mild Moderate Severe Not Documented

Tricuspid Valve Disease: VDTTr (1780) Yes No

(If Yes→) Tricuspid Stenosis: VDStenT (1785) Yes No

Pulmonic Valve

Pulmonic Insufficiency: None Trivial/Trace Mild Moderate Severe Not Documented

VDInsufP (1820)

Pulmonic Valve Disease: Yes No

VDPulm (1825)

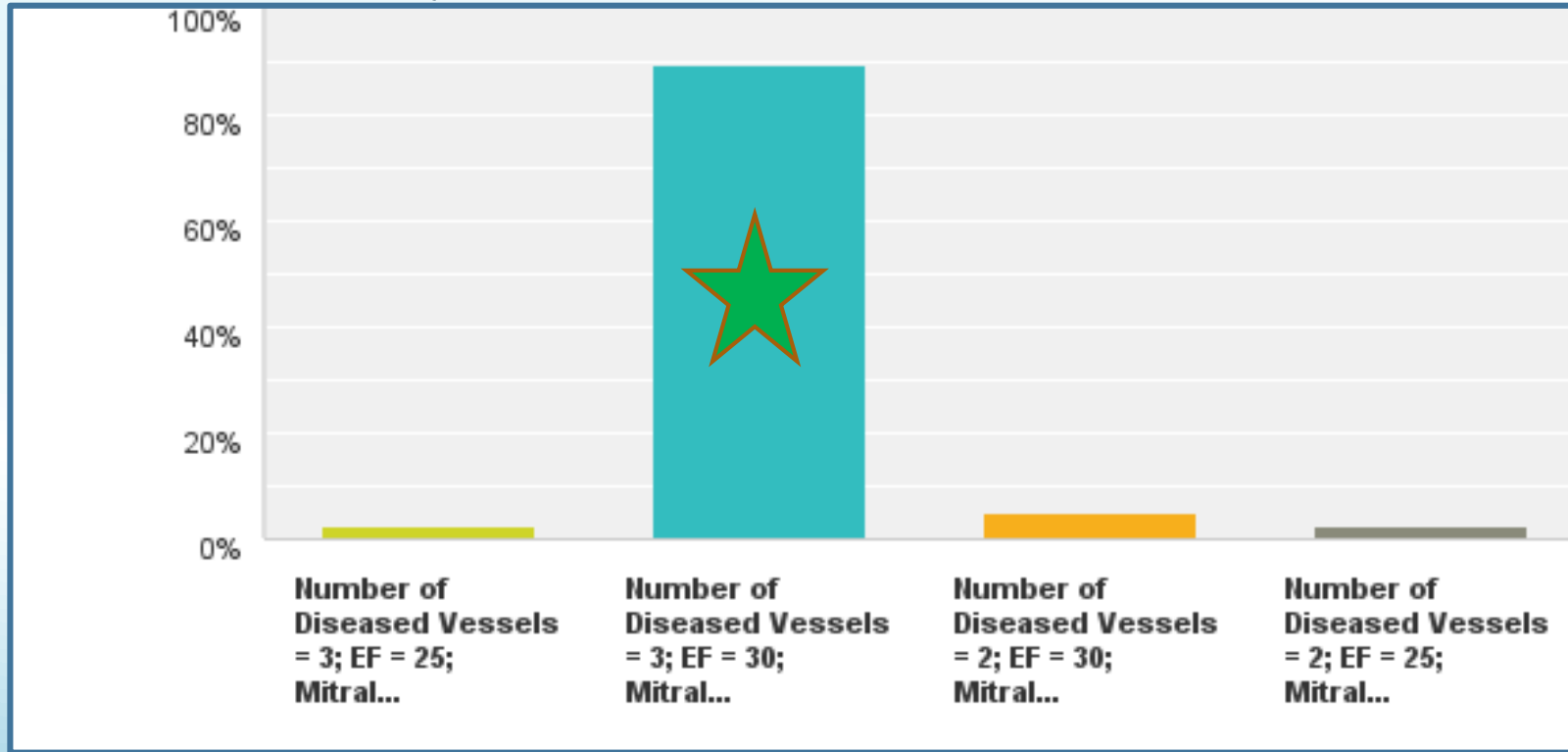
Question 4 Selections:

- Number of Diseased Vessels = 3; EF = 25; Mitral Insufficiency = Severe; Aortic, Tricuspid, and Pulmonic Insufficiency = None



- Number of Diseased Vessels = 3; EF = 30; Mitral Insufficiency = Severe; Aortic Insufficiency = Not Documented.
- Number of Diseased Vessels = 2; EF = 30; Mitral Insufficiency = Moderate; Aortic Insufficiency = Not Documented.
- Number of Diseased Vessels = 2; EF = 25; Mitral Insufficiency = Moderate; Aortic Insufficiency = None.

Question 4 Results



Answer Choices	Responses
Number of Diseased Vessels = 3; EF = 25; Mitral Insufficiency = Severe; Aortic, Tricuspid, and Pulmonic Insufficiency = None	2.56% 1
Number of Diseased Vessels = 3; EF = 30; Mitral Insufficiency = Severe; Aortic Insufficiency = Not Documented.	89.74% 35
Number of Diseased Vessels = 2; EF = 30; Mitral Insufficiency = Moderate; Aortic Insufficiency = Not Documented.	5.13% 2
Number of Diseased Vessels = 2; EF = 25; Mitral Insufficiency = Moderate; Aortic Insufficiency = None.	2.56% 1
Total	39

Question 5 Selections:

- Status = Urgent; Reason = PCI Incomplete w/o deterioration; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent)
- Status = Emergent; Reason = Pulmonary Edema; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged)
- Status = Urgent; Reason = AMI; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged).
- Status = Emergent; Reason = CHF; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent).

Data Points to Consider

- What is the patient's Operative Status?
- What is the best Reason for Operative Status?
- What surgical procedures were performed?
- Was this a "Combined" case? What constitutes "Combined"?

Status

Seq. #: 1975

Long Name: Status; **Short Name:** Status

Definition: Indicate the clinical status of the patient prior to entering the operating room.

Elective: The patient's cardiac function has been stable in the days or weeks prior to the operation. The *procedure could be deferred without increased risk of compromised cardiac outcome.*

Urgent: Any of the conditions that require that *the patient remain in the hospital until surgery can take place, but the patient is able to wait for surgery until the next available OR schedule time.* Delay in the operation may be necessitated by attempts to improve the patient's condition, availability of a spouse or parent for informed consent, availability of blood products, or the availability of results of essential laboratory procedures or tests.

Emergent: An emergency operation is one in which there should be *no delay in providing operative intervention.* Patients requiring emergency operations will have ongoing, refractory (difficult, complicated, and/or unmanageable) cardiac compromise, with or without hemodynamic instability, and not responsive to any form of therapy except cardiac surgery.

