



# **MSTCVS Quality Collaborative Summer Meeting 2017 Traverse City, MI**

Survey  
Monkey

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These slides are intended for use by MSTCVS surgeon champions and data managers for the development and evaluation of quality improvement plans.

Each slide includes the MSTCVS confidentiality statement.

# Scenario #1

A 76 yr. male with a 18 month medical history of A-Fib, rate controlled with beta blockade and Xarelto, presents to the ER at 23:00 on Sunday evening complaining of chest pain. His wife states the pain had started earlier that evening after a long day of yard work. Suspecting muscle pain, and having nothing else in the house, she made him eat four 80 mg. aspirin. The pain subsided somewhat, but then he began to complain of shortness of breath, and noticed that his heart was "fluttering like before". In the ER, sublingual NTG relieved the patient's chest pain, while a Heparin drip was started and his Xarelto was held.

Upon exam, the EKG confirmed A-Fib, and ruled out ST changes. TTE reported "inconclusive results", but suggested moderate MR with bi-leaflet thickening and an EF calculated at 45%. X-ray identified small bilateral pleural effusions with a slightly enlarged cardiac silhouette. Rales and mild pedal edema were also noted, prompting diuretic therapy. Following the diagnosis of heart failure, the patient was admitted for cardiac evaluation.

A cardiac cath performed Monday afternoon revealed triple vessel disease, mild MR and an estimated EF of 50%. After a discussion of the results and the surgical risks involved, the patient is scheduled for CAB, with possible MV Repair, and possible Maze procedure for Wednesday afternoon.

Upon entry to the OR, the TEE reveals the heart to be in NSR with mild MR, minimal anterior leaflet prolapse, and a measured EF of 48%. Based on these findings, the patient undergoes CAB x 4, an epicardial Maze and LAA exclusion via Atriclip.

Post-operatively, the patient reverts back to A-Fib, but converts to NSR with resumption of his home medications, and is discharged on POD #6 on ASA, Statin, BB, and Xarelto. Unfortunately, he is re-admitted on POD #33 with recurrent A-Fib. Following cardioversion and medication adjustment, he is discharged 4 days later.



## #1: Does this patient suffer from Heart Failure (#911) in version 2.9?

### Choice of Answers:

- No.
- Yes. Timing is Chronic. Type is Both.
- Yes. Timing is Acute. Type is Unavailable.
- Yes. Timing is Both. Type is Both.

### Points to Consider:

- Re-visit Heart Failure definition.
- Heart Failure Timing (re-design).
- Heart Failure Type (new).





**SEQ. #:** 911

**Long Name:** Heart Failure

**Short Name:** HeartFail

**Definition:** Indicate whether there is physician documentation or report that the patient has been in a state of heart failure.

**Intent/Clarification:**

Heart failure is 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.

**F. Preoperative Cardiac Status**

Prior Myocardial Infarction:  Yes  No  Unknown (If Yes ↓)

PrevMI (885)

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		

Heart Failure:  Yes  No  Unknown (If Yes →)

HeartFail (911)

Timing:  Acute  Chronic  Both

HeartFailTmg (912)

Type:  Systolic  Diastolic  Both  Unavailable

HeartFailType (913)

Classification-NYHA:  Class I  Class II  Class III  Class IV  Not Documented

ClassNYH (915)

**SEQ. #:** 912

**Long Name:** Heart Failure Timing

**Short Name:** HeartFailTmg

**Definition:** Indicate whether heart failure is acute, chronic or both (acute on chronic)

**Intent/Clarification:**

- Acute heart failure is the rapid onset of symptoms and signs of heart failure and may occur with or without previous cardiac disease. Acute decompensated heart failure is a sudden worsening of the signs and symptoms of heart failure, which typically includes difficulty breathing (dyspnea), leg or feet swelling, and fatigue.
- Chronic heart failure develops gradually over time with symptoms of shortness of breath, lower extremity swelling and fatigue without an acute exacerbation.
- Both involves patients with chronic heart failure who presents with acute symptoms.

**SEQ. #:** 913

**Long Name:** Heart Failure Type

**Short Name:** HeartFailType

**Definition:** Indicate the type of heart failure.

**Intent/Clarification:**

- Systolic: The left ventricle lacks the force to push enough blood into the circulation.
- Diastolic: The left ventricle is stiff and fails to relax sufficiently to allow adequate filling.
- Both: Components of both systolic and diastolic failure exist.
- Unavailable: The type of heart failure is not documented in the medical record.



**SEQ. #: 911**

**Long Name:** Heart Failure

**Short Name:** HeartFail

**Definition:** Indicate whether there is physician documentation or report that the patient has been in a state of heart failure.

**Intent/Clarification:**

Heart failure is 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.

**F. Preoperative Cardiac Status**

Prior Myocardial Infarction:  Yes  No  Unknown (If Yes ↓)

PrevMI (885)

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		

Heart Failure  Yes  No  Unknown (If Yes →)

HeartFail (911)

Timing:  Acute  Chronic  Both

HeartFailTmg (912)

Type:  Systolic  Diastolic  Both  Unavailable

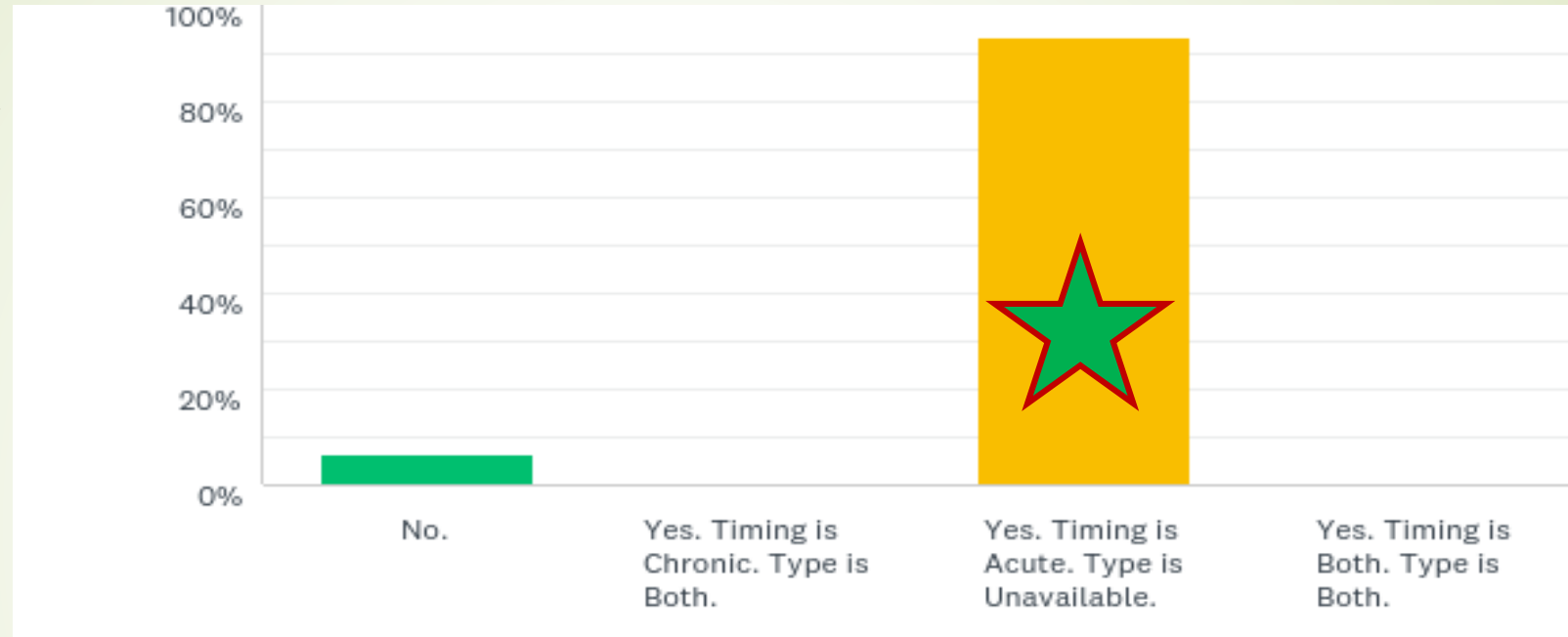
HeartFailType (913)

Classification-NYHA:  Class I  Class II  Class III  Class IV  Not Documented

ClassNYH (915)



# #1: Does this patient suffer from Heart Failure (#911) in version 2.9?



Answer Choices
No.
Yes. Timing is Chronic. Type is Both.
Yes. Timing is Acute. Type is Unavailable.
Yes. Timing is Both. Type is Both.

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## #2: What Type of Atrial Fibrillation (#962) does this patient suffer from?

### Choice of Answers:

- Paroxysmal
- Persistent
- Longstanding Persistent
- Permanent

**SEQ. #:** 962

**Long Name:** Cardiac Arrhythmia - Atrial Fibrillation - Type

**Short Name:** ArrhythAFib

**Definition:** Indicate whether arrhythmia was atrial fibrillation and if so, which type.

### Intent/Clarification:

If the diagnosis of atrial fibrillation is present code the type:

- Paroxysmal: Recurrent AF (> 2 episodes). Terminates spontaneously within 7 days.
- Persistent: Sustained episode > 7 days, or lasting < 7 days, but necessitating pharmacologic or electrical cardioversion.
- Long-Standing Persistent: Continuous episode of > 1 year duration.
- Permanent: Continuous episode of > 1 year duration.

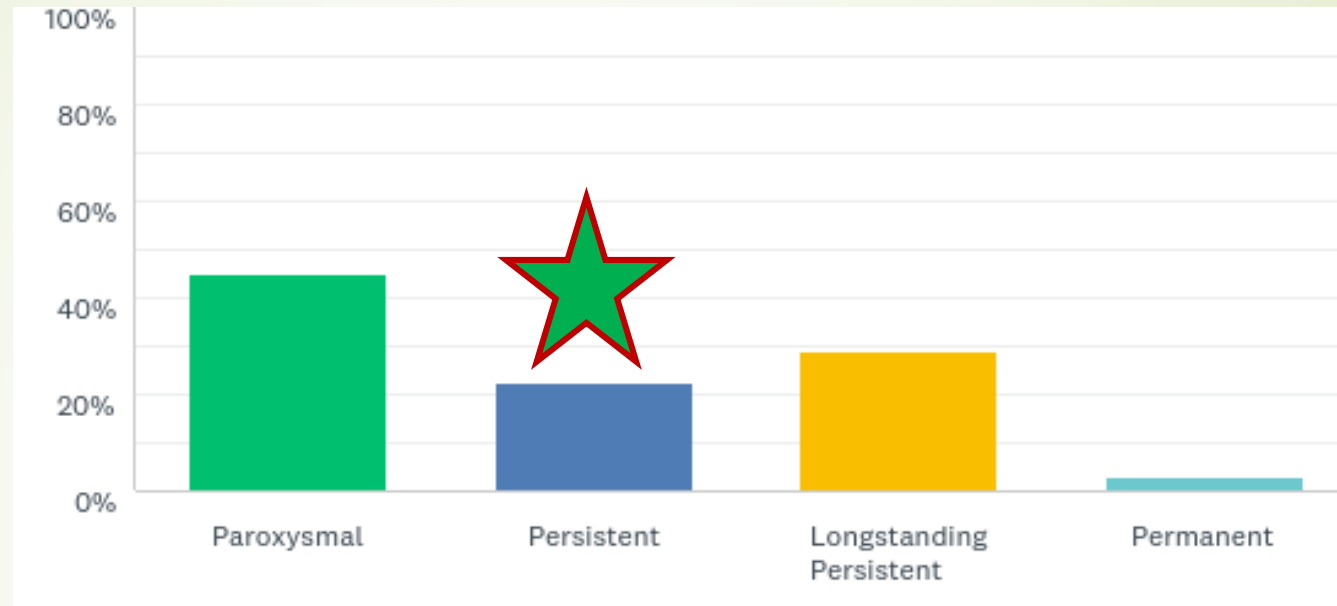
### Ver. 2.8

Paroxysmal	Recurrent AF (> 2 episodes). Terminates spontaneously within 7 days
Continuous/Persistent	Sustained episode > 7 days, or lasting < 7 days, but necessitating pharmacologic or electrical cardioversion
Long-Standing Persistent	Continuous episode of > 1 year duration
Permanent	AF at a point in which no further treatment of any kind is considered.

F. Preoperative Cardiac Status						
Prior Myocardial Infarction: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes ↓) PrevMI (885)						
MI When: <input type="checkbox"/> ≤6 Hrs. <input type="checkbox"/> >6 Hrs. but <24 Hrs. <input type="checkbox"/> 1 to 7 Days <input type="checkbox"/> 8 to 21 Days <input type="checkbox"/> >21 Days MIWhen (890)						
Cardiac Presentation/Symptoms: (Choose <u>one</u> 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						
Heart Failure: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes →)    Timing: <input type="checkbox"/> Acute <input type="checkbox"/> Chronic <input type="checkbox"/> Both    Type: <input type="checkbox"/> Systolic <input type="checkbox"/> Diastolic <input type="checkbox"/> Both <input type="checkbox"/> Unavailable HeartFail (911)    HeartFailTmg (912)    HeartFailType (913)						
Classification-NYHA: <input type="checkbox"/> Class I <input type="checkbox"/> Class II <input type="checkbox"/> Class III <input type="checkbox"/> Class IV <input type="checkbox"/> Not Documented ClassNYH (915)						
Cardiogenic Shock : <input type="checkbox"/> Yes, at the time of the procedure <input type="checkbox"/> Yes, not at the time of the procedure but within prior 24 hours <input type="checkbox"/> No CarShock (930)						
Resuscitation: <input type="checkbox"/> Yes - Within 1 hour of the start of the procedure <input type="checkbox"/> Yes - More than 1 hour but less than 24 hours of the start of the procedure <input type="checkbox"/> No Resusc (935)						
Arrhythmia: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No Arrhythmia (945)						
Permanently Paced Rhythm: <input type="checkbox"/> Yes <input type="checkbox"/> No ArrhythPPaced (947)						
(If Arrhythmia = Yes →)						
(If Yes , choose one response below for each rhythm →)	VTach/VFib ArrhythVV (950)	Sick Sinus Syndrome ArrhythSSS (955)	AFlutter ArrhythAFlutter (960)	AFibrillation ArrhythAtrFib (961)	Second Degree Heart Block ArrhythSecond (965)	Third Degree Heart Block ArrhythThird (970)
None						
Remote (> 30 days preop)				<input checked="" type="checkbox"/>		
Recent (≤= 30 days preop)						
(If AFibrillation not 'None' →)	Atrial Fibrillation Type: <input type="checkbox"/> Paroxysmal <input checked="" type="checkbox"/> Persistent <input type="checkbox"/> Longstanding Persistent <input type="checkbox"/> Permanent ArrhythAFib (962)					

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## #2: What Type of Atrial Fibrillation (#962) does this patient suffer from?