Emergent/Salvage: The patient is *undergoing CPR in route to the OR* prior to anesthesia induction *or has ongoing ECMO to maintain life.*

Status:

Elective



Urgent

Emergent


Emergent Salvage

Status (1975)

UrgEmergRsn (1990)

(If Urgent or Emergent choose one reason↓)

Urgent/ Emergent reason:

- | | |
|--|---|
| <input type="checkbox"/>  AMI | <input type="checkbox"/> PCI Incomplete without clinical deterioration |
| <input type="checkbox"/> Anatomy | <input type="checkbox"/> PCI or attempted PCI with Clinical Deterioration |
| <input type="checkbox"/> Aortic Aneurysm | <input type="checkbox"/> Pulmonary Edema |
| <input type="checkbox"/> Aortic Dissection | <input type="checkbox"/> Pulmonary Embolus |
| <input type="checkbox"/> CHF | <input type="checkbox"/> Rest Angina |
| <input type="checkbox"/> Device Failure | <input type="checkbox"/> Shock Circulatory Support |
| <input type="checkbox"/> Diagnostic/Interventional Procedure Complication | <input type="checkbox"/> Shock No Circulatory Support |
| <input type="checkbox"/> Endocarditis | <input type="checkbox"/> Syncope |
| <input type="checkbox"/> Failed Transcatheter Valve Therapy | <input type="checkbox"/> Transplant |
| <input type="checkbox"/> IABP | <input type="checkbox"/> Trauma |
| <input type="checkbox"/> Infected Device | <input type="checkbox"/> USA |
| <input type="checkbox"/> Intracardiac mass or thrombus | <input type="checkbox"/> Valve Dysfunction |
| <input type="checkbox"/> Ongoing Ischemia | <input type="checkbox"/> Worsening CP |
| | <input type="checkbox"/> Other |

Seq. #: 1990

Long Name: Urgent Or Emergent Reason; Short Name: UrgEmergRsn

Definition: Choose one reason from the list below that best describes why this operation was considered urgent or emergent.

Intent/Clarification: See list for options. There may be multiple reasons, **choose one that best describes this patient's clinical state.**

Operative Procedure

Approach converted during procedure: Yes, planned Yes, unplanned No

ApproachCon (2105)

Robot Used: Yes No (If Yes →) Used for entire operation Used for part of the operation

Robotic (2110)

RobotTim (2115)

Coronary Artery Bypass: Yes, planned Yes, unplanned due to surgical complication

Yes, unplanned due to unsuspected disease or anatomy No (If "Yes" complete Section J) OpCAB (2120)

Valve Surgery: Yes No (If "Yes" complete Section K)

OpValve (2125)

VAD Implanted or Removed: Yes No

VADProc (2130)

Other Cardiac Procedure: Yes No (If "Yes" complete Section M)

OpOCard (2140)

Other Cardiac Procedure, AFib: Yes No (If "Yes" complete Section M-1)

AFibProc (2145)

Other Cardiac Procedure, Aortic: Yes, planned Yes, unplanned due to surgical complication

Yes, unplanned due to unsuspected disease or anatomy No (If "Yes" complete Section M-2) AortProc (2150)

Other Non-Cardiac Procedure: Yes No (If "Yes" complete Section N)

Combined Procedures

Seq. #: 2585

Long Name: Combined Cardiac Surgery and PCI Performed; **Short Name:** CombCardPCI

Definition: Indicate whether a cardiac surgical procedure was performed in addition to a PCI during this hospitalization.

Intent/Clarification:

This includes planned and unplanned combinations of cardiac surgery procedures and percutaneous coronary interventions.

Seq. #: 2590

Long Name: Combined Cardiac and PCI Procedures Performed; **Short Name:** CombProcs

Definition: Indicate which procedures were performed during this hospitalization.

Intent/Clarification:

- PCI + CAB
- PCI + Valve
- PCI + Aortic
- PCI + Other

Seq. #: 2595

Long Name: Combined Cardiac Surgery and PCI Procedure Status; **Short Name:** CombProcsStatus

Definition: Indicate whether the procedures were performed concurrently or staged.

Intent/Clarification:

- Concurrent - same setting
- Staged - PCI followed by surgery
- Staged - surgery followed by PCI

Combined cardiac surgery and PCI Performed: Yes No (If Yes ↓)

CombCardPCI (2585)

Procedures: PCI + CAB PCI + Valve PCI + Aortic PCI + Other

CombProcs (2590)

Status: Concurrent- same setting Staged - PCI followed by surgery Staged - Surgery followed by PCI

CombProcsStatus (2595)

PCI Procedure: Angioplasty Stent Angioplasty and Stent Attempted PCI

CombProcsPCI (2600)

(If Stent or Angioplasty & Stent →) Stent Type: Bare metal Drug-eluting Bioresorbable Multiple Not documented

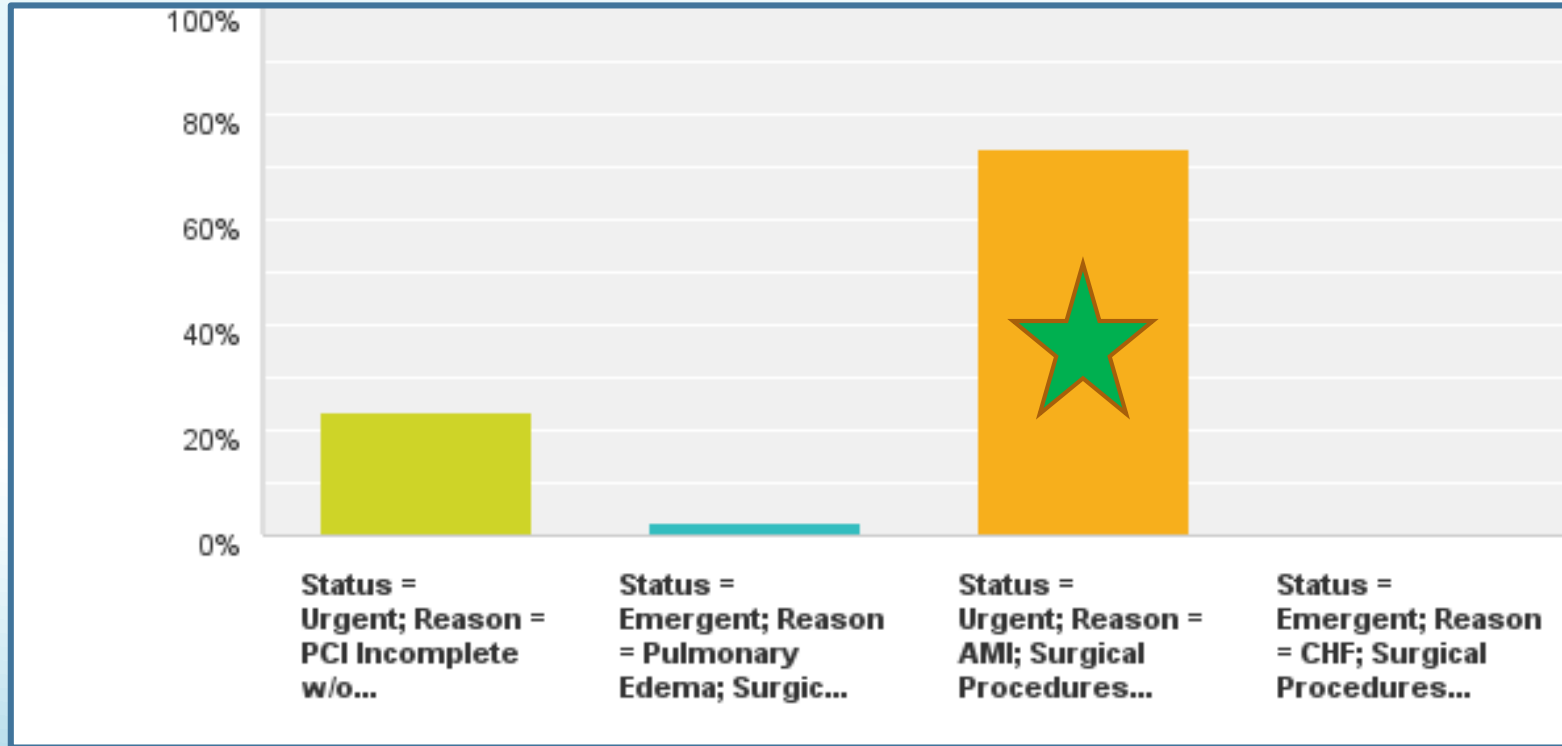
CombProcsStentTy (2605)

Question 5 Selections:

- Status = Urgent; Reason = PCI Incomplete w/o deterioration; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent)
- Status = Emergent; Reason = Pulmonary Edema; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged)
- Status = Urgent; Reason = AMI; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged).
- Status = Emergent; Reason = CHF; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent).



Question 5 Results



Answer Choices	Responses
Status = Urgent; Reason = PCI Incomplete w/o deterioration; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent)	23.68% 9
Status = Emergent; Reason = Pulmonary Edema; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged)	2.63% 1
Status = Urgent; Reason = AMI; Surgical Procedures include: CAB, Valve Surgery, Other Cardiac; Combined Surgery and PCI = Yes (Staged).	73.68% 28
Status = Emergent; Reason = CHF; Surgical Procedures include: CAB, Valve Surgery; Combined Surgery and PCI = Yes (Concurrent).	0.00% 0
Total	38

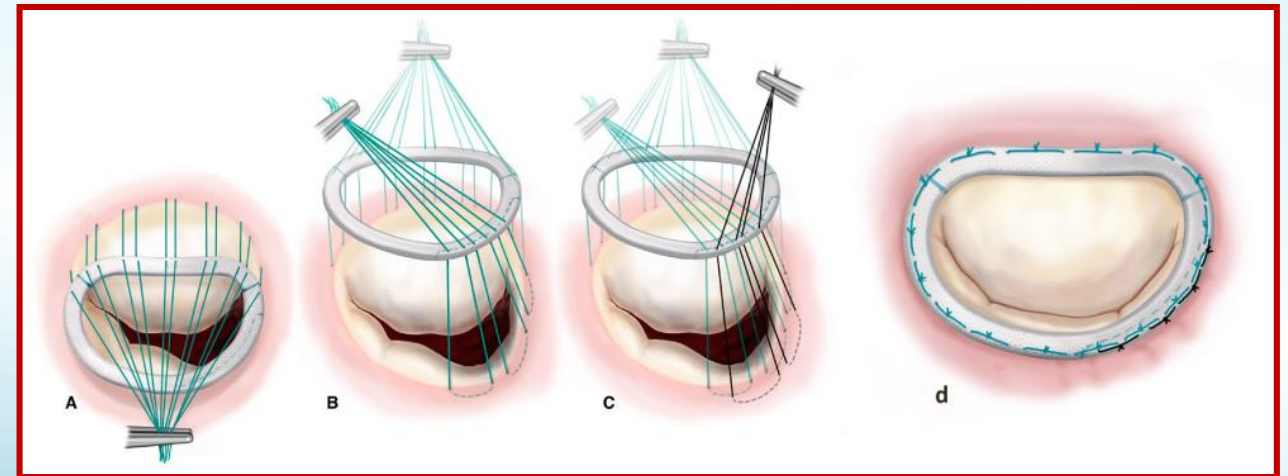
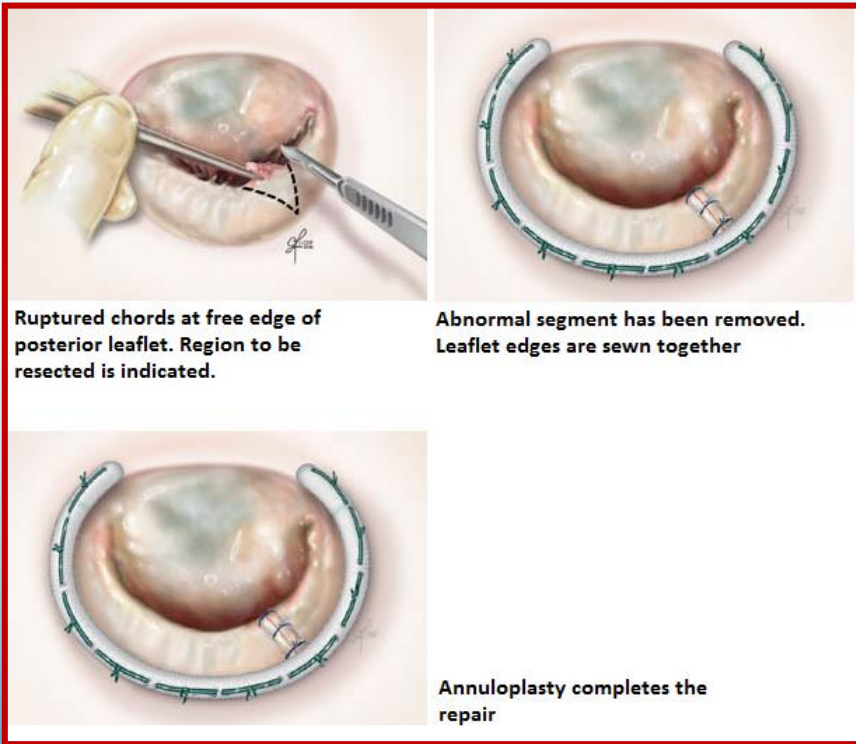
Question 6 Selections:

- Mitral Valve Procedure performed = Repair, Planned; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, CPB Wean.
- Mitral Valve Procedure performed = Repair, Unplanned disease or anatomy; Repair Type = Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, Hemodynamic Instability.
- Mitral Valve Procedure performed = Repair, Planned; Repair Type = Annuloplasty, Sliding Plasty; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, Procedural Support.
- Mitral Valve Procedure performed = Repair, Unplanned disease or anatomy; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, CPB Wean.

Data Points to Consider

- What Mitral Procedure was performed?
- What type of procedure was performed?
- What was the Other Cardiac Procedure performed?
- Why was an IABP inserted?

Mitral Valve Repair



Annuloplasty Only

Leaflet Resection Posterior,
Triangular

Mitral Valve Procedure Performed: Yes, planned Yes, unplanned due to surgical complication Yes, unplanned due to unsuspected disease or anatomy No (If Yes ↓) **VSMV (3495)**

Procedure Performed: **VSMVPr (3500)**
Repair

(If Repair→) Repair Type: (Select all that apply↓)

Annuloplasty Yes No

VSMitRAnnulo (3505)

Leaflet Resection Yes No (If Yes↓)

VSMitRLeafRes (3510)

Resection Type: Triangular Quadrangular Other

VSLeafResTyp (3515)

Location: Anterior Posterior Both Anterior and Posterior

VSLeafRepLoc (3520)

Leaflet Plication Yes No

VSMitRLeafPlic (3525)

Leaflet Debridement Yes No

VSMitRLeafDeb (3530)

Folding Plasty Yes No

VSMitRFold (3535)

Sliding Plasty Yes No

VSMitRSlidP (3540)

Annular decalcification/debridement Yes No

VSMitRADecalc (3545)

Neochords (PTFE) Yes No

VSMitRPTFE (3550)

(If Yes→) # of neochords inserted: _____

VSNeoChNum (3555)

Other Cardiac Procedure

M. Other Cardiac Procedure (If Other Cardiac Procedure = Yes ↓)

These procedures do not impact isolated category

AFib Epicardial lesions (complete M-1) Yes No

OCarAFibEpLes (4070)

ASD repair- PFO type Yes No

OCarASDPFO (4075)

Atrial Appendage procedure: RAA LAA Both No

OCarAAProc (4080)

Arrhythmia Device: OCarACD (4085)

These procedures move the case out of isolated category

AFib Intracardiac lesions (complete M-1) Yes No

OCarAFibIntraLes (4105)

ASD Repair- secundum or sinus venosus Yes No

OCarASDSec (4110)

Lead Extraction Yes, planned

OCarACDLE (4120)