### Answer Choices

Paroxysmal

Persistent

Longstanding Persistent

Permanent

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### #3: How would you abstract this patient's use of ASA (#1070)?

#### Choice of Answers:

- ASA is Yes; Discontinuation is 3 days; One Time Dose is No.
- ASA is Yes; Discontinuation is 3 days; One Time Dose is Yes.

G. Preoperative Medications		
Medication	Timeframe	Administration
ACE or ARB <b>MedACEI48 (1020)</b>	Within 48 hours	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
Amiodarone <b>MedAmiodarone (1025)</b>	Prior to surgery	<input type="checkbox"/> Yes, on home therapy <input type="checkbox"/> Yes, therapy started this admission <input type="checkbox"/> No <input type="checkbox"/> Unknown
Antianginal	Beta Blocker <b>MedBeta (1030)</b>	Within 24 hours <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated
	Beta Blocker <b>MedBetaTher (1035)</b>	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Calcium Channel Blocker <b>MedCCChanTher (1040)</b>	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Long-acting Nitrate <b>MedLongActNit (1045)</b>	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Nitrates, intravenous <b>MedNitIV (1050)</b>	Within 24 hours <input type="checkbox"/> Yes <input type="checkbox"/> No
	Other Antianginal <b>MedOthAntiang (1055)</b>	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
Antiplatelet	ADP Inhibitor (includes P2Y12) <b>MedADP5Days (1060)</b>	Within 5 days <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown (If Yes→)ADP Inhibitors Discontinuation: _____ (# days prior to surgery) <b>MedADPIDis (1065)</b>
	Aspirin <b>MedASA (1070)</b>	Within 5 days <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
		(If Yes→) Aspirin Discontinuation: _____ (# days prior to surgery) <b>MedASADis (1071)</b> Aspirin one time dose: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>MedASAOnce (1072)</b>

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**SEQ. #:** 1070

**Long Name:** Meds-Aspirin Within Five Days

**Short Name:** MedASA

**Definition:** Indicate whether or not the patient received Aspirin or Ecotrin within 5 days preceding surgery.

**Intent/Clarification:**

Anti-inflammatory, analgesic and antiplatelet action. Half-life of aspirin products is 5-7 days. Aspirin use may predispose patient to post op bleeding.

- **Yes** - Capture those who are prescribed to take Aspirin or Ecotrin on a regular schedule and are presumed to be at a therapeutic level, 5 days preceding surgery (entry into the OR) - The minimum dose should be at least 75 mg (i.e. Aggrenox, which is only 25mg, should not be included). Do Not Include a one-time dose.
- **No** – Patient did not receive Aspirin within 5 days preceding surgery.
- **Contraindicated** - Documented evidence of contraindication: If a contraindication

**SEQ. #:** 1071

**Long Name:** Meds-Aspirin Discontinuation

**Short Name:** MedASADis

**Definition:** Indicate the number of days prior to surgery Aspirin use was discontinued. If less than 24 hours, enter "0".

**Intent/Clarification:** -

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**SEQ. #:** 1072

**Long Name:** Meds-Aspirin One-Time Dose

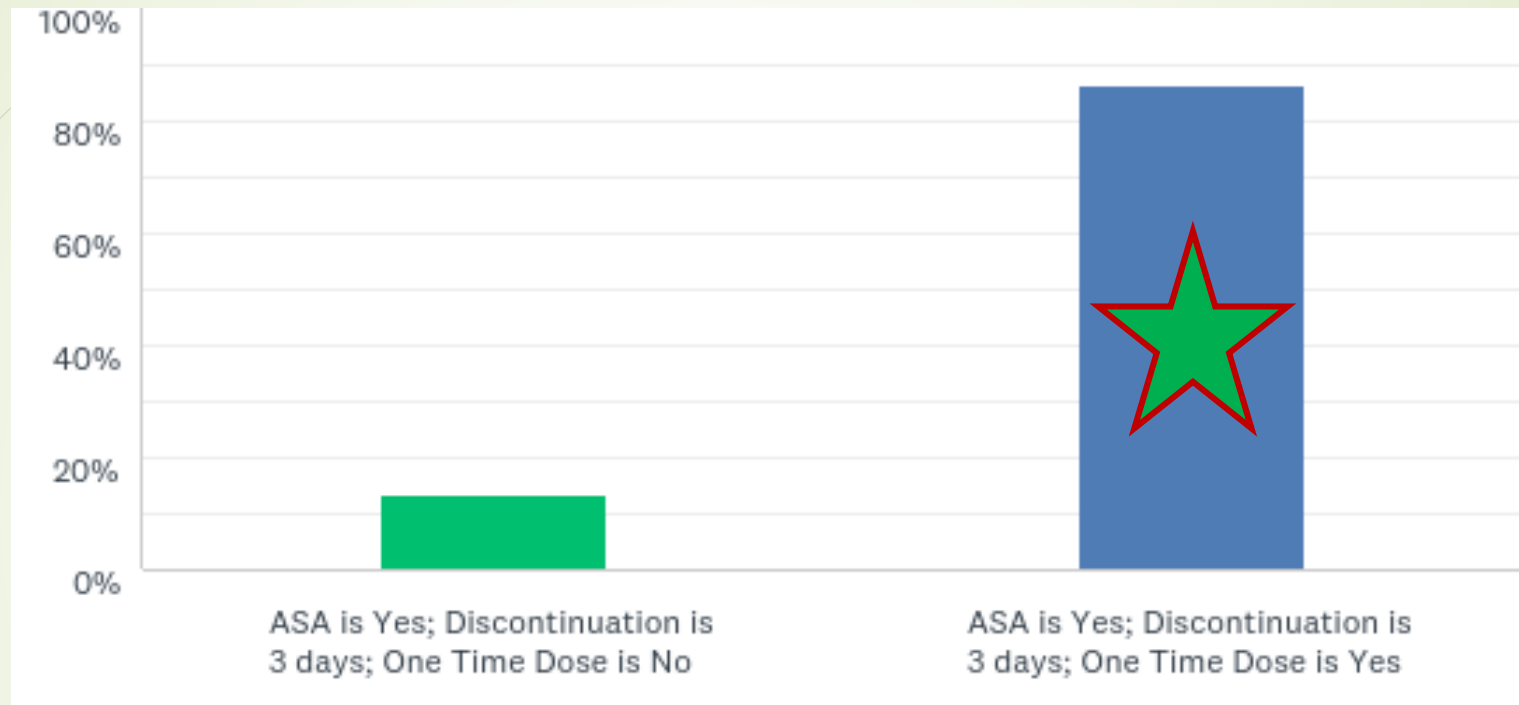
**Short Name:** MedASAOnce

**Definition:** Indicate whether the patient received a one-time dose of Aspirin and is not on daily aspirin.

G. Preoperative Medications		
Medication	Timeframe	Administration
ACE or ARB MedACEI48 (1020)	Within 48 hours	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
Amiodarone MedAmiodarone (1025)	Prior to surgery	<input type="checkbox"/> Yes, on home therapy <input type="checkbox"/> Yes, therapy started this admission <input type="checkbox"/> No <input type="checkbox"/> Unknown
Antianginal	Beta Blocker MedBeta (1030)	Within 24 hours <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated
	Beta Blocker MedBetaTher (1035)	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Calcium Channel Blocker MedCChanTher (1040)	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Long-acting Nitrate MedLongActNit (1045)	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
	Nitrates, intravenous MedNitIV (1050)	Within 24 hours <input type="checkbox"/> Yes <input type="checkbox"/> No
	Other Antianginal MedOthAntiang (1055)	On therapy for $\geq 2$ weeks prior to surgery <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
Antiplatelet	ADP Inhibitor (includes P2Y12) MedADP5Days (1060)	Within 5 days <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown (If Yes $\rightarrow$ )ADP Inhibitors Discontinuation: _____ (# days prior to surgery) MedADPIDis (1065)
	Aspirin MedASA (1070)	Within 5 days <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Contraindicated <input type="checkbox"/> Unknown
		(If Yes $\rightarrow$ ) Aspirin Discontinuation: <u>3</u> (# days prior to surgery) MedASADis (1071) Aspirin one time dose: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No MedASAOnce (1072)

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### #3: How would you abstract this patient's use of ASA (#1070)?



#### Answer Choices

ASA is Yes; Discontinuation is 3 days; One Time Dose is No

ASA is Yes; Discontinuation is 3 days; One Time Dose is Yes

# What's a NOAC?

**NOAC = Novel Oral AntiCoagulant**

- **NOAC = DOAC (Direct Oral AntiCoagulant)**
- **Developed and introduced to address several drawbacks and limitations associated with Warfarin (Coumadin) use.**
- **Indicated for prevention of stroke in patients with Non-Valvular AFib (NVAF).**
- **Do not require frequent monitoring and/or dose adjustment associated with Warfarin.**
- **All have short half-lives, BUT, with the exception of Dabigatran, reversal agents are currently not available.**
- **Currently, there are only four medications designated as NOAC's. A fifth was recently released, and more are under development.**



# Thrombin Inhibitors & Factor Xa Inhibitors

Within 5 Days of OR (Brand Names in Parentheses if Applicable)

## Direct Thrombin Inhibitors

Argatroban (**Acova, Argata, Novastan, Arganova, Exembol**)

Bivalirudin (**Angiomax**)

Desirudin (**Ipravask, Revasc**)

Hirudin

Lepirudin (**Refludan**)

Dabigatran (**Pradaxa**) (**NOAC**)

## Factor Xa Inhibitors

Apixaban (**Equilis**) (**NOAC**)

Betrixaban (**BEVYXXA**) **Newest NOAC (?)**

Edoxaban (**Lixiana, Savaysa**) (**NOAC**)

Fondaparinux (**Arixtra**)

Rivaroxaban (**Xarelto**) (**NOAC**)

Others are in development

**July 2017:** Not an Inclusive List/Use with Caution/ Work in Progress!

MSTCVS Quality Collaborative Data Managers

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## #4: How should we abstract this patient's pre-op use of Xarelto?

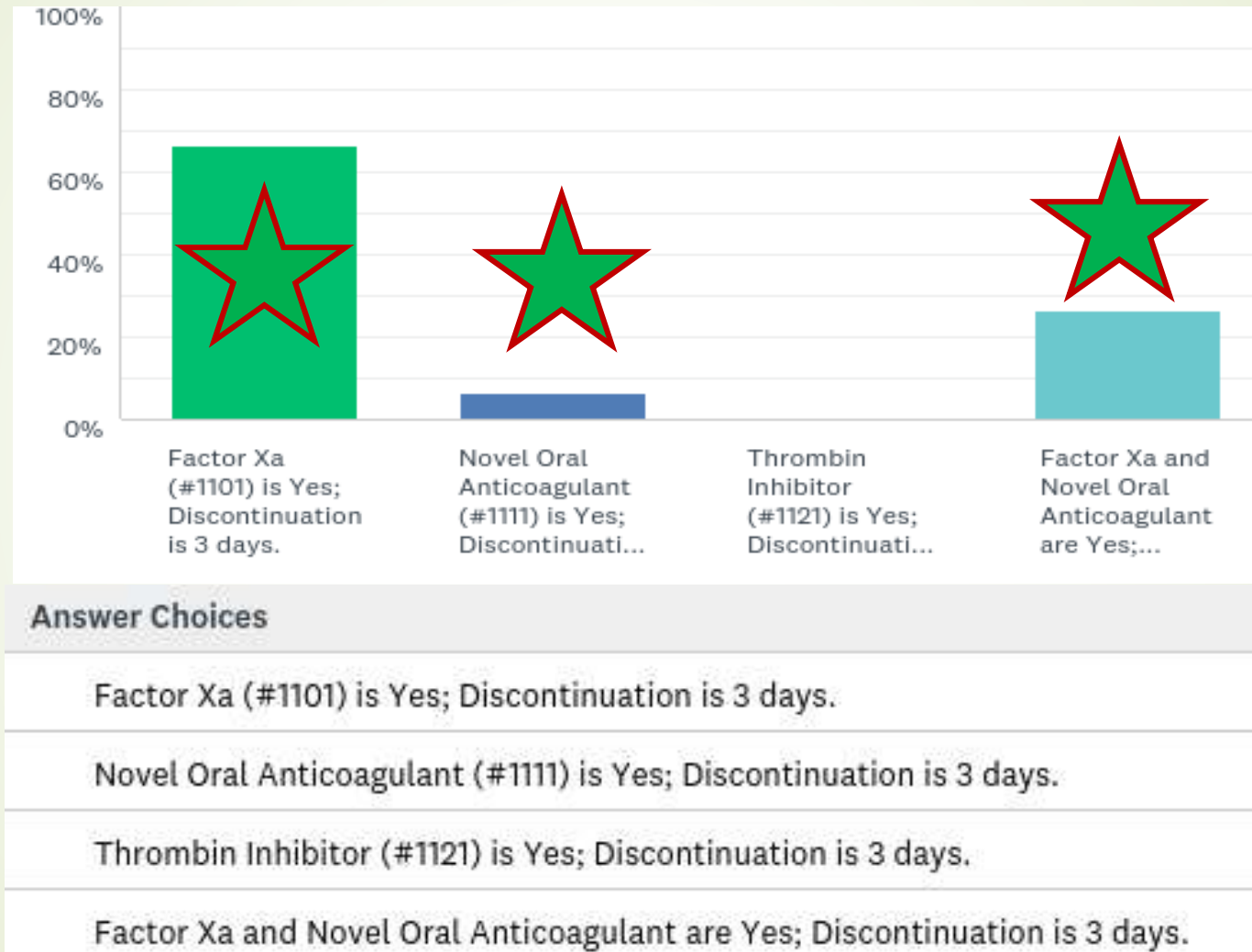
### Choice of Answers:

- Factor Xa (#1101) is Yes; Discontinuation is 3 days.
- Novel Oral Anticoagulant (#1111) is Yes; Discontinuation is 3 days.
- Thrombin Inhibitor (#1121) is Yes; Discontinuation is 3 days.
- Factor Xa and Novel Oral Anticoagulant are Yes; Discontinuation is 3 days.

	Glycoprotein IIb/IIIa MedGP (1073)	Within 24 hours	<input type="checkbox"/> Yes <input type="checkbox"/> No
Anticoagulant	Anticoagulants (Intravenous/ SubQ) MedACoag (1075)	Within 48 hours	<input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes→) Medication: <input type="checkbox"/> Heparin (Unfractionated) <input type="checkbox"/> Heparin (Low Molecular) <input type="checkbox"/> Both <input type="checkbox"/> Other MedACMN (1080)
	Warfarin (Coumadin) MedCouv5Days (1091)	Within 5 days	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes→) Coumadin Discontinuation: _____ (# days prior to surgery) MedCouv5Dis (1092)
	Factor Xa inhibitors MedXa5Days (1101)	Within 5 days	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes→) Factor Xa Discontinuation: _____ (# days prior to surgery) MedXa5DDis (1102)
	Novel Oral Anticoagulant MedNOAC5Days (1111)	Within 5 days	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes→) NOAC Discontinuation: _____ (# days prior to surgery) MedNOACDisc (1112)
	Thrombin Inhibitors MedThromIn5Days (1121)	Within 5 days	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes→) Thrombin Inhibitor Discontinuation: _____ (# days prior to surgery) MedThromInDisc (1122)
	Thrombolytics MedThrom (1125)	Within 48 hours	<input type="checkbox"/> Yes <input type="checkbox"/> No

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## #4: How should you abstract this patient's pre-op use of Xarelto?



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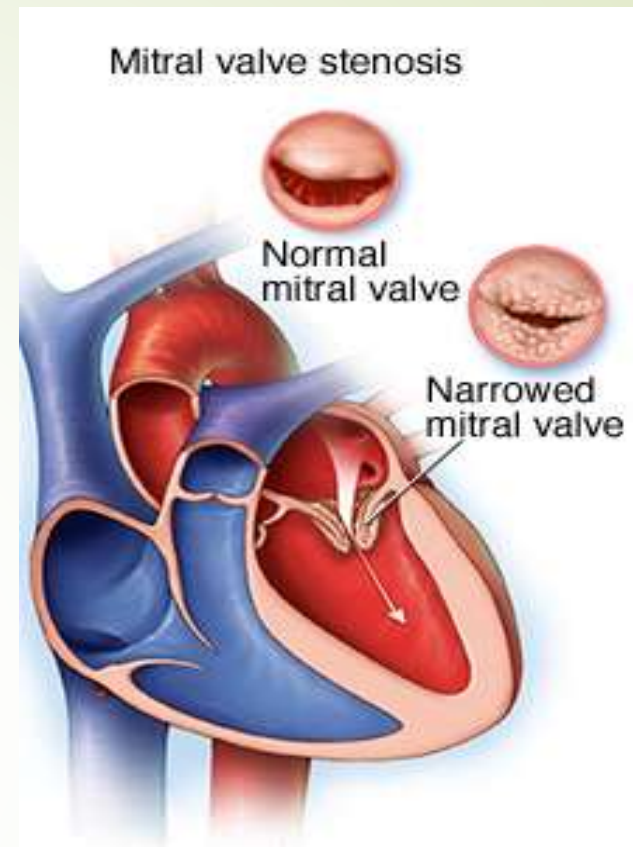
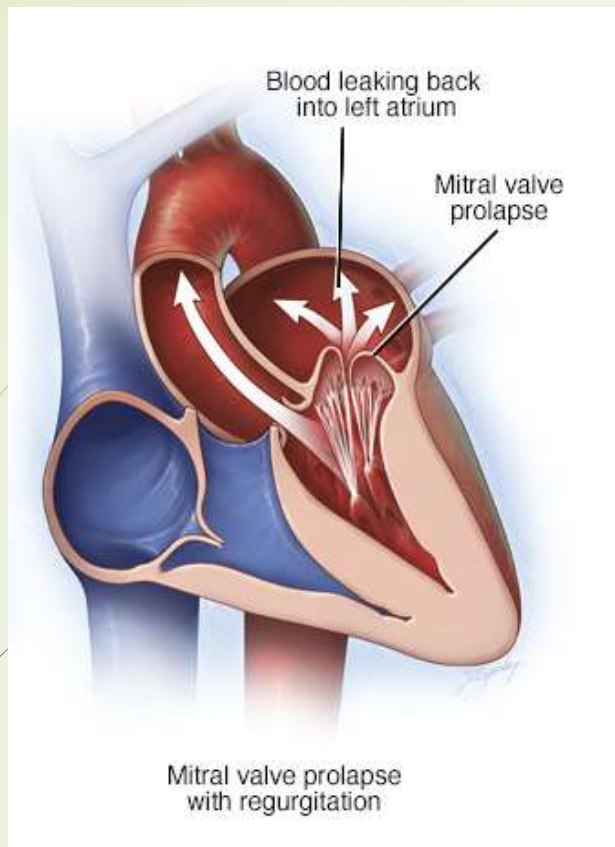
# Mitral Valve Etiology

MV Disease Etiology Choose PRIMARY Etiology (one): VDMiPrimEt (1731)	
<input type="checkbox"/> Myxomatous degeneration/prolapse	<input type="checkbox"/> Tumor, Papillary fibroelastoma
<input type="checkbox"/> Rheumatic	<input type="checkbox"/> Tumor, Other
<input type="checkbox"/> Ischemic- acute, post infarction (MI $\leq$ 21 days)	<input type="checkbox"/> Carcinoid
<input type="checkbox"/> Ischemic- chronic (MI $>$ 21 days)	<input type="checkbox"/> Trauma
<input type="checkbox"/> Non-ischemic Cardiomyopathy	<input type="checkbox"/> Congenital
<input type="checkbox"/> Endocarditis	<input type="checkbox"/> Pure annular dilatation
<input type="checkbox"/> Hypertrophic Obstructive Cardiomyopathy (HOCM)	<input type="checkbox"/> Reoperation-Failure of previous MV repair or replacement
<input type="checkbox"/> Tumor, Carcinoid	<input type="checkbox"/> Mixed Etiology
<input type="checkbox"/> Tumor, Myxoma	<input type="checkbox"/> Not Documented

- **Designed to address both types of Mitral Disease: Insufficiency and Stenosis.**
- **Etiology is the “why” that causes the disease state.**
- **New to v2.9, choice is limited to one (Primary).**
- **While there is no hierarchy, some etiologies are more common than others.**
- **“Mixed Etiology” definition is unclear.**

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**Myxomatous Degeneration/Prolapse**  
**Rheumatic**  
**Endocarditis**  
**Ischemic**  
**Cardiomyopathy**

**Rheumatic**

# Myxomatous Degeneration

- **A pathological weakening of connective tissue.**
- **The fibrous collagen layer of the valve thins and mucoid (myxomatous) material accumulates.**
- **The chordae tendinae become longer and thinner and the valve leaflets enlarge and become rubbery.**
- **These changes result in floppy valve leaflets that balloon back (prolapse) into the left atrium when the left ventricle contracts, leading to regurgitation.**
- **Rupture of a degenerated chord can allow part of the valve leaflet to flail into the atrium, causing severe regurgitation.**

# Mitral Valve Lesion

MV Lesion Choose PRIMARY Lesion (one):	
VDMiPrimLes (1746)	
<input type="checkbox"/> Leaflet prolapse, posterior	<input type="checkbox"/> Papillary muscle elongation
<input type="checkbox"/> Leaflet prolapse, bileaflet	<input type="checkbox"/> Papillary muscle rupture
<input type="checkbox"/> Leaflet prolapse, anterior	<input type="checkbox"/> Leaflet thickening
<input type="checkbox"/> Leaflet prolapse, unspecified	<input type="checkbox"/> Leaflet retraction
<input type="checkbox"/> Elongated/ruptured chord(s)/Flail	<input type="checkbox"/> Chordal tethering
<input type="checkbox"/> Annular dilatation	<input type="checkbox"/> Chordal thickening/retraction/fusion
<input type="checkbox"/> Leaflet calcification	<input type="checkbox"/> Commissural fusion
<input type="checkbox"/> Leaflet perforation/hole	<input type="checkbox"/> Mixed lesion
<input type="checkbox"/> Mitral annular calcification	<input type="checkbox"/> Not Documented

- Again, “Lesion” pertains to both disease states.
- Lesion is the “what” that is contributing to the disease process.
- As with Etiology, only one choice (Primary) applies.
- “Mixed Lesion” is unclear.

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# #5: How do we abstract this patient's Mitral Etiology (1731) and Mitral Lesion (1746)?

## Answer Choices:

- Mitral Etiology is Myxomatous Degeneration/Prolapse; Mitral Lesion is Leaflet Prolapse, Bi-Leaflet and Leaflet Thickening.
- Mitral Etiology is Rheumatic; Mitral Lesion is Leaflet Prolapse, Unspecified.
- Mitral Etiology is Mixed Etiology; Mitral Lesion is Mixed Lesion.
- Mitral Etiology is Myxomatous Degeneration/Prolapse; Mitral Lesion is Leaflet Prolapse, Anterior

MV Disease Etiology Choose PRIMARY Etiology (one):	
VDMiPrimEt (1731)	
<input checked="" type="checkbox"/> Myxomatous degeneration/prolapse	<input type="checkbox"/> Tumor, Papillary fibroelastoma
<input type="checkbox"/> Rheumatic	<input type="checkbox"/> Tumor, Other
<input type="checkbox"/> Ischemic- acute, post infarction (MI ≤ 21 days)	<input type="checkbox"/> Carcinoid
<input type="checkbox"/> Ischemic- chronic (MI > 21 days)	<input type="checkbox"/> Trauma
<input type="checkbox"/> Non-ischemic Cardiomyopathy	<input type="checkbox"/> Congenital
<input type="checkbox"/> Endocarditis	<input type="checkbox"/> Pure annular dilatation
<input type="checkbox"/> Hypertrophic Obstructive Cardiomyopathy (HOCM)	<input type="checkbox"/> Reoperation-Failure of previous MV repair or replacement
<input type="checkbox"/> Tumor, Carcinoid	<input type="checkbox"/> Mixed Etiology
<input type="checkbox"/> Tumor, Myxoma	<input type="checkbox"/> Not Documented

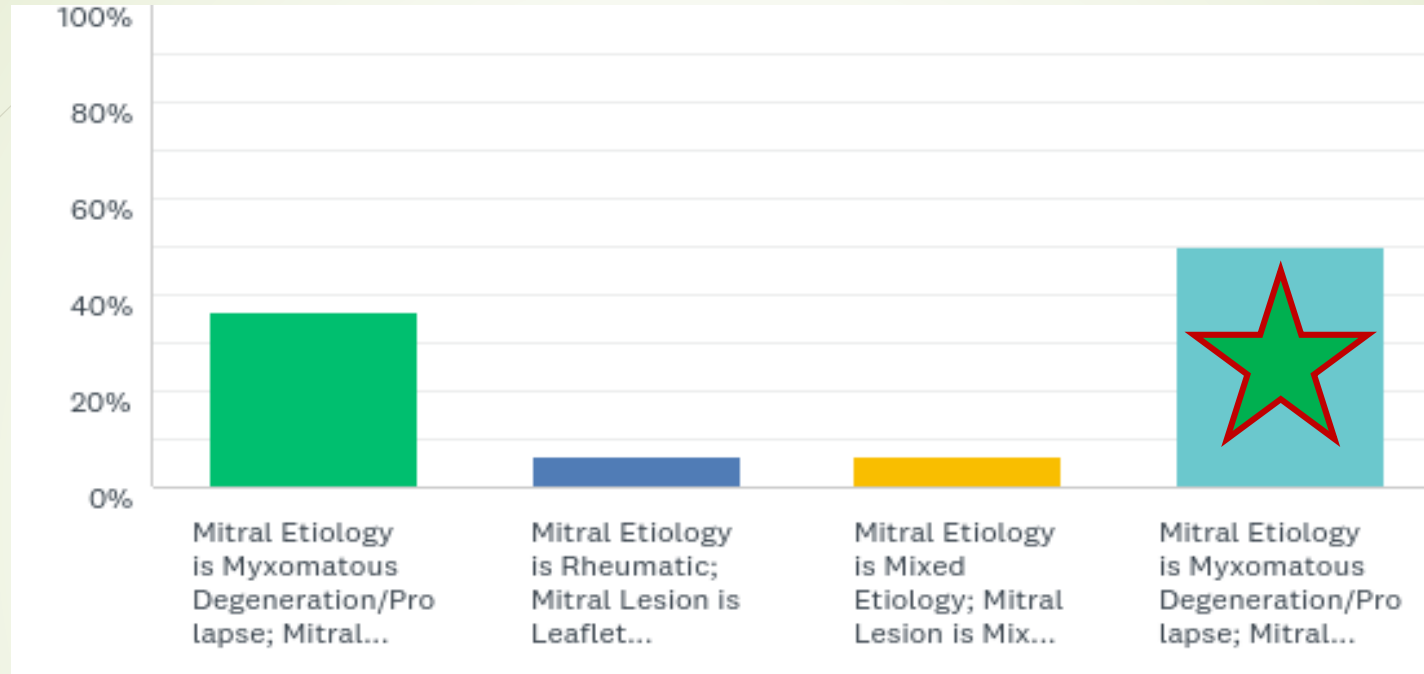
  

MV Lesion Choose PRIMARY Lesion (one):	
VDMiPrimLes (1746)	
<input type="checkbox"/> Leaflet prolapse, posterior	<input type="checkbox"/> Papillary muscle elongation
<input type="checkbox"/> Leaflet prolapse, bileaflet	<input type="checkbox"/> Papillary muscle rupture
<input checked="" type="checkbox"/> Leaflet prolapse, anterior	<input type="checkbox"/> Leaflet thickening
<input type="checkbox"/> Leaflet prolapse, unspecified	<input type="checkbox"/> Leaflet retraction
<input type="checkbox"/> Elongated/ruptured chord(s)/Flail	<input type="checkbox"/> Chordal tethering
<input type="checkbox"/> Annular dilatation	<input type="checkbox"/> Chordal thickening/retraction/fusion
<input type="checkbox"/> Leaflet calcification	<input type="checkbox"/> Commissural fusion
<input type="checkbox"/> Leaflet perforation/hole	<input type="checkbox"/> Mixed lesion
<input type="checkbox"/> Mitral annular calcification	<input type="checkbox"/> Not Documented

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# #5: How would you abstract this patient's Mitral Etiology (1731) and Mitral Lesion (1746)?



## Answer Choices

Mitral Etiology is Myxomatous Degeneration/Prolapse; Mitral Lesion is Leaflet Prolapse, Bi-Leaflet and Leaflet Thickening.

Mitral Etiology is Rheumatic; Mitral Lesion is Leaflet Prolapse, Unspecified.

Mitral Etiology is Mixed Etiology; Mitral Lesion is Mixed Lesion.

Mitral Etiology is Myxomatous Degeneration/Prolapse; Mitral Lesion is Leaflet Prolapse, Anterior.

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## #6: Which Ejection Fraction (1545) should be abstracted?