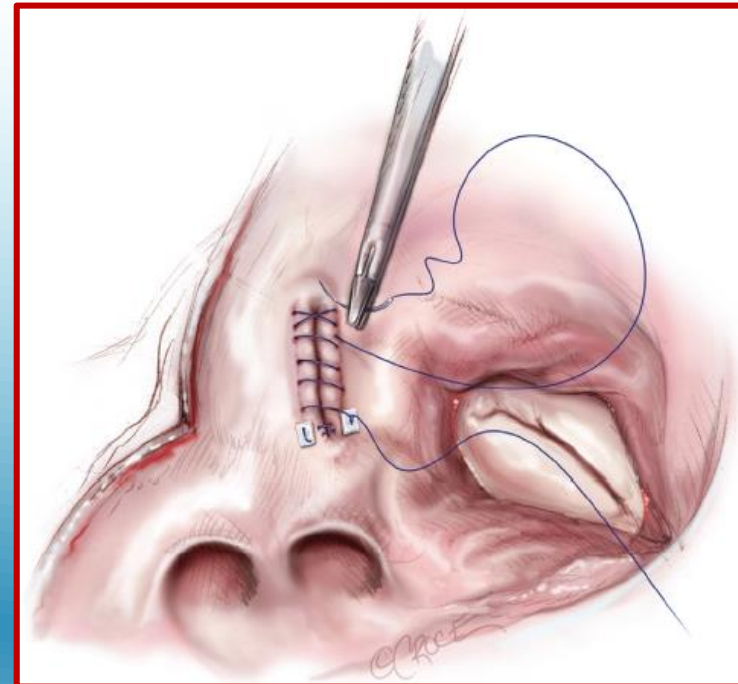
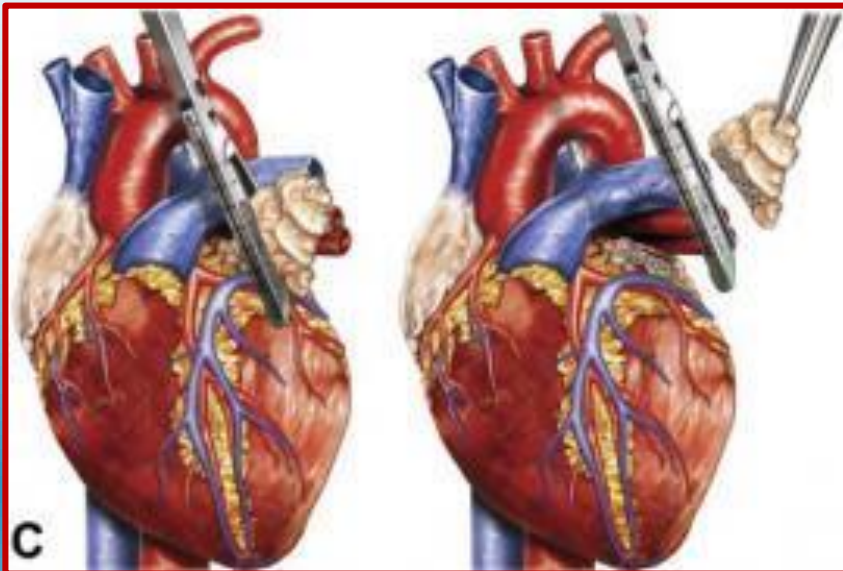
Yes, unplanned due to surgical complication

Yes, unplanned due to unsuspected disease or anatomy

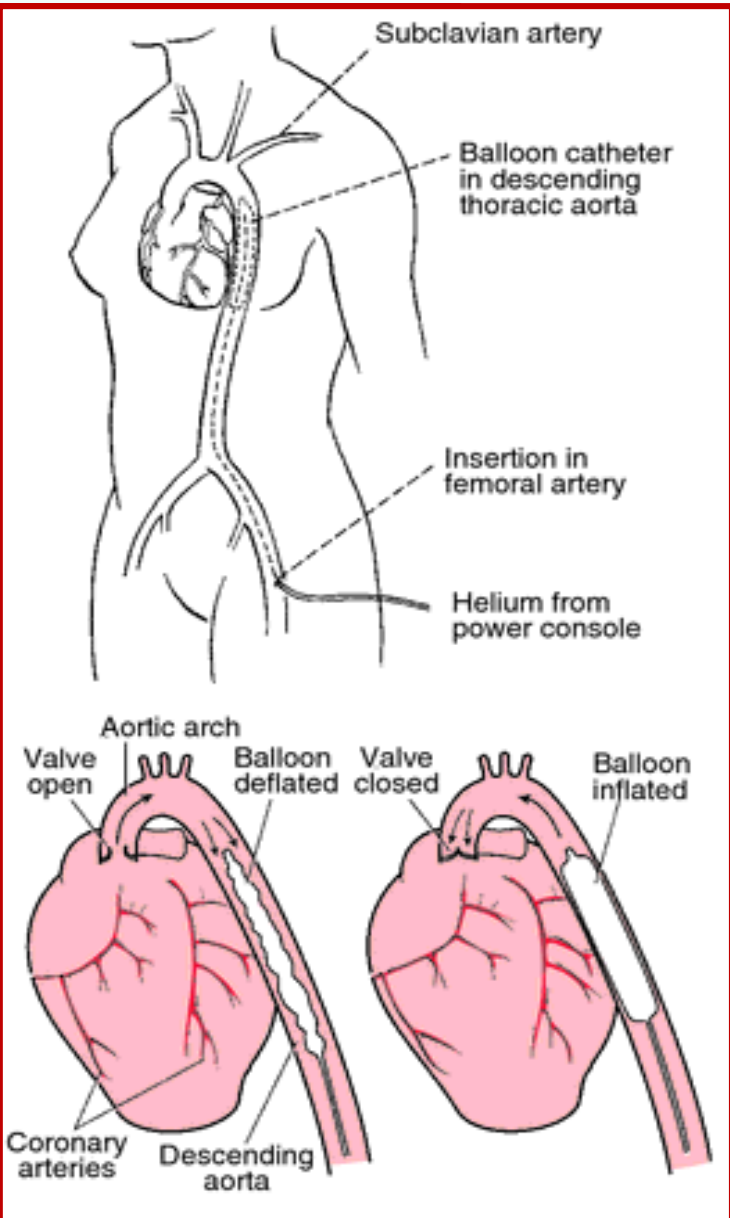
No

LV Aneurysm Repair:

Yes No



INDICATIONS
<ul style="list-style-type: none"> • Left ventricular failure or Cardiogenic Shock a. Myocardial infarction (MI) b. Myocarditis c. Cardiomyopathy d. Severe Myocardial contusion e. Septic shock f. Drug induced
<ul style="list-style-type: none"> • Mechanical Complications of Acute MI
<ul style="list-style-type: none"> • Post - Myocardial Infarction Ventricular Irritability
<ul style="list-style-type: none"> • Unstable Angina refractory to medical therapy
<ul style="list-style-type: none"> • Support for High risk PTCA - Patients
<ul style="list-style-type: none"> • Failed PTCA
<ul style="list-style-type: none"> • Thrombolytic Therapy of Acute MI
<ul style="list-style-type: none"> • Failure to wean from Cardiopulmonary Bypass
<ul style="list-style-type: none"> • Low - Output Syndrome • Stabilization of High - Risk Patients undergoing General Anesthesia • Bridge to Transplant • Stunned Myocardium



IABP(INTRA-AORTIC BALLOON PUMP)

TABLE 54-33. Indications for IABP During Cardiac Surgery
Inability to discontinue bypass: multiple interventions Inadequate hemodynamics: after $\uparrow\uparrow$ inotropic support $\downarrow\downarrow$ Systolic blood pressure \downarrow 80 mm. Hg $\downarrow\downarrow$ Cardiac index \downarrow 2.0 L./min./sq. m. $\uparrow\uparrow$ Left atrial pressure \uparrow 20 mm. Hg $\uparrow\uparrow$ Vascular resistance \uparrow 2500 dynes/sec./cm. ⁻⁵ Large doses of multiple inotropic drugs Continued refractory ventricular arrhythmias

L. Mechanical Cardiac Assist Devices

Intra-Aortic Balloon Pump (IABP): Yes No (If Yes \downarrow)