### Intent/Clarification:

- Use the **most recent determination prior to the induction of anesthesia** documented on a diagnostic report, regardless of the diagnostic procedure to obtain it.
- If no diagnostic report specifying an ejection fraction (EF) is in the medical record, a value documented in the progress record is acceptable.
- **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.**
- Use the surgeon's documentation if more than one value is reported as this was likely used to plan operative care.
- Time Frame: **Collect the last value closest to incision, not greater than 6 months.**

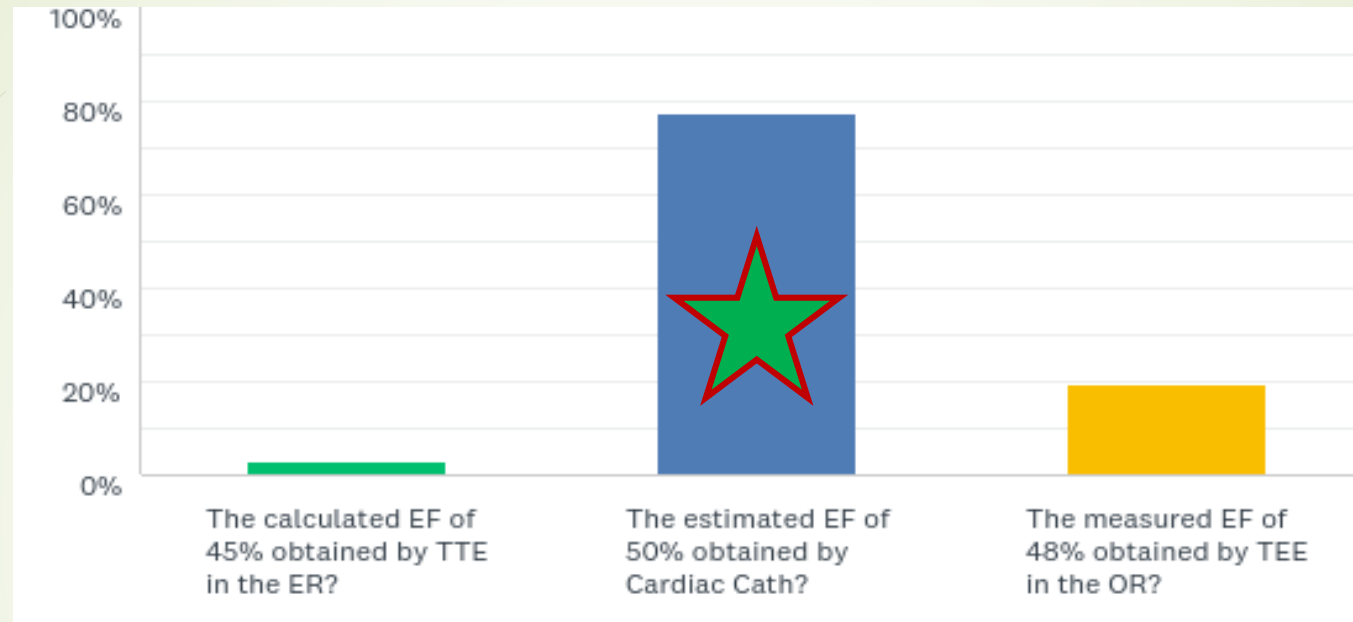
## #6: Which Ejection Fraction (1545) should be abstracted?

### Answer Choices:

- The *calculated* EF of 45% obtained by TTE in the ER?
- The *estimated* EF of 50% obtained by Cardiac Cath?
- The *measured* EF of 48% obtained by TEE in the OR?

Stress Test: <input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes →) Result: <input type="checkbox"/> Negative (Normal) <input type="checkbox"/> Positive (Abnormal) <input type="checkbox"/> Not Documented		
StressTst (1525) StrsTstRes (1531)		
Ejection Fraction Done: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No (If Yes →)		Ejection Fraction: <u>50</u> (%)
HDEFD (1540)		HDEF (1545)
Dimensions Available: <input type="checkbox"/> Yes <input type="checkbox"/> No (If Yes →)	LV End-Systolic Dimension: _____ (mm)	LV End-Diastolic Dimension: _____ (mm)
DimAvail (1555)	LVSD (1560)	LVEDD (1565)

## #6: Which Ejection Fraction (1545) should be abstracted?



### Answer Choices

The calculated EF of 45% obtained by TTE in the ER?

The estimated EF of 50% obtained by Cardiac Cath?

The measured EF of 48% obtained by TEE in the OR?



## #7: Given the patient's pre-op history of AFib, does this patient suffer from post-op AFib (6930)?

**SEQ. #: 6930 Long Name: Post-Op-Other-A Fib Short Name: COtAFib Definition:** Indicate whether the patient experienced atrial fibrillation/flutter (AF) requiring treatment. Exclude patients who were in AFib at the start of surgery.

**Intent/Clarification:** Include any episode of A-Fib lasting longer than one hour and/or requiring treatment. **Capture event(s) in all patients who were not in A-Fib at the start of surgery.**

**Ver. 2.8 Example:** A patient is on a protocol preoperatively; the patient then goes in to atrial fibrillation (AF) postoperatively and the protocol is not adjusted: **If the patient was in sinus rhythm and then develops AF postoperatively, this should be coded "Yes" as a post op event.**

## #7: Given the patient's pre-op history of AFib, does this patient suffer from post-op AFib (6930)?

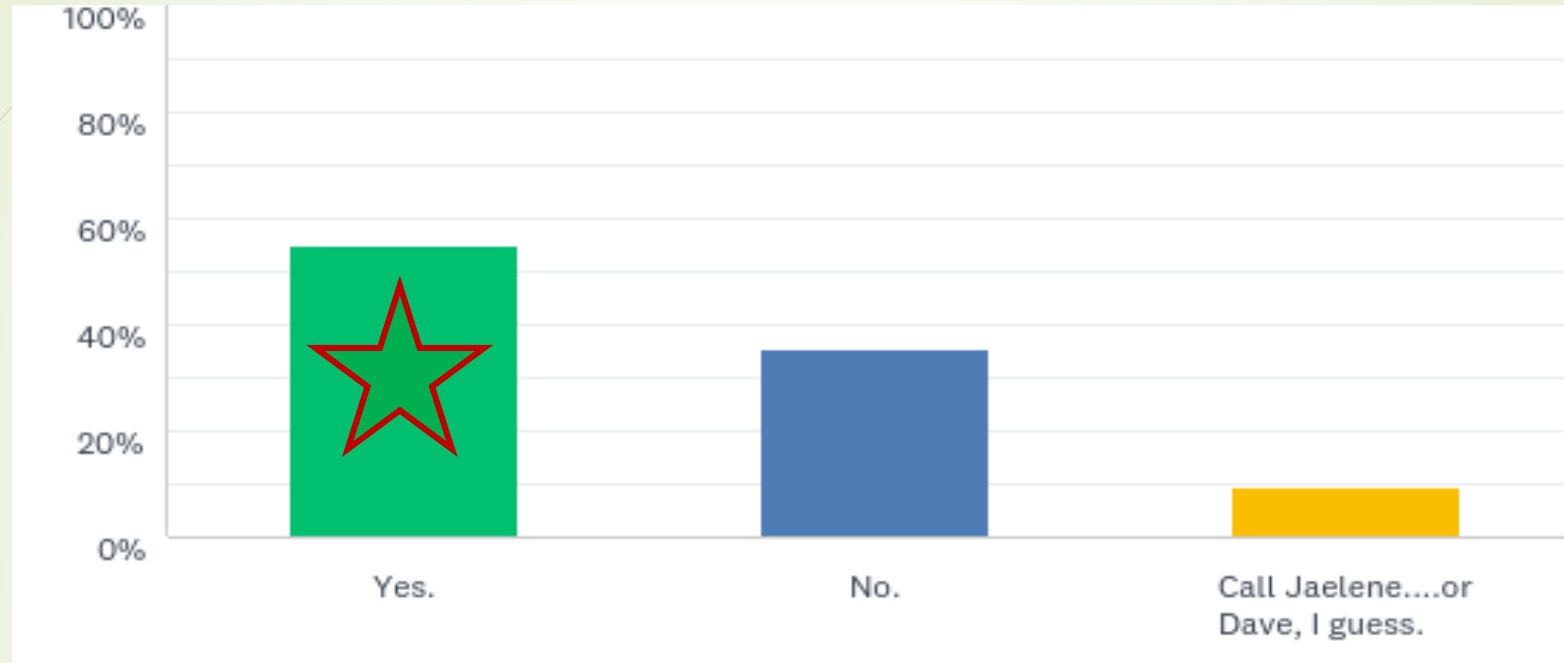
### Answer Choices:

- Yes.
- No.
- Call Jaelene....or Dave, I guess.

<b>Other</b>
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 <b>CRhythmDis (6900)</b>
Cardiac Arrest: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTarrst (6905)</b>
Post Op Aortic Endoleak: <input type="checkbox"/> Yes <input type="checkbox"/> No (if yes→) Type: <input type="checkbox"/> Ia <input type="checkbox"/> Ib <input type="checkbox"/> II <input type="checkbox"/> III <input type="checkbox"/> IV <input type="checkbox"/> V <b>COTaortEndo (6906)</b> <b>COTaortEndoTy (6907)</b>
Aortic Rupture: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTaortRupt (6908)</b>
Aortic Dissection: <input type="checkbox"/> Yes <input type="checkbox"/> No (if yes→) Type: <input type="checkbox"/> Antegrade <input type="checkbox"/> Retrograde <input type="checkbox"/> Both <b>CVaAoDis (6909)</b> <b>CVaAoDisTy (6910)</b>
Aortic Side Branch malperfusion: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTaortSide (6911)</b>
Aortic stent graft induced entry tear: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTaortTear (6912)</b>
Anticoagulant Event: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTCoag (6914)</b>
Pericardiocentesis: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTamp (6915)</b>
Gastro-Intestinal Event: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTGI (6920)</b>
Liver Dysfunction/ Failure: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTLiver (6921)</b>
Multi-System Failure: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTMSF (6925)</b>
Atrial Fibrillation: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <b>COTAFib (6930)</b>
Other: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>COTOther (6950)</b>

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## #7: Given the patient's pre-op history of A-Fib, does this patient suffer from post-op A-Fib (6930)?



Answer Choices
Yes.
No.
Call Jaelene....or Dave, I guess.

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## #8: How would you code this patient's Readmission (7140)?

**SEQ. #:** 7140

**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.

**Intent/Clarification:**

**This is not part of the composite score.**

The intent is to capture inpatient readmissions to acute care and primary care facilities only where the patient **status is listed as "In-Patient"**.

- Obtain information as close to 30 days from date of discharge as possible.
- It is understood that some readmissions are planned; these are still counted as readmissions.
  
- To code "Yes", readmissions do not need to be at same institution where the initial surgical procedure was done.
- Discharge and readmission to a psychiatric care facilities, where the patient is considered an in-patient are to be considered as readmissions.
- Do not include Emergency Department visits or observation status visits unless the ED visits leads to status of in-patient.
- If a patient is readmitted to an in-patient rehabilitation hospital, code "No".
- If a patient is readmitted to an LTAC, code "No".
- Do not code transfers to higher level of care, this is considered an extension of the same acute care admission. If the patient was discharged to the "Acute Rehab" floor of the same hospital and then readmitted back as an in-patient back into a nursing floor, code "Yes" to admission as an inpatient is considered "Yes."
- To align with CMS, 30 day readmission should not be coded for patients who remain in observation units, no matter the duration.



## #8: How would you code this patient's Readmission (7140)?

### Answer Choices:

- Readmission is No.
- Readmission is Yes; Reason is Arrhythmia/Heart Block; Procedure is Other Procedure.
- Readmission is Yes; Reason is Arrhythmia/Heart Block; Procedure is No Procedure Performed.

#### R. Readmission

(If Discharge/Mortality Status = "Discharged alive, last know status=alive" or "Discharged alive, died after discharge" ↓)

Readmit:  Yes  No  Unknown (If Yes ↓)

Readmit (7140)

Readmit Date:    /    /    (mm/dd/yyyy)

# Expanded Readmission Reasons

Readmit Primary Reason:

ReadmRsn (7160)

- |   |  |
|---|--|
| <input type="checkbox"/> Angina   | <input type="checkbox"/> Pericardial Effusion and/or Tamponade   |
| <input type="checkbox"/> Anticoagulation Complication - Pharmacological | <input type="checkbox"/> Pericarditis/Post Cardiotomy Syndrome   |
| <input type="checkbox"/> Anticoagulation Complication – Valvular        | <input type="checkbox"/> Pleural effusion requiring intervention |
| <input type="checkbox"/> Aortic Complication                            | <input type="checkbox"/> Pneumonia                               |
| <input checked="" type="checkbox"/> Arrhythmia or Heart Block           | <input type="checkbox"/> Renal Failure                           |
| <input type="checkbox"/> Blood Pressure (hyper or hypotension)          | <input type="checkbox"/> Renal Insufficiency                     |
| <input type="checkbox"/> Chest pain, noncardiac                         | <input type="checkbox"/> Respiratory complication, Other         |
| <input type="checkbox"/> Congestive Heart Failure                       | <input type="checkbox"/> Sepsis                                  |
| <input type="checkbox"/> Coronary Artery/Graft Dysfunction              | <input type="checkbox"/> Stroke                                  |
| <input type="checkbox"/> Depression/psychiatric issue                   | <input type="checkbox"/> TIA                                     |
| <input type="checkbox"/> DVT  | <input type="checkbox"/> Transfusion                             |
| <input type="checkbox"/> Electrolyte imbalance                          | <input type="checkbox"/> Transplant Rejection                    |
| <input type="checkbox"/> Endocarditis                                   | <input type="checkbox"/> VAD Complication                        |
| <input type="checkbox"/> Failure to thrive                              | <input type="checkbox"/> Valve Dysfunction                       |
| <input type="checkbox"/> GI issue                                       | <input type="checkbox"/> Vascular Complication, acute            |
| <input type="checkbox"/> Infection, Conduit Harvest Site                | <input type="checkbox"/> Wound , other (drainage, cellulitis)    |
| <input type="checkbox"/> Infection, Deep Sternum / Mediastinitis        | <input type="checkbox"/> Other – Related Readmission             |
| <input type="checkbox"/> Mental status changes                          | <input type="checkbox"/> Other – Nonrelated Readmission          |
| <input type="checkbox"/> Myocardial Infarction                          | <input type="checkbox"/> Other – Planned Readmission             |
| <input type="checkbox"/> PE   | <input type="checkbox"/> Unknown                                 |

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# Expanded Readmission Procedures

Readmit Primary Procedure:

**ReadmPro (7165)**

- |   |  |
|---|--|
| <input type="checkbox"/> No Procedure Performed                   | <input type="checkbox"/> OR for Vascular Procedure           |
| <input type="checkbox"/> Cath lab for Valve Intervention          | <input type="checkbox"/> OR for Aorta Intervention           |
| <input type="checkbox"/> Cath lab for Coronary Intervention (PCI) | <input type="checkbox"/> Pacemaker Insertion / AICD          |
| <input type="checkbox"/> Dialysis                                 | <input type="checkbox"/> Pericardiotomy / Pericardiocentesis |
| <input type="checkbox"/> OR for Bleeding                          | <input type="checkbox"/> Planned noncardiac procedure        |
| <input type="checkbox"/> OR for Coronary Artery Intervention      | <input type="checkbox"/> Thoracentesis/ Chest tube insertion |
| <input type="checkbox"/> OR for Sternal Debridement / Muscle Flap | <input type="checkbox"/> Wound vac                           |
| <input type="checkbox"/> OR for Valve Intervention                | <input checked="" type="checkbox"/> Other Procedure          |
|   | <input type="checkbox"/> Unknown                             |

(if OR for Aorta intervention→)

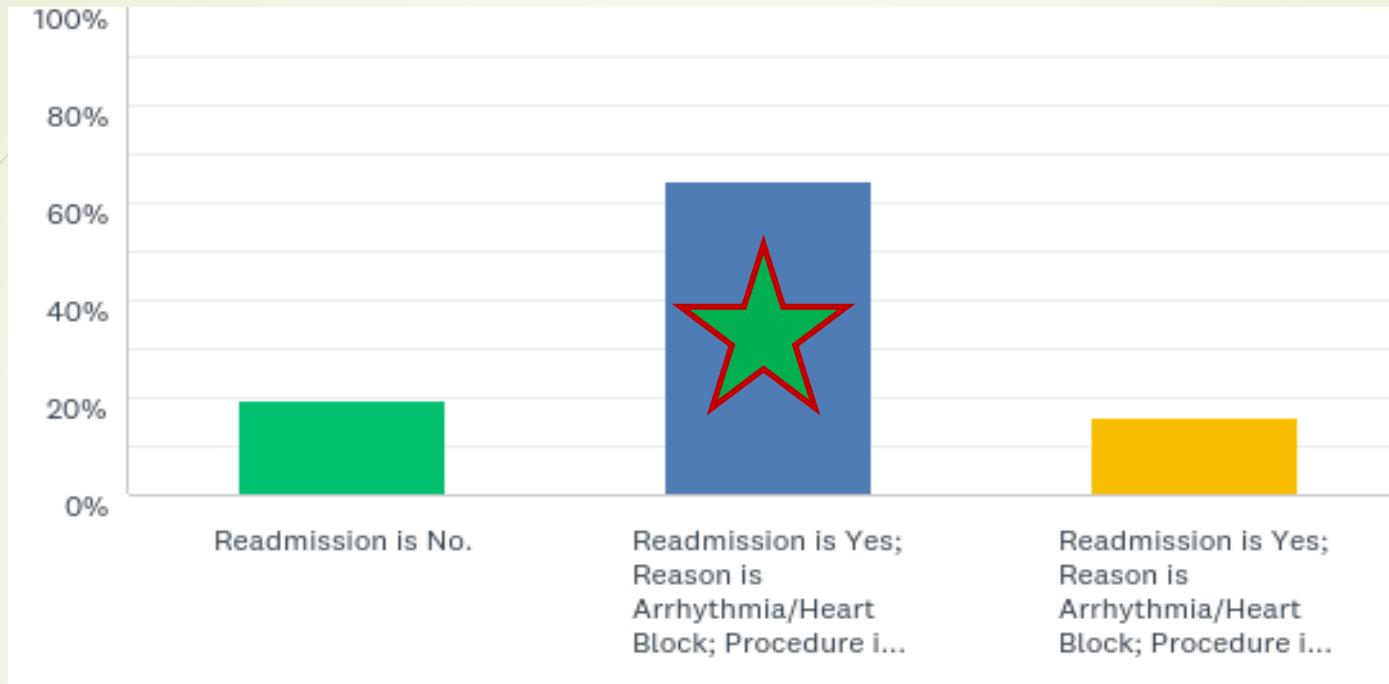
Type:  Open  Endovascular

**ReadmAortIntTy (7166)**

Indication:  Rupture  Endoleak  Infection  Dissection  Expansion  Loss of side branch patency  Other

**ReadmAortIntInd (7167)**

## #8: How would you code this patient's Readmission (7140)?



### Answer Choices

Readmission is No.

Readmission is Yes; Reason is Arrhythmia/Heart Block; Procedure is Other Procedure.

Readmission is Yes; Reason is Arrhythmia/Heart Block; Procedure is No Procedure Performed.

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# Scenario #2

An elderly patient is brought to the ER as a result of a syncopal episode at home. The patient had regained consciousness shortly after arrival, but began to complain of severe back pain, and was taken for CT scans of the head, neck, chest, and abdomen, with the following results:

*"Dissection of the ascending thoracic aorta which originates in the ascending thoracic aorta just above the sinotubular junction, and extends throughout the aorta and into the iliac vasculature. A secondary tear is evident in the descending thoracic aorta 2 cm. above the diaphragm. The dissection compromises the innominate, left common and right common carotid arteries, with up to 75% diminished flow of the true lumen of the vessels."*

The patient is taken for emergent surgery, where the surgeon replaces the ascending aorta and hemi-arch with a 30 mm. dacron graft while undergoing circulatory arrest. After an extended and complicated recovery, the patient is discharged to an acute rehab facility on POD #24. On POD #27, the facility phones to report the patient had suffered a cardiac arrest, and could not be revived.

## #9: Which of the following Risk Factors (section D.) would you abstract for this patient?

### Answer Choices:

- Thoracic Artery Disease (510).
- Thoracic Artery Disease (510), Cerebrovascular Disease, Carotid Stenosis, Both (545).
- Thoracic Artery Disease (510), Peripheral Artery Disease (505) and Cerebrovascular Disease, TIA (540).
- All of the above. #'s 505, 510, 540, 545.

**SEQ. #:** 510

**Long Name:** RF-Thoracic Aorta Disease

**Short Name:** ThAoDisease

**Definition:** Indicate whether the patient has a history of disease of the thoracic or thoracoabdominal aorta.

Abdominal aortic disease without thoracic involvement is captured in peripheral artery disease.

**Intent/Clarification:**

Code "Yes" to aortic aneurysms, aortic dissection/rupture. Fusiform ascending thoracic aneurysm is more likely to dissect when the aortic cross clamp is applied and should be coded as thoracic aorta disease.

**SEQ. #: 505**

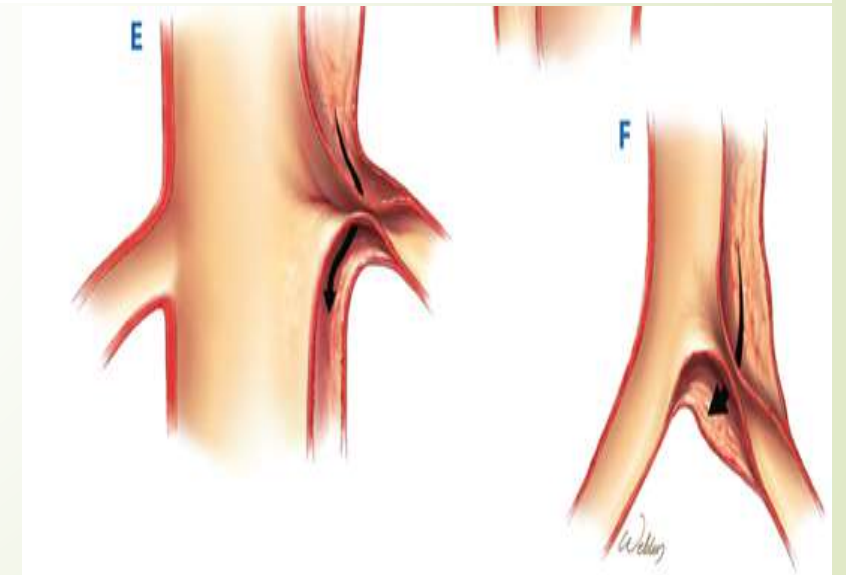
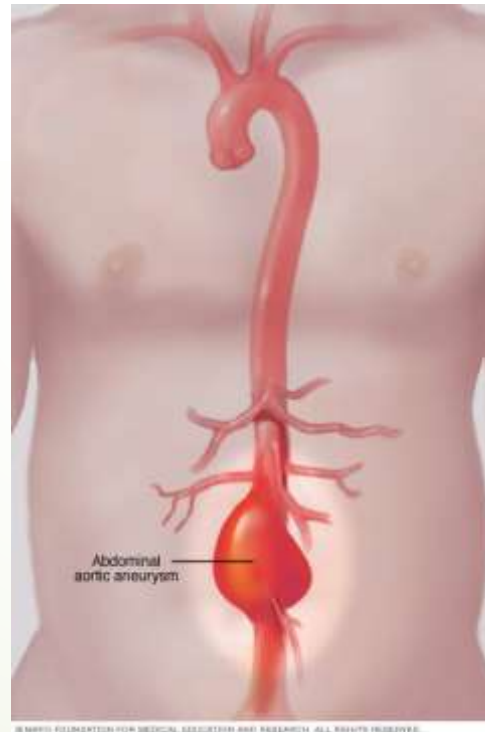
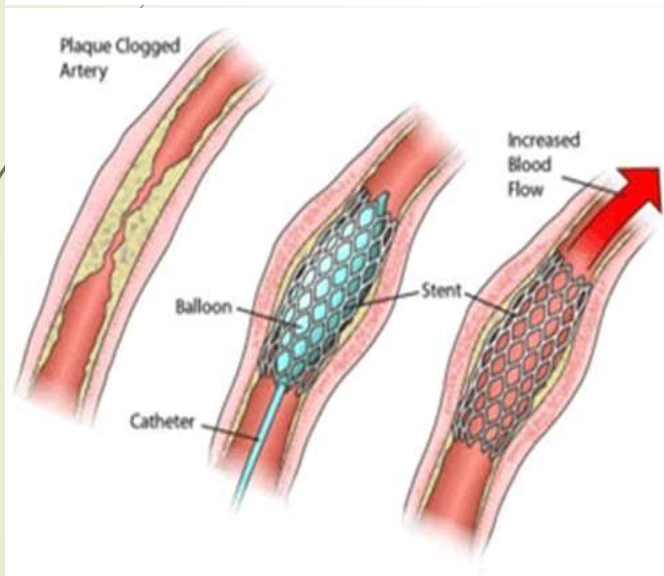
**Long Name: RF-Peripheral Arterial Disease**

**Short Name: PVD**

**Definition:** Indicate whether the patient has a history of peripheral arterial disease (includes upper and lower extremity, renal, mesenteric, and abdominal aortic systems).

Peripheral arterial disease excludes disease in the carotid, cerebrovascular arteries or thoracic aorta.

PVD does not include DVT.



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SEQ. #: 525

Long Name: RF-Cerebrovascular Dis

Short Name: CVD

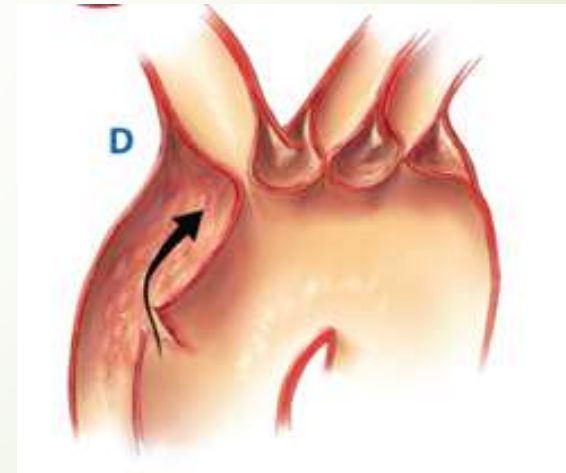
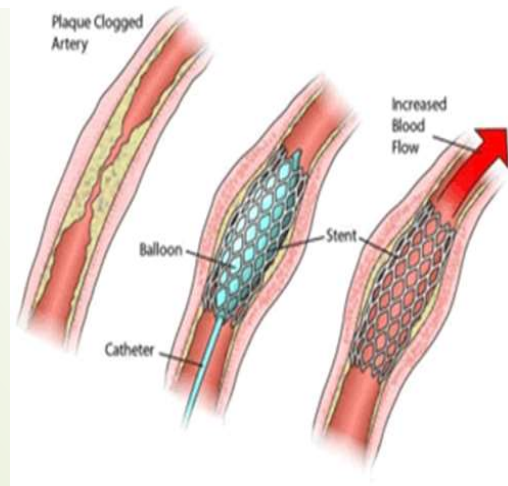
**Definition:** Indicate whether the patient has a current or previous history of any of the following:

A. Stroke: Stroke is an acute episode of focal or global neurological dysfunction caused by brain, spinal cord, or retinal vascular injury as a result of hemorrhage or infarction, where the neurological dysfunction lasts for greater than 24 hours.

B. TIA: is defined as a transient episode of focal neurological dysfunction caused by brain, spinal cord, or retinal ischemia, without acute infarction, where the neurological dysfunction resolves within 24 hours.