IABP (3725)


IABP Insertion: Preop Intraop Postop

IABP When (3730)

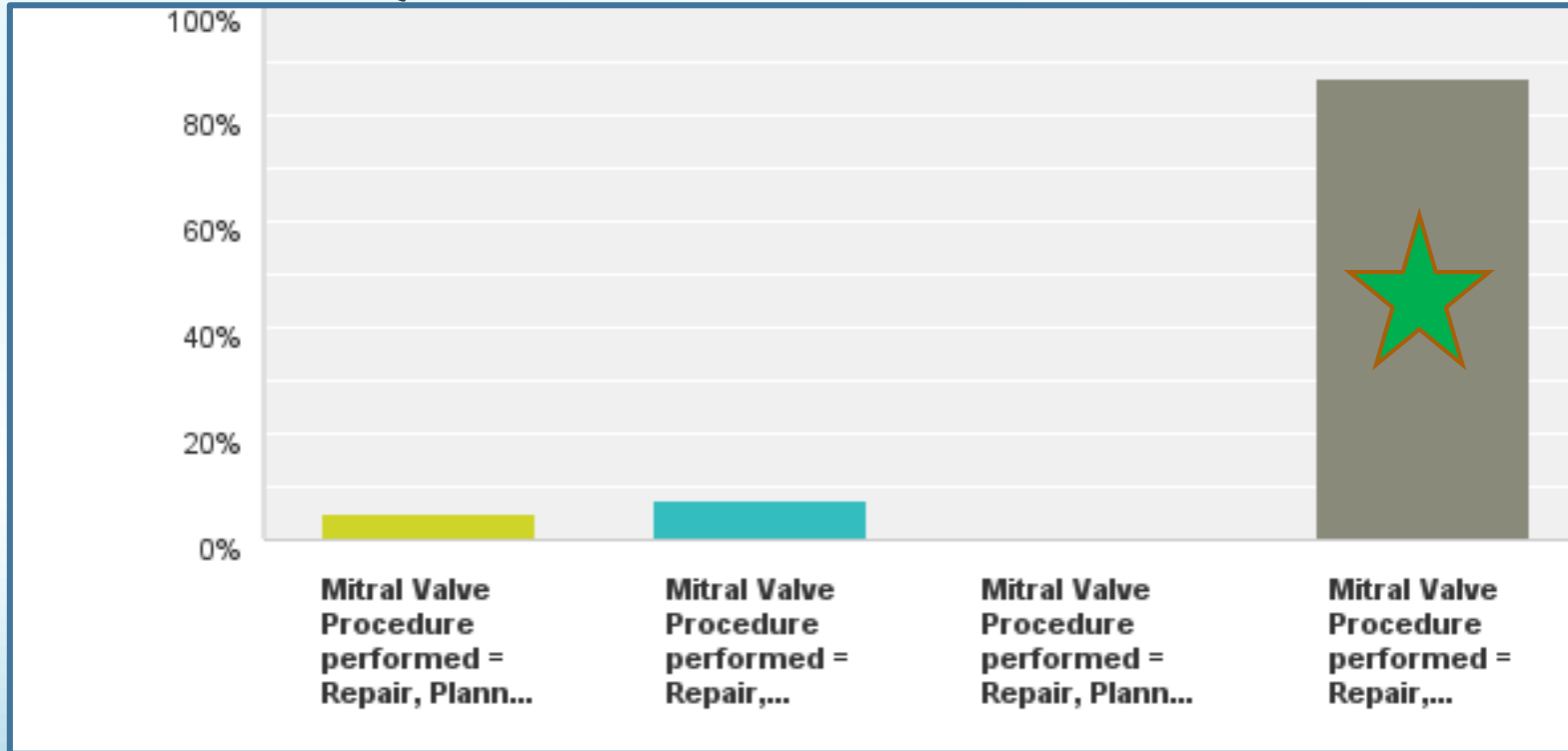
Primary Reason for Insertion: Hemodynamic Instability Procedural Support Unstable Angina

IABP Ind (3735) CPB Weaning Failure Prophylactic Other

Question 6 Selections:

- Mitral Valve Procedure performed = Repair, Planned; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, CPB Wean.
- Mitral Valve Procedure performed = Repair, Unplanned disease or anatomy; Repair Type = Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, Hemodynamic Instability.
- Mitral Valve Procedure performed = Repair, Planned; Repair Type = Annuloplasty, Sliding Plasty; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, Procedural Support.
-  Mitral Valve Procedure performed = Repair, Unplanned disease or anatomy; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, CPB Wean.

Question 6 Results



Answer Choices	Responses
Mitral Valve Procedure performed = Repair , Planned; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, CPB Wean.	5.13% 2
Mitral Valve Procedure performed = Repair , Unplanned disease or anatomy; Repair Type = Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, Hemodynamic Instability.	7.69% 3
Mitral Valve Procedure performed = Repair , Planned; Repair Type = Annuloplasty, Sliding Plasty; Other Cardiac Procedure = AFib Intracardiac lesion; IABP = Intraoperative, Procedural Support.	0.00% 0
Mitral Valve Procedure performed = Repair , Unplanned disease or anatomy; Repair Type = Annuloplasty, Leaflet Resection; Other Cardiac Procedure = Atrial Appendage procedure, LAA; IABP = Intraoperative, CPB Wean.	87.18% 34
Total	39

Question 7 Selections:

- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Late; Encephalopathy = Yes.
- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Acute; Atrial Fib = Yes.
- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of injury; Re-Op for Bleeding/Tamponade = Yes, Late; Pleural Effusion requiring drainage = Yes.
- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of injury; Re-Op for Bleeding/Tamponade = Yes, Acute; Prolonged Ventilation = Yes.

Data Points to Consider

- What were the Postoperative Echo results?
- What is an Imaging Study?
- What Postoperative Events did the patient encounter?

Postoperative Echo

Post Op Echo Performed to evaluate valve(s):	<input type="checkbox"/> Yes	<input type="checkbox"/> No (If Yes ↓)	POpTTEch (4625)				
Highest level aortic insufficiency found:	<input type="checkbox"/> None	<input type="checkbox"/> Trace/trivial	<input type="checkbox"/> Mild	<input type="checkbox"/> Moderate	<input type="checkbox"/> Severe	<input type="checkbox"/> Not Reported	POpTTAR (4630)
Highest level mitral insufficiency found:	<input type="checkbox"/> None	<input type="checkbox"/> Trace/trivial	<input type="checkbox"/> Mild	<input type="checkbox"/> Moderate	<input type="checkbox"/> Severe	<input type="checkbox"/> Not Reported	POpTTMR (4635)
Highest level tricuspid insufficiency found:	<input type="checkbox"/> None	<input type="checkbox"/> Trace/trivial	<input type="checkbox"/> Mild	<input type="checkbox"/> Moderate	<input type="checkbox"/> Severe	<input type="checkbox"/> Not Reported	POpTTTR (4640)
Highest level pulmonic insufficiency found:	<input type="checkbox"/> None	<input type="checkbox"/> Trace/trivial	<input type="checkbox"/> Mild	<input type="checkbox"/> Moderate	<input type="checkbox"/> Severe	<input type="checkbox"/> Not Reported	POpTTPu (4645)

Seq. #: 4625

Long Name: Postop Echo; Short Name: POpTTEch

Definition: Indicate whether an echo was performed postoperatively to evaluate valvular function prior to discharge.

- Capture echos performed after the patient leaves the operating room but prior to hospital discharge. - Code the exam **closest** to discharge.
- Indicate the highest/worst level found.
- Mild-to-moderate should be coded as moderate; moderate to severe should be coded as severe.
- If the report for an echo does not address valve disease, code "not reported".
- Use the following to categorize the level of insufficiency/regurgitation:
 - None = 0
 - Trace/trivial = 1+
 - Mild = 2+
 - Moderate = 3+
 - Severe = 4+

"Not Reported" = "Not Documented"

Mitral Valve
Mitral Insufficiency: VDInsufM (1680) <input type="checkbox"/> None <input type="checkbox"/> Trivial/Trace <input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe <input type="checkbox"/> Not Documented

Imaging Study

Imaging Study for Myocardial Injury : ~~POplmagStdy (4685)~~ *...for any reason*

- Not performed
- Angiographic evidence of new thrombosis or occlusion of graft or native coronary
- Imaging evidence of new loss of viable myocardium
- No evidence of new myocardial injury
- Other

Seq. #: 4685

Long Name: Postop Imaging Study; **Short Name:** POplmagStdy

Definition: Indicate the post procedure imaging study findings, if performed.

- This does not imply that post op imaging is expected to be performed on all patients; the intent is to *capture results if an exam was performed....for any reason.*
- Studies may include *echo, cardiac cath, CT, MRI (does not include CXR).*
- If more than one study is done following surgery, *capture the last study done prior to discharge.*

Postoperative Events

P. Postoperative Events

Surgical Site Infection within 30 days of operation: Yes No (If Yes ↓) **SurSInf (4690)**

Sternal Superficial Wound Infection: Yes, within 30 days of procedure Yes, >30 days after procedure but during hosp. for surgery No
CSternalSupInf (4695)

Deep Sternal Infection/ Mediastinitis: **DeepStemInf (4700)**

Yes, within 30 days of procedure Yes, >30 days after procedure but during hosp. for surgery No

(If either Yes value →) Diagnosis Date: ___/___/___ (mm/dd/yyyy) **DeepStemInfDt (4705)**

Thoracotomy: Yes, within 30 days of procedure Yes, >30 days after procedure but during hosp. for surgery No **CTThor (4710)**

Conduit Harvest: Yes, within 30 days of procedure Yes, >30 days after procedure but during hosp. for surgery No **ConduitHarv (4715)**

Cannulation Site: Yes, within 30 days of procedure Yes, >30 days after procedure but during hosp. for surgery No **CanSite (4720)**

Wound Intervention/Procedure: Yes No (If Yes ↓) **WoundInter (4725)**

Wound Intervention – Open with Packing/Irrigation: Yes, primary incision Yes, secondary incision Both No
WoundIntOpen (4730)

Wound Intervention – Wound Vac: **WoundIntVac (4735)** Yes, primary incision Yes, secondary incision Both No

Secondary Procedure Muscle Flap: **WoundIntMuscle (4740)** Yes, primary incision Yes, secondary incision Both No

Secondary Procedure Omental Flap: **WoundIntOmental (4745)** Yes No

Other In Hospital Postoperative Event Occurred: Yes No (If Yes ↓) **Complics (4750)**

Operative

ReOp for Bleeding /Tamponade: Yes No **COpReBld (4755)** (If Yes →) Bleed Timing: Acute Late **COpReBldTim (4760)**

ReOp for Valvular Dysfunction: Yes, surgical Yes, transcatheter No **COpReVlv (4765)**

ReOp for Graft Occlusion: Yes, surgical Yes, PCI No **COpReGft (4770)**

ReOp for Other Cardiac Reasons: Yes No **COpReOth (4775)**

ReOp for Other Non-Cardiac Reasons: Yes No **COpReNon (4780)**

Open chest with planned delayed sternal closure: Yes No **COpPlndDelay (4785)**


Sternotomy Issue: Yes No **CSternal (4790)** (If Yes →) Sternal instability/dehiscence (sterile): Yes No **CSternalDehis (4795)**

Infection

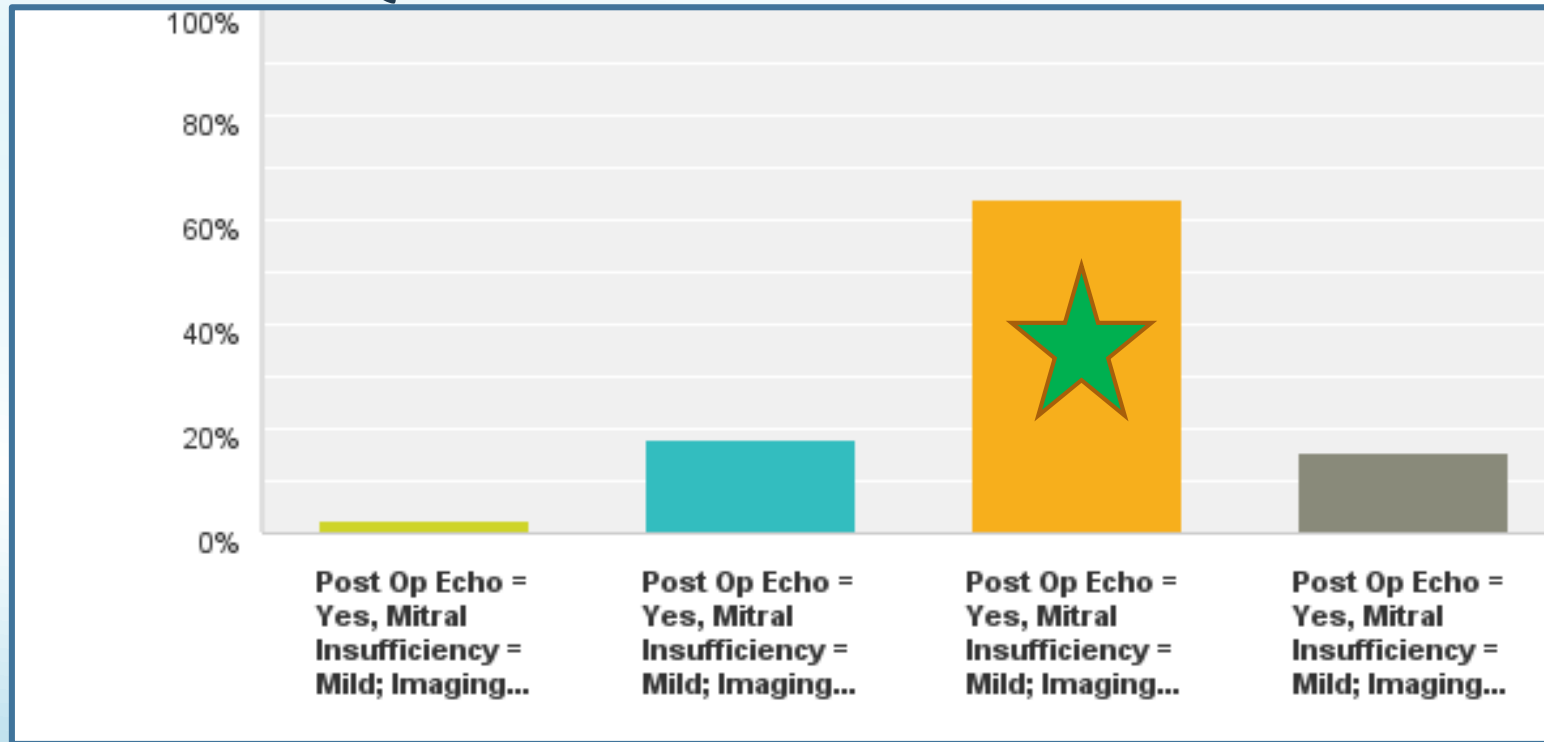
Sepsis: Yes No **CSepsis (4800)** (If Yes →) Positive Blood Cultures: Yes No **CSepsisPBC (4805)**