C. Noninvasive or invasive arterial imaging test demonstrating  $\geq 50\%$  stenosis of any of the major extracranial or intracranial vessels of the brain

D. Previous cervical or cerebral artery revascularization surgery or percutaneous intervention



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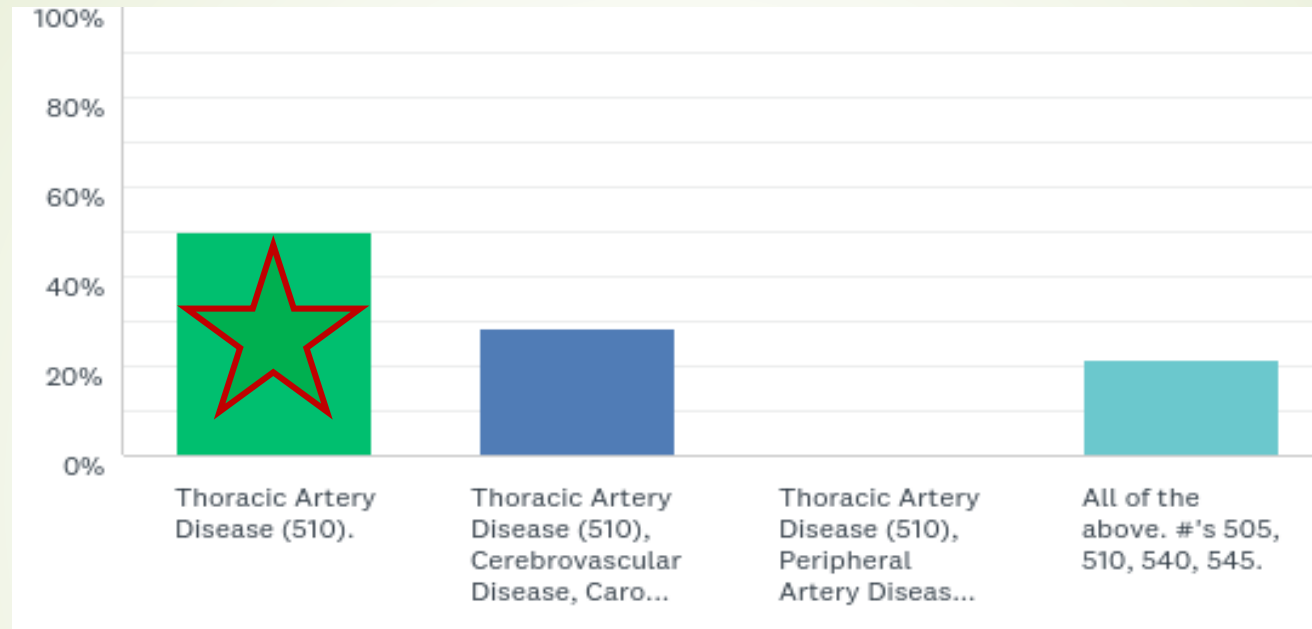
## Answer Choices:

- Thoracic Artery Disease (510).
- Thoracic Artery Disease (510), Cerebrovascular Disease, Carotid Stenosis, Both (545).
- Thoracic Artery Disease (510), Peripheral Artery Disease (505) and Cerebrovascular Disease, TIA (540).
- All of the above. #'s 505, 510, 540, 545.

Immunocompromise Present: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>ImmSupp (490)</b>	Mediastinal Radiation: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>MediastRad (495)</b>
Cancer Within 5 Years: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>Cancer (500)</b>	Peripheral Artery Disease: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>PVD (505)</b>
Thoracic Aorta Disease: <input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>ThAoDisease (510)</b>	Syncope: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>Syncope (515)</b>
Cerebrovascular Disease: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>CVD (525)</b>	
(If Yes →) Prior CVA: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes →) <b>CVA (530)</b>	Prior CVA-When: <input type="checkbox"/> ≤ 30 days <input type="checkbox"/> > 30 days <b>CVAWhen (535)</b>
CVD TIA: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <b>CVDTIA (540)</b>	
CVD Carotid stenosis: <input type="checkbox"/> Right <input type="checkbox"/> Left <input type="checkbox"/> Both <input type="checkbox"/> None <input type="checkbox"/> Not Documented <b>CVDCarSten (545)</b>	
(If "Right" or "Both" →)	Severity of stenosis on the right carotid artery: <input type="checkbox"/> 50-79% <input type="checkbox"/> 80 – 99% <input type="checkbox"/> 100% <input type="checkbox"/> Not documented <b>CVDCarStenRt (550)</b>
(If "Left" or "Both" →)	Severity of stenosis on the left carotid artery: <input type="checkbox"/> 50-79% <input type="checkbox"/> 80 – 99% <input type="checkbox"/> 100% <input type="checkbox"/> Not documented <b>CVDCarStenLft (555)</b>
History of previous carotid artery surgery and/or stenting: <input type="checkbox"/> Yes <input type="checkbox"/> No <b>CVDPCarSurg (560)</b>	



## #9: Which of the following Risk Factors (section D.) would you abstract for this patient?



### Answer Choices

Thoracic Artery Disease (510).

Thoracic Artery Disease (510), Cerebrovascular Disease, Carotid Stenosis, Both (545).

Thoracic Artery Disease (510), Peripheral Artery Disease (505) and Cerebrovascular Disease, TIA (540).

All of the above. #'s 505, 510, 540, 545.

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### STS Aorta Surgery Worksheet V2.9

**Identify procedural location using graph letters A-N →**

**For Aneurysm:**

Aneurysm/Primary Tear Location: \_\_\_\_\_

**For Dissections:**

(Select all that apply and fill in location)

Secondary Tear Location: \_\_\_\_\_

Retrograde Extension Location: \_\_\_\_\_

Distal Extension Location: \_\_\_\_\_

Rupture Location: \_\_\_\_\_

Rupture Contained:  Yes  No

**For Open Descending Thoracic Aorta or Thoracoabdominal Procedures:**

Proximal Location: \_\_\_\_\_  Reverse Heel

Distal Location: \_\_\_\_\_

**For Endovascular Procedures:**

Proximal Location: \_\_\_\_\_

Distal Location: \_\_\_\_\_ **Intra-OP!**

Unintentional Rupture of dissection septum

Location: \_\_\_\_\_

**[ PLEASE COMPLETE THE FOLLOWING SECTIONS ]**

Presentation	Endoleak:	<input type="checkbox"/> Type I (leak at graft attachment site) → Location: <input type="checkbox"/> Ia-proximal <input type="checkbox"/> Ib-distal <input type="checkbox"/> Ic-iliac occluded <input type="checkbox"/> Type II (aneurysm sac filling via branch vessel) → Number of vessel: <input type="checkbox"/> IIa - single vessel <input type="checkbox"/> IIb - two vessels or more <input type="checkbox"/> Type III (leak defect in graft) → Graft defect type: <input type="checkbox"/> IIIa - junctional separation of modular components <input type="checkbox"/> IIIb - endograft fractures/holes <input type="checkbox"/> Type IV (leak through graft fabric) <input type="checkbox"/> Type V (endoleak - expansion aneurysm sac without leak)
	Aneurysm:	<b>Etiology:</b> <input type="checkbox"/> Atherosclerosis <input type="checkbox"/> Infection <input type="checkbox"/> Inflammatory <input type="checkbox"/> Connective Tissue Disorder <input type="checkbox"/> Penetrating Ulcer <input type="checkbox"/> Pseudoaneurysm <input type="checkbox"/> Mycotic <input type="checkbox"/> Traumatic transection <input type="checkbox"/> Intercostal visceral patch <input type="checkbox"/> Anastomotic site <b>Type:</b> <input type="checkbox"/> Fusiform <input type="checkbox"/> Saccular <b>Rupture:</b> <input type="checkbox"/> Yes <input type="checkbox"/> No
	Dissection:	<b>Timing:</b> <input type="checkbox"/> Hyper-acute (<48hrs) <input type="checkbox"/> Acute (48hrs-2wks) <input type="checkbox"/> Sub-acute (>2wks-90days) <input type="checkbox"/> Chronic (>90days) <input type="checkbox"/> Acute on Chronic <b>Malperfusion:</b> <input type="checkbox"/> Yes (If Yes ↓) <input type="checkbox"/> No If Yes → <b>Subclavian</b> → <input type="checkbox"/> Right <input type="checkbox"/> Left <b>Common Carotid</b> → <input type="checkbox"/> Right <input type="checkbox"/> Left <b>Renal</b> → <input type="checkbox"/> Right <input type="checkbox"/> Left <input type="checkbox"/> Coronary <input type="checkbox"/> Celiac <input type="checkbox"/> Superior Mesenteric <input type="checkbox"/> Iliofemoral <input type="checkbox"/> Spinal
		<b>Infection (if yes →):</b> <input type="checkbox"/> Graft infection <input type="checkbox"/> Valvular endocarditis <input type="checkbox"/> Nonvalvular endocarditis <input type="checkbox"/> Native aorta <input type="checkbox"/> Multiple infection types <b>Trauma (if yes →):</b> <input type="checkbox"/> Rupture <input type="checkbox"/> Ascending <input type="checkbox"/> Arch <input type="checkbox"/> Descending <input type="checkbox"/> Thoracoabdominal <input type="checkbox"/> Abdominal
	<b>Root:</b> <input type="checkbox"/> Aorto-annular ectasia <input type="checkbox"/> Asymmetric Root Dilatation (if yes →) <input type="checkbox"/> Right <input type="checkbox"/> Left <input type="checkbox"/> Non-coronary <input type="checkbox"/> Sinus of Valsalva aneurysm (if yes →) <input type="checkbox"/> Right <input type="checkbox"/> Left <input type="checkbox"/> Non-coronary	
	<b>Arch Type:</b> <input type="checkbox"/> Right <input type="checkbox"/> Left <input type="checkbox"/> Aberrant subclavian → <input type="checkbox"/> Right <input type="checkbox"/> Left / <input type="checkbox"/> Kommerell / <input type="checkbox"/> Bovine / <input type="checkbox"/> Variant vertebral origin / <input type="checkbox"/> Patent IMA bypass graft	
	<b>Ascending:</b> <input type="checkbox"/> Asymmetric Dilatation <input type="checkbox"/> Proximal coronary artery bypass grafts	

### STS Aorta Surgery Worksheet V2.9

**Intervention:**

Planned stage hybrid  
 Open Arch Procedure (If Yes ↓)  
 Distal Technique:  Open  Clamped  
 Site →  Ascending Aorta  Femoral  Zone 1  Zone 2  Zone 3  Zone 4 (Refer to graph on 1<sup>st</sup> page)  
 Extension →  Elephant Trunk  Frozen Elephant Trunk  No  
 Arch Branch Reimplantation (If Yes →) **Subclavian** →  Right  Left **Common Carotid** →  Right  Left  
 Innominate  Left Vertebral  Other

**Open Descending Thoracic Aorta or Thoracoabdominal Procedure:**

Intercostal reimplantation  
 Visceral vessel intervention (If Yes →)  Celiac →  Reimplantation  Branch Graft  
 Superior mesenteric →  Reimplantation  Branch Graft  
 Right renal →  Reimplantation  Branch Graft  
 Left renal →  Reimplantation  Branch Graft

**Endovascular Procedure:**

Access (If Yes →)  Femoral  Iliac  Abdominal Aorta  Lt. Subclavian  Rt. Subclavian  Ascending Aorta  LV Apex  
 Percutaneous Access  TAVR (for combination procedures)  Ascending TEAVR (If Yes →)  Dedicated IDE  Off-label stent  No

**Arch Vessel Management:**

Innominate →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass (If Yes →)  Aorta-Innominate  Aorta-Rt Carotid  Aorta Rt Subclavian  Rt Carotid-Rt Subclavian  Other  
 Left Carotid →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass (If Yes →)  Aorta-Lt Carotid  Innominate-Lt Carotid  Rt Carotid-Lt Carotid  Other  
 Left Subclavian →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass (If Yes →)  Aorta-Lt Subclavian  Lt Carotid-Lt Subclavian  Other  
 Other Arch Vessel(s) Extra-anatomic bypass (If Yes →)  Innominate-Carotid  Innominate-Subclavian  Subclavian-Subclavian  Other

**Visceral Vessel Management:**

Celiac →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass →  Aorta-celiac  Iliac-celiac  Other  
 Superior mesenteric →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass →  Aorta-superior mesenteric  Iliac-superior mesenteric  Other  
 Right renal →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass →  Aorta-right renal  Iliac-right renal  Other  
 Left renal →  Native Flow  Endovascular Branch Graft  Endovascular Parallel Graft  Fenestrated  
 Extra-anatomic Bypass →  Aorta-left renal  Iliac-left renal  Other  
 Right Iliac →  Native Flow  Bifurcated Graft  Extra-anatomic Bypass →  Fem-Fem  Other  
 Left Iliac →  Native Flow  Bifurcated Graft  Extra-anatomic Bypass →  Fem-Fem  Other  
 Internal Iliac Preserved →  Rt Iliac only  Lt Iliac only  Both  No  
 Other Visceral Vessel(s) →  Extra-anatomic Bypass →  Aorta-Other  Iliac-Other  Other

**Intra-Op (Check all that apply):**

Dissection proximal entry tear covered  
 Endoleak at end of procedure → Type:  Ia  Ib  II  III  IV  V  
 Conversion to open →  Deployment failure  Endoleak  Rupture  Occlusion/loss of branch  
 Intra-Op Dissection Extension →  None  Antegrade  Retrograde  Both  
 Spinal drain placement →  Pre-Aortic procedure  Post-Aortic procedure  
 IntraOp Motor Evoked Potential → Documented MEP abnormality →  Yes  No  
 IntraOp Somatosensory Evoked Potential → Documented SEP abnormality →  Yes  No  
 IntraOp EEG → Documented EEG abnormality →  Yes  No  Unknown  
 IVUS Performed Intra-Op  
 IntraOp Transcatheter Doppler Performed Intra-Op  
 IntraOp Angiogram → Volume of Contrast \_\_\_\_\_ ml Flow time \_\_\_\_\_ min

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## #10: Based on the CT scan results, how would you abstract the following fields in Section M2?

### Answer Choices:

- Primary Tear Location (4750) is Below STJ (Zone A); Secondary Tear (4755) is Zone 6.
- Primary Tear Location (4750) is STJ to mid-ascending (Zone B); Secondary Tear (4755) is Zone 5.
- Primary Tear Location (4750) is mid-ascending to distal ascending (Zone C); Secondary Tear (4755) is Zone 5.

### Points to Consider:

- Primary Tear
- Secondary Tear
- Location, Location, Location

***"Dissection of the ascending thoracic aorta which originates in the ascending thoracic aorta just above the sinotubular junction, and extends throughout the aorta and into the iliac vasculature. A secondary tear is evident in the descending thoracic aorta 2 cm. above the diaphragm. The dissection compromises the innominate, left common and right common carotid arteries, with up to 75% diminished flow of the true lumen of the vessels."***



**SEQ. #:** 4750

**Long Name:** Dissection - Primry Tear Location **Short Name:** DisTearLoc

**Definition:** Indicate location of the primary tear

**Intent/Clarification:**

The intent is to identify the primary entry tear for the dissection. As most dissections include multiple re-entry tears it may be difficult to confirm the primary site and the surgeon MUST be the final arbiter of this definition. This is the site identified by the surgeon at an open operation or judged by the surgeon from imaging as the primary site to be covered by endovascular stent. If the radiology report names a primary entry point and the surgeon concurs, report this location.