Neurologic	
<input checked="" type="checkbox"/>	Postoperative Stroke: <input type="checkbox"/> Yes, hemorrhagic <input type="checkbox"/> Yes, embolic <input type="checkbox"/> Yes, undetermined type <input type="checkbox"/> No CNStrokP (4810)
<input checked="" type="checkbox"/>	Transient Ischemic Attack (TIA): <input type="checkbox"/> Yes <input type="checkbox"/> No CNStrokTTIA (4815)
<input checked="" type="checkbox"/>	Encephalopathy: <input type="checkbox"/> None <input type="checkbox"/> Anoxic <input type="checkbox"/> Embolic <input type="checkbox"/> Drug <input type="checkbox"/> Metabolic <input type="checkbox"/> Intracranial Bleeding <input type="checkbox"/> Other <input type="checkbox"/> Unknown CNComaEnceph (4820)
<input checked="" type="checkbox"/>	Paralysis: <input type="checkbox"/> Yes <input type="checkbox"/> No CNParal (4825) (If Yes →) Paralysis Type: <input type="checkbox"/> Transient <input type="checkbox"/> Permanent CNParalTy (4830)
Pulmonary	
<input checked="" type="checkbox"/>	Prolonged Ventilation: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No (OR exit time until initial extubation, plus any additional reintubation hours) CPVntLng (4835)
<input checked="" type="checkbox"/>	Pneumonia: <input type="checkbox"/> Yes <input type="checkbox"/> No CPPneum (4840)
<input checked="" type="checkbox"/>	Venous Thromboembolism – VTE: <input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes ↓) CVTE (4845)
<input checked="" type="checkbox"/>	Pulmonary Thromboembolism: <input type="checkbox"/> Yes <input type="checkbox"/> No PulmEmb (4850)
<input checked="" type="checkbox"/>	Deep Venous Thrombosis: <input type="checkbox"/> Yes <input type="checkbox"/> No DVT (4855)
<input checked="" type="checkbox"/>	Pleural Effusion Requiring Drainage: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No CPIEff (4860)
<input checked="" type="checkbox"/>	Pneumothorax Requiring Intervention: <input type="checkbox"/> Yes <input type="checkbox"/> No PostOpPneumo (4865)
Renal	
<input checked="" type="checkbox"/>	Renal Failure: <input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes ↓) CRenFail (4870)
<input checked="" type="checkbox"/>	Dialysis (Newly Required): <input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes →) Required after Hospital Discharge: <input type="checkbox"/> Yes <input type="checkbox"/> No CRenDial (4875) DialDur (4880)
<input checked="" type="checkbox"/>	Ultra Filtration Required: <input type="checkbox"/> Yes <input type="checkbox"/> No CUltraFil (4885)
Vascular	
<input checked="" type="checkbox"/>	Iliac/Femoral Dissection: <input type="checkbox"/> Yes <input type="checkbox"/> No CVaIlFem (4890)
<input checked="" type="checkbox"/>	Acute Limb Ischemia: <input type="checkbox"/> Yes <input type="checkbox"/> No CVaLbIs (4895)
Other	
<input checked="" type="checkbox"/>	Rhythm Disturbance Requiring Permanent Device: <input type="checkbox"/> Pacemaker <input type="checkbox"/> ICD <input type="checkbox"/> Pacemaker/ICD <input type="checkbox"/> Other <input type="checkbox"/> None CRhythmDis (4900)
<input checked="" type="checkbox"/>	Cardiac Arrest: <input type="checkbox"/> Yes <input type="checkbox"/> No COtArrst (4905)
<input checked="" type="checkbox"/>	Anticoagulant Event: <input type="checkbox"/> Yes <input type="checkbox"/> No COtCoag (4910)
<input checked="" type="checkbox"/>	Tamponade (Non-Surgical Intervention): <input type="checkbox"/> Yes <input type="checkbox"/> No COtTamp (4915)
<input checked="" type="checkbox"/>	Gastro-Intestinal Event: <input type="checkbox"/> Yes <input type="checkbox"/> No COtGI (4920)
<input checked="" type="checkbox"/>	Multi-System Failure: <input type="checkbox"/> Yes <input type="checkbox"/> No COtMSF (4925)
<input checked="" type="checkbox"/>	Atrial Fibrillation: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No COtAFib (4930)
<input checked="" type="checkbox"/>	Aortic Dissection: <input type="checkbox"/> Yes <input type="checkbox"/> No CVaAoDis (4935)
<input checked="" type="checkbox"/>	Recurrent Laryngeal Nerve Injury: <input type="checkbox"/> Yes <input type="checkbox"/> No RecLarynNrvInj (4940)
<input checked="" type="checkbox"/>	Phrenic Nerve Injury: <input type="checkbox"/> Yes <input type="checkbox"/> No PhrenNrvInj (4945)
<input checked="" type="checkbox"/>	Other: <input type="checkbox"/> Yes <input type="checkbox"/> No COtOther (4950)

Question 7 Selections:

- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Late; Encephalopathy = Yes.
- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Acute; Atrial Fib = Yes.
-  Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of injury; Re-Op for Bleeding/Tamponade = Yes, Late; Pleural Effusion requiring drainage = Yes.
- Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of injury; Re-Op for Bleeding/Tamponade = Yes, Acute; Prolonged Ventilation = Yes.

Question 7 Results



Answer Choices	Responses
Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Late; Encephalopathy = Yes.	2.56% 1
Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = Not performed; Re-Op for Bleeding/Tamponade = Yes, Acute; Atrial Fib = Yes.	17.95% 7
Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of new injury; Re-Op for Bleeding/Tamponade = Yes, Late; Pleural Effusion requiring drainage = Yes.	64.10% 25
Post Op Echo = Yes, Mitral Insufficiency = Mild; Imaging Study = No evidence of new injury; Re-Op for Bleeding/Tamponade = Yes, Acute; Prolonged Ventilation = Yes.	15.38% 6
Total	39

Question 8 Selections:

- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.
- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = No.
- Mortality = Yes; Operative Death = No, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.
- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.

Data Points to Consider

- Was this mortality an Operative Death?
- Was this patient considered a “Readmission”?

Mortality and Operative Death

Seq. #: 5005

Long Name: Mort-Mortality; **Short Name:** Mortalty

Definition: Indicate whether the patient has been declared dead **within this hospitalization or any time after discharge** from this hospitalization. This includes all causes of death, including those causes clearly unrelated to the operation. . This could be while the patient is in the hospital for the current procedure, within 30 days of the procedure, or **"long term"**, meaning **six months, five years, or anytime in the future.**

Seq. #: 5025

Long Name: Mort-Op Death; **Short Name:** MtOpD

Definition: Operative Mortality includes: (1) **ALL deaths, regardless of cause, occurring during the hospitalization in which the operation was performed, even if after 30 days (including patients transferred to other acute care facilities);** and (2) **ALL deaths, regardless of cause, occurring after discharge from the hospital, but before the end of the thirtieth postoperative day.**

Q. Mortality		
Mortality <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No Mortality (5005)	Discharge Status: <input checked="" type="checkbox"/> Alive <input type="checkbox"/> Dead MtDCStat (5010)	Status at 30 days After Surgery: <input type="checkbox"/> Alive <input checked="" type="checkbox"/> Dead <input type="checkbox"/> Unknown Mt30Stat (5015)
Primary method used to verify 30-day status: Mt30StatMeth (5020)		
<input type="checkbox"/> Phone call to patient or family	<input type="checkbox"/> Medical record	<input type="checkbox"/> Social Security Death Master File /NDI
<input type="checkbox"/> Letter from medical provider	<input type="checkbox"/> Office visit >= 30 days after procedure	<input type="checkbox"/> Other
(If Mortality = Yes ↓)		
Operative Death: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No MtOpD (5025)	Mortality - Date ___/___/___ (mm/dd/yyyy)	MtDate (5030)
Location of Death: <input checked="" type="checkbox"/> OR During Initial Surgery <input checked="" type="checkbox"/> Hospital (Other than OR)	<input type="checkbox"/> Home	<input type="checkbox"/> Extended Care Facility
MtLocatn (5035)	<input type="checkbox"/> Hospice <input type="checkbox"/> Acute Rehabilitation	<input type="checkbox"/> OR During Reoperation <input type="checkbox"/> Unknown <input type="checkbox"/> Other
Primary Cause of Death (select only one) MtCause (5040)		
<input checked="" type="checkbox"/> Cardiac	<input type="checkbox"/> Neurologic	<input type="checkbox"/> Renal <input type="checkbox"/> Vascular <input type="checkbox"/> Infection <input type="checkbox"/> Pulmonary <input type="checkbox"/> Unknown <input type="checkbox"/> Other

Readmission

Seq. #: 5140

Long Name: Readmission; **Short Name:** Readmit

Definition: Indicate whether the patient was readmitted to the hospital within 30 days of discharge from hospitalization for this surgery. Code yes for inpatient admission to an acute care facility. **Do not capture ED or outpatient visits or admission to a skilled facility or nursing home.**

- It is understood that some readmissions are planned; these are **still counted as readmissions**.
- Readmission **does not need to be at same institution** as surgical procedure. Obtain information as close to 30 days from date of discharge as possible.
- **Do not include Emergency Dept. visits or observation unless the ED visits lead to admission.** The intent is to capture inpatient readmissions to acute care and primary care institutions only.
- If a patient is readmitted to an inpatient rehabilitation hospital, code "No".
- To align with CMS, **30 day readmission should not be coded for patients who remain in observation units, no matter the duration.**
- On occasion a patient is readmitted twice within the 30 day time frame from the date of the procedure. Any time the patient is readmitted to a hospital \leq 30 days from the date of discharge regardless if the readmission was planned or unplanned, related or unrelated. You **code the first readmission only**.

S. Readmission

(If Discharge Status = Alive.)

Readmit : Yes No Unknown (If Yes ↓) **Readmit (5140)**

Readmit Date: ___/___/____ (mm/dd/yyyy) **ReadmitDt (5145)**

Readmit Primary Reason: **ReadmRsn (5160)**

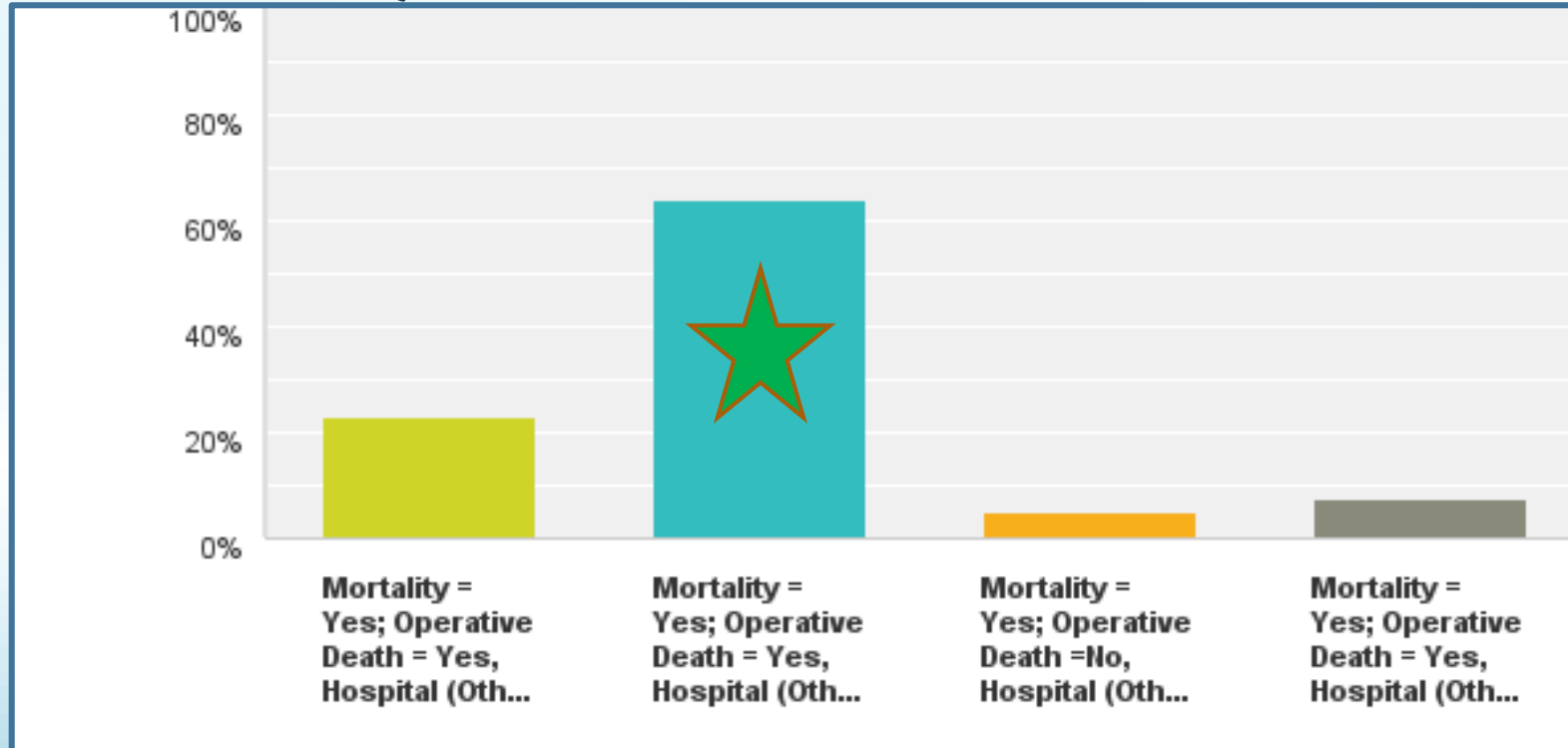
- | | |
|---|--|
| <input type="checkbox"/> Anticoagulation Complication - Pharmacological | <input type="checkbox"/> Pneumonia |
| <input type="checkbox"/> Anticoagulation Complication – Valvular | <input type="checkbox"/> Renal Failure |
| <input type="checkbox"/> Arrhythmia/Heart Block | <input type="checkbox"/> Respiratory complication, Other |
| <input type="checkbox"/> Congestive Heart Failure | <input type="checkbox"/> Stroke |
| <input type="checkbox"/> Coronary Artery/Graft Dysfunction | <input type="checkbox"/> TIA |
| <input type="checkbox"/> DVT | <input type="checkbox"/> Transplant Rejection |
| <input type="checkbox"/> Endocarditis | <input type="checkbox"/> VAD Complication |
| <input type="checkbox"/> Infection, Conduit Harvest Site | <input type="checkbox"/> Valve Dysfunction |
| <input type="checkbox"/> Infection, Deep Sternum / Mediastinitis | <input type="checkbox"/> Vascular Complication, acute |
| <input type="checkbox"/> Myocardial Infarction and/or Recurrent Angina | <input type="checkbox"/> Other – Related Readmission |
| <input type="checkbox"/> PE | <input type="checkbox"/> Other – Nonrelated Readmission |
| <input type="checkbox"/> Pericardial Effusion and/or Tamponade | <input type="checkbox"/> Other – Planned Readmission |
| <input type="checkbox"/> Pleural effusion requiring intervention | <input type="checkbox"/> Unknown |

Question 8 Selections:

- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.
- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = No.
- Mortality = Yes; Operative Death = No, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.
- Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.



Question 8 Results



Answer Choices	Responses
Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.	23.08% 9
Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = No.	64.10% 25
Mortality = Yes; Operative Death = No, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.	5.13% 2
Mortality = Yes; Operative Death = Yes, Hospital (Other than the OR); Discharge Location = Extended/Transitional Care; Readmission = Yes.	7.69% 3
Total	39



Thanks!

Any Question?