**SEQ. #:** 4755

**Long Name:** Dissection - Secondary Tear Location **Short Name:** DisSecLoc

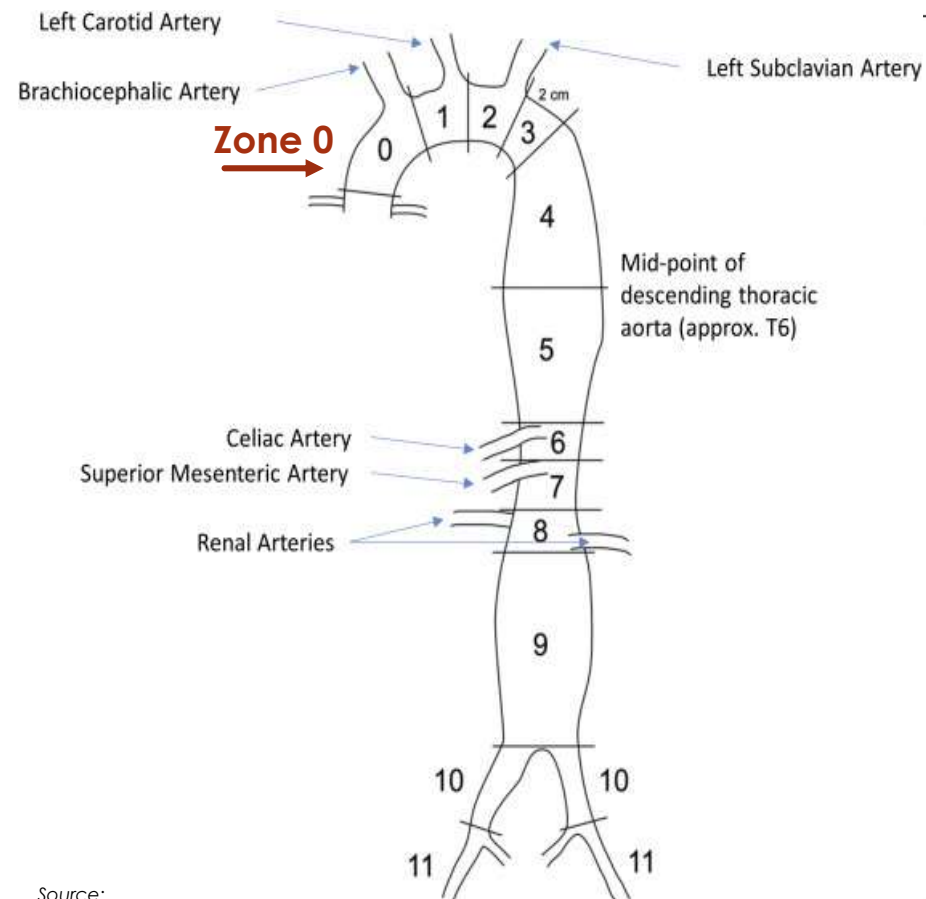
**Definition:** Indicate location of secondary tear

**Intent/Clarification:**

The intent is to identify any secondary tear for the dissection. This would be a re-entry site resulting from flow within the false lumen returning to the true lumen. The surgeon MUST be the final arbiter of this definition. This is the site identified by the surgeon at open operation or judged by the surgeon from imaging as a secondary site to be covered by endovascular stent.

## Intent/Clarification:

Zone 0 is the Ascending Aorta and includes letter A-C. Verify exact location with CV Surgeon. Aortic Root (letter A) is below sinotubular junction.

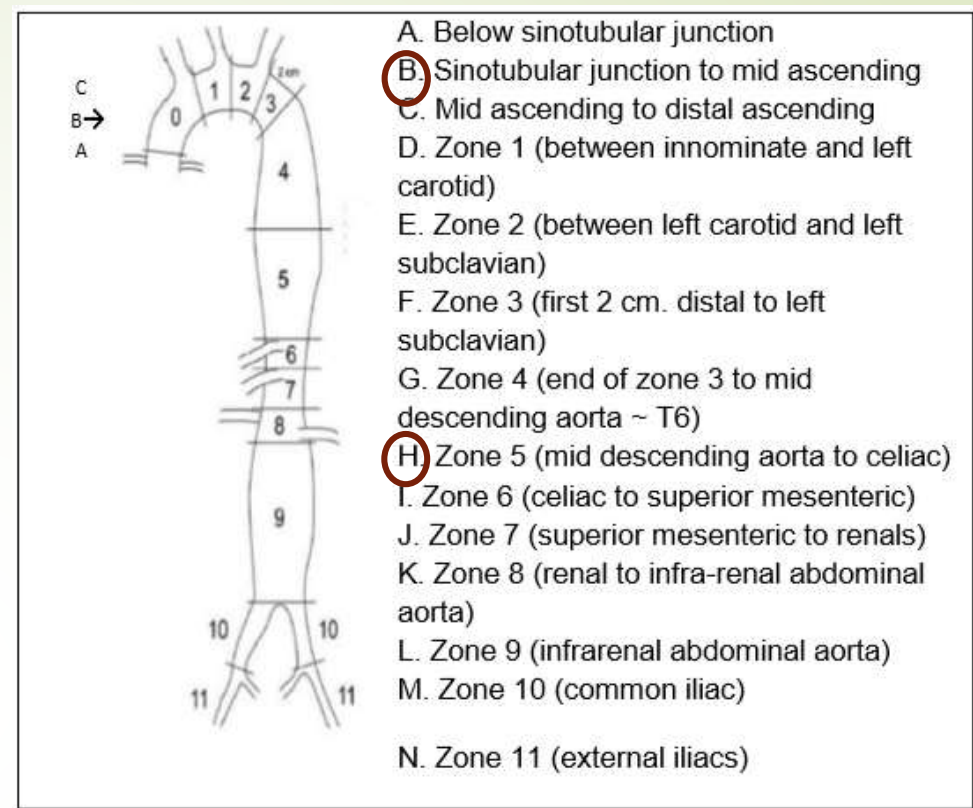
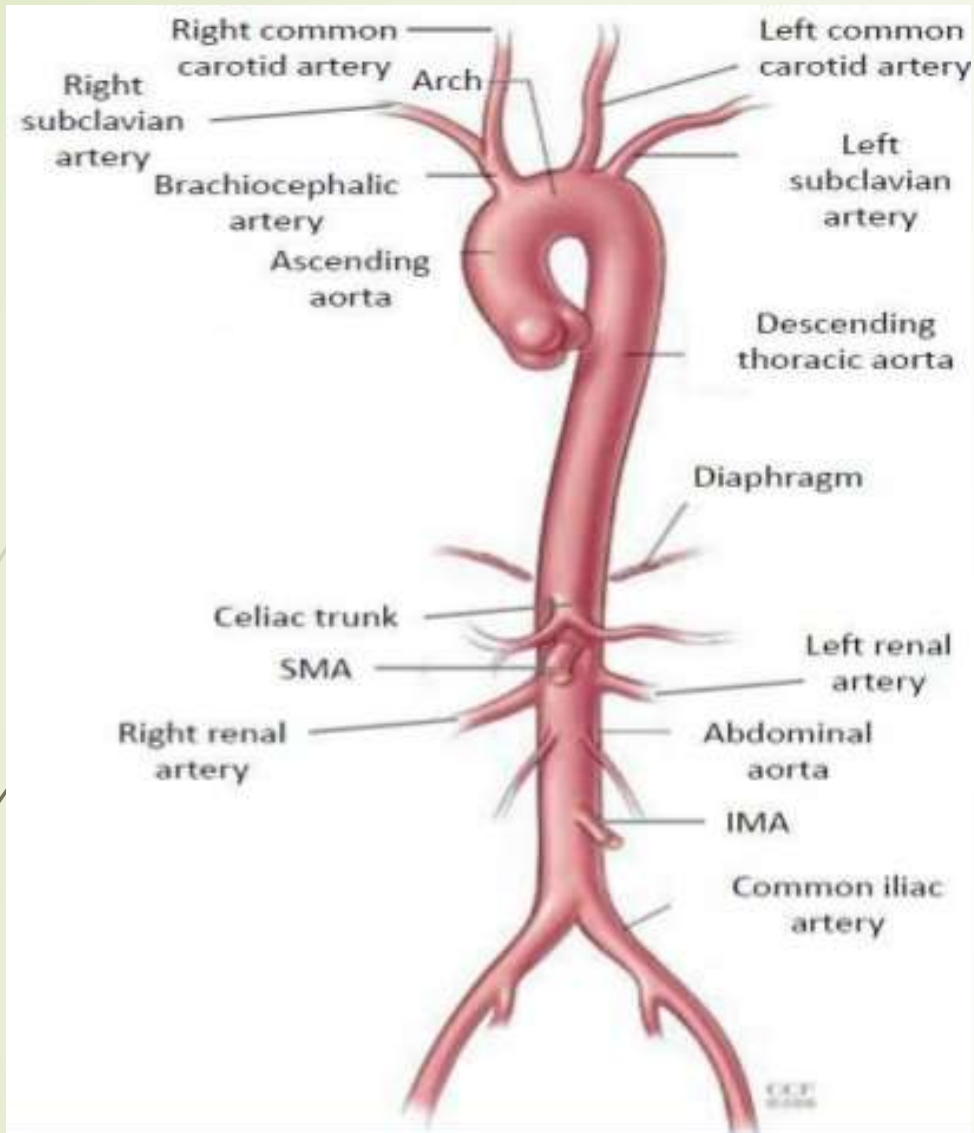


- A. Below sinotubular junction
- B. Sinotubular junction to mid ascending
- C. Mid ascending to distal ascending
- D. Zone 1 (between innominate and left carotid)
- E. Zone 2 (between left carotid and left subclavian)
- F. Zone 3 (first 2 cm. distal to left subclavian)
- G. Zone 4 (end of zone 3 to mid descending aorta ~ T6)
- H. Zone 5 (mid descending aorta to celiac)
- I. Zone 6 (celiac to superior mesenteric)
- J. Zone 7 (superior mesenteric to renals)
- K. Zone 8 (renal to infra-renal abdominal aorta)
- L. Zone 9 (infra-renal abdominal aorta)
- M. Zone 10 (common iliac)
- N. Zone 11 (external iliacs)

Source:  
Society of Thoracic Surgeons Training Manual

Source:  
Canadian Cardiovascular Society





## #10: Based on the CT scan results, how would you abstract the following fields in Section M2?

### Answer Choices:

- Primary Tear Location (4750) is Below STJ (Zone A); Secondary Tear (4755) is Zone 6.
- Primary Tear Location (4750) is STJ to mid-ascending (Zone B); Secondary Tear (4755) is Zone 5.
- Primary Tear Location (4750) is mid-ascending to distal ascending (Zone C); Secondary Tear (4755) is Zone 5.

Primary tear location:

DisTearLoc (4750)

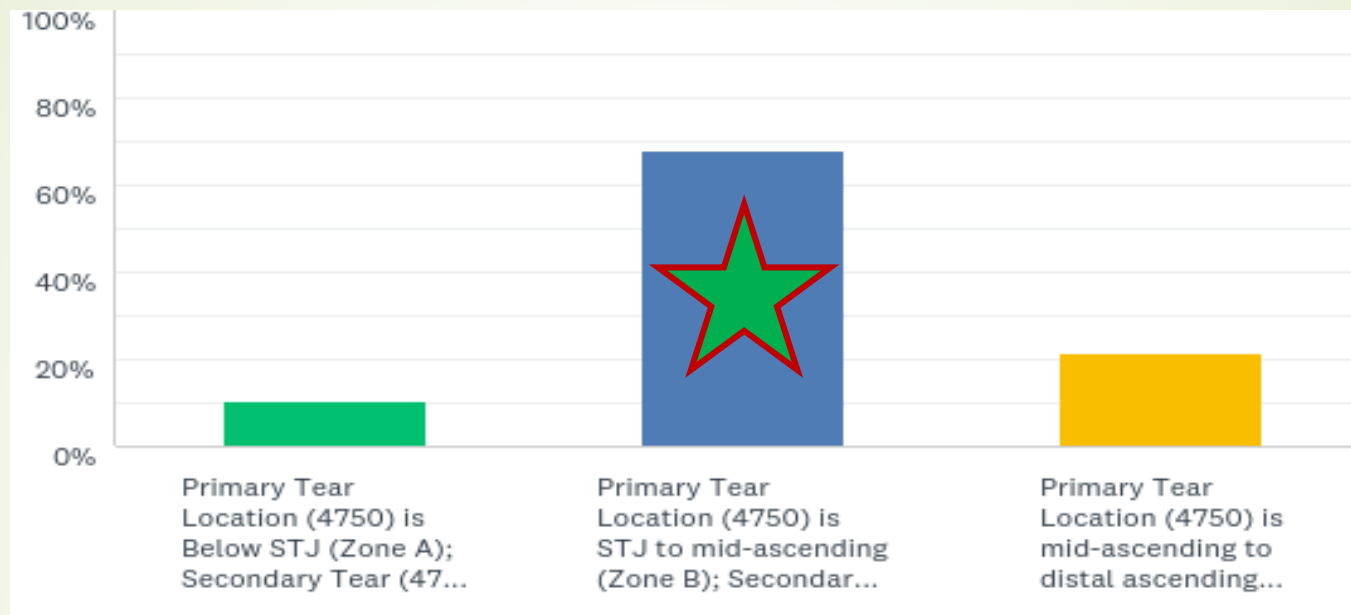
- Below STJ  STJ-midascending  Midascending to distal ascending  
 Zone 1  Zone 2  Zone 3  Zone 4  Zone 5  Zone 6  Zone 7  Zone 8  Zone 9  Zone 10  Zone 11

Secondary tear location:

DisSecLoc (4755)

- Below STJ  STJ-midascending  Midascending to distal ascending  
 Zone 1  Zone 2  Zone 3  Zone 4  Zone 5  Zone 6  Zone 7  Zone 8  Zone 9  Zone 10  Zone 11

## #10: Based on the CT scan results, how would you abstract the following fields in Section M2?



### Answer Choices

Primary Tear Location (4750) is Below STJ (Zone A); Secondary Tear (4755) is Zone 6.

Primary Tear Location (4750) is STJ to mid-ascending (Zone B); Secondary Tear (4755) is Zone 5.

Primary Tear Location (4750) is mid-ascending to distal ascending (Zone C); Secondary Tear (4755) is Zone 5.

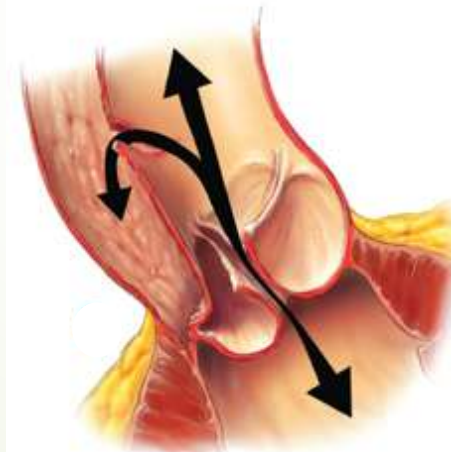
## #11: Based on the CT scan results, how would you abstract these subsequent fields?

### Answer Choices:

- Retrograde extension (4760) is Yes; Retrograde Location (4765) is Below STJ; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.
- Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 5.
- Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.

### Points to Consider:

- Retrograde Extension
- Retrograde Location
- Distal Extension
- Distal Location





**SEQ. #:** 4760

**Long Name:** Dissection - Retrograde Extension **Short Name:** DisRetExt

**Definition:** Indicate whether there was retrograde extension

**Intent/Clarification:**

The intent is to determine whether the dissection propagates proximal (toward the aortic valve) from the primary tear location. Report yes if imaging indicates an extension of the false lumen proximal (toward the aortic valve) to the primary tear location.

-----  
-----

**SEQ. #:** 4765

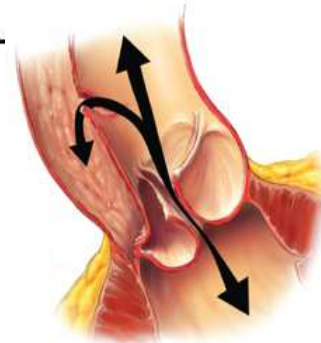
**Long Name:** Dissection - Retrograde Location

**Short Name:** DisRetLoc

**Definition:** Indicate location of retrograde extension

**Intent/Clarification:**

The intent is to define how far the retrograde dissection extends toward the aortic valve. This would be the point at which the false lumen comes closest to the aortic valve. The surgeon or radiologist can be the final arbiter of this definition. Refer to the image showing the zones and note that zone "0" is subdivided into 3 sections:





**SEQ. #: 4775**

**Long Name:** Dissection - Distal Extension

**Short Name:** DistalExt

**Definition:** Indicate whether there is distal extension

**Intent/Clarification:**

The intent is to identify where distal (antegrade) dissection occurred or extended.

**SEQ. #: 4780**

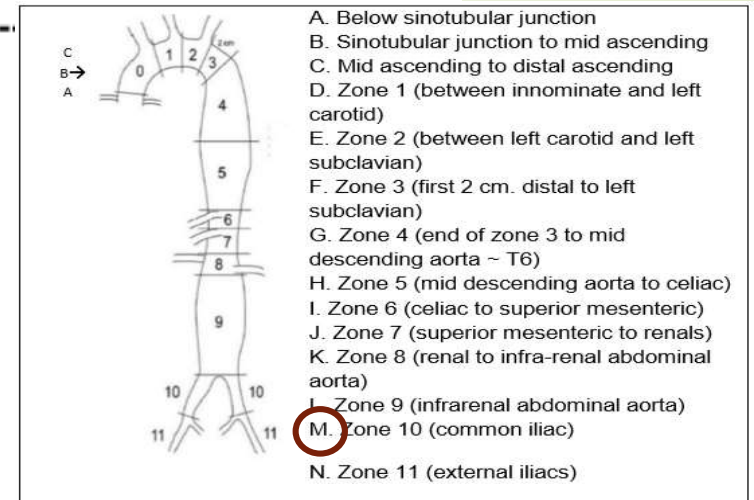
**Long Name:** Dissection - Distal Extension Location

**Short Name:** DistalExtLoc

**Definition:** Indicate location of distal extension

**Intent/Clarification:**

The intent is to define the how far along the aorta (away from the valve) any new or extended dissection goes. Refer to the image showing the zones and report the most distal (highest # zone) extent of the false lumen.



# #11: Based on the CT scan results, how would you abstract these subsequent fields?

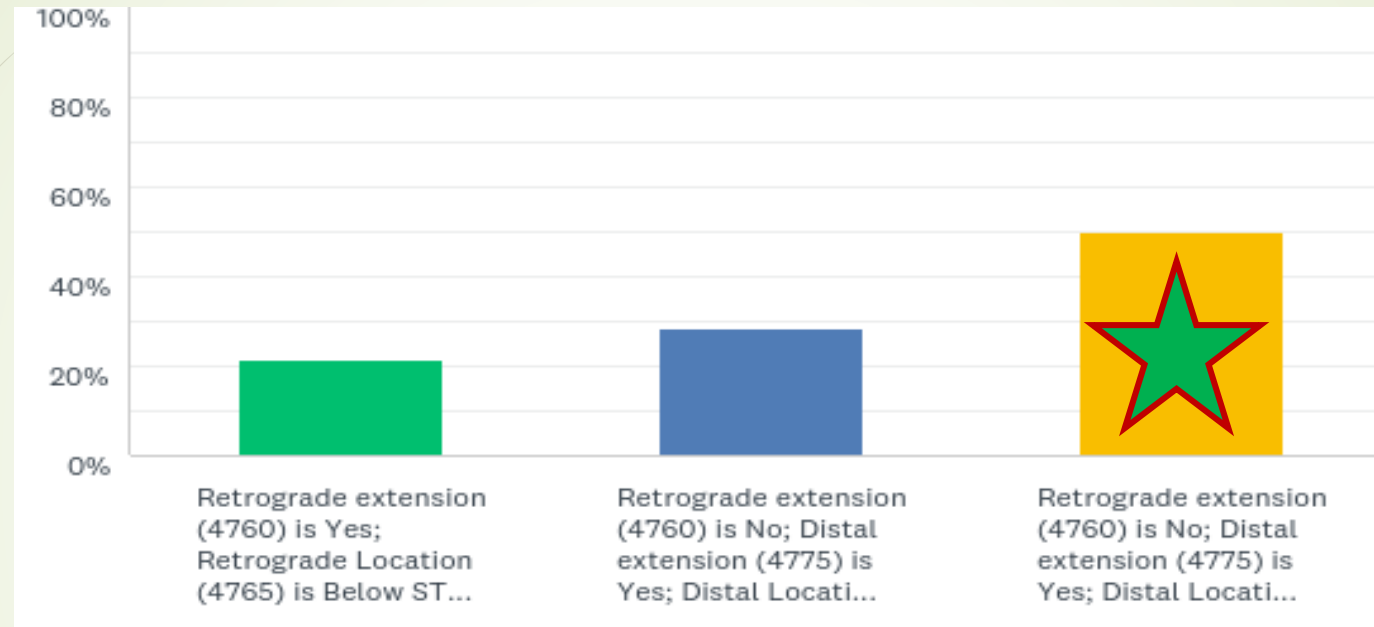
## Answer Choices:

- Retrograde extension (4760) is Yes; Retrograde Location (4765) is Below STJ; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.
- Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 5.
- Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.

Primary tear location: <b>DisTearLoc (4750)</b>	<input type="checkbox"/> Below STJ <input type="checkbox"/> STJ-midascending <input type="checkbox"/> Midascending to distal ascending <input type="checkbox"/> Zone 1 <input type="checkbox"/> Zone 2 <input type="checkbox"/> Zone 3 <input type="checkbox"/> Zone 4 <input type="checkbox"/> Zone 5 <input type="checkbox"/> Zone 6 <input type="checkbox"/> Zone 7 <input type="checkbox"/> Zone 8 <input type="checkbox"/> Zone 9 <input type="checkbox"/> Zone 10 <input type="checkbox"/> Zone 11
Secondary tear location: <b>DisSecLoc (4755)</b>	<input type="checkbox"/> Below STJ <input type="checkbox"/> STJ-midascending <input type="checkbox"/> Midascending to distal ascending <input type="checkbox"/> Zone 1 <input type="checkbox"/> Zone 2 <input type="checkbox"/> Zone 3 <input type="checkbox"/> Zone 4 <input type="checkbox"/> Zone 5 <input type="checkbox"/> Zone 6 <input type="checkbox"/> Zone 7 <input type="checkbox"/> Zone 8 <input type="checkbox"/> Zone 9 <input type="checkbox"/> Zone 10 <input type="checkbox"/> Zone 11
Retrograde extension: <b>DisRetExt (4760)</b>	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes ↓)
Retrograde Location: <b>DisRetLoc (4765)</b>	<input type="checkbox"/> Below STJ <input type="checkbox"/> STJ-midascending <input type="checkbox"/> Midascending to distal ascending <input type="checkbox"/> Zone 1 <input type="checkbox"/> Zone 2 <input type="checkbox"/> Zone 3 <input type="checkbox"/> Zone 4
Post TEVAR: <b>DisPosTEVAR (4770)</b>	<input type="checkbox"/> Yes <input type="checkbox"/> No
Distal extension: <b>DistalExt (4775)</b>	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown (If Yes ↓)
Distal Extension Location: <b>DistalExtLoc (4780)</b>	<input type="checkbox"/> Below STJ <input type="checkbox"/> STJ-midascending <input type="checkbox"/> Midascending to distal ascending <input checked="" type="checkbox"/> Zone 1 <input type="checkbox"/> Zone 2 <input type="checkbox"/> Zone 3 <input type="checkbox"/> Zone 4 <input type="checkbox"/> Zone 5 <input type="checkbox"/> Zone 6 <input type="checkbox"/> Zone 7 <input type="checkbox"/> Zone 8 <input type="checkbox"/> Zone 9 <input checked="" type="checkbox"/> Zone 10 <input type="checkbox"/> Zone 11

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# #11: Based on the CT scan results, how would you abstract these subsequent fields?



## Answer Choices

Retrograde extension (4760) is Yes; Retrograde Location (4765) is Below STJ; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.

Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 5.

Retrograde extension (4760) is No; Distal extension (4775) is Yes; Distal Location (4780) is Zone 10.

## #12: What sort of Open Arch Procedure (4975) did this patient undergo?

### Answer Choices:

- Distal Technique (4980) is Open; Distal Site (4985) is Hemi-Arch; Distal Extension (4990) is Elephant Trunk; Arch Branch Re-implantation (4995) is Yes.
- Distal Technique (4980) is Clamped; Distal Site (4985) is Zone 3; Distal Extension (4990) is Frozen Elephant Trunk; Arch Branch Re-implantation (4995) is No.
- Distal Technique (4980) is Open; Distal Site (4985) is Hemi-Arch; Distal Extension is No; Arch Branch Re-implantation (4995) is No.

### Points to Consider:

- Distal Technique
- Distal Site
- Distal Extension
- Arch Branch Re-Implantation

### Remember:

- Proximal = *Toward* the Heart
- Distal = *Away* from the Heart



**SEQ. #:** 4975

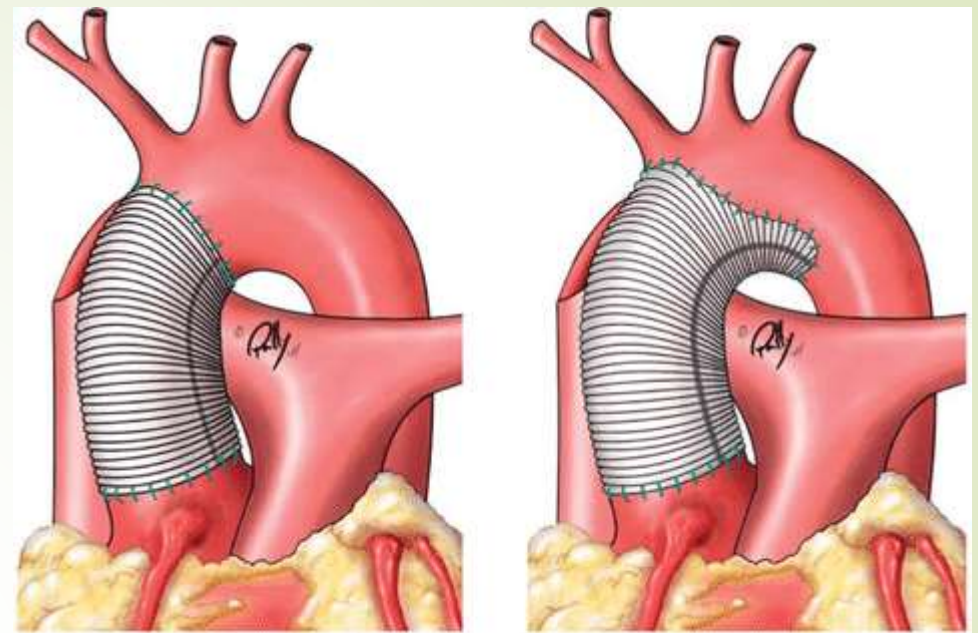
**Long Name:** Open Arch Procedure

**Short Name:** ArchProc

**Definition:** Indicate whether there was an open arch procedure

**Intent/Clarification:**

The intent is to identify procedures with replacement of or connection to the arch of the aorta. Anything from the base of the innominate through the subclavian takeoff would be included.



**a** Supracommissural replacement

**b** Hemiarch replacement

**SEQ. #:** 4980

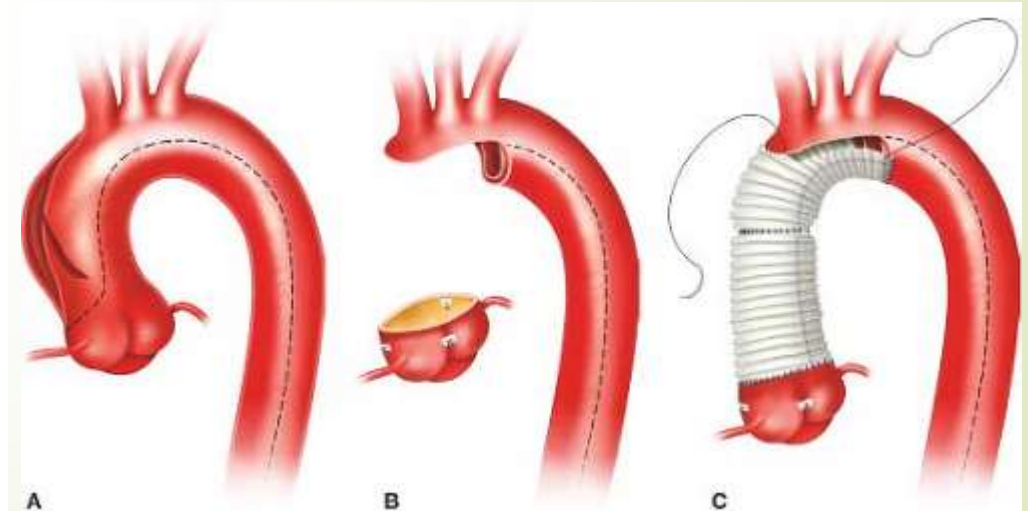
**Long Name:** Open Arch Procedure - Distal Technique

**Short Name:** ArchDisTech

**Definition:** Indicate the distal technique for the arch procedure

**Intent/Clarification:**

The intent is to define that the distal anastomosis was done with or without a clamp. Many arch procedures are done with the clamp removed, sewing to the aorta looking down the barrel of the vessel. This of course requires circulatory arrest. The clamp means that the aorta is clamped with an instrument and the anastomosis is completed proximal (close to the heart) to that part of the aorta.



**A**

**B**

**C**



SEQ. #: 4985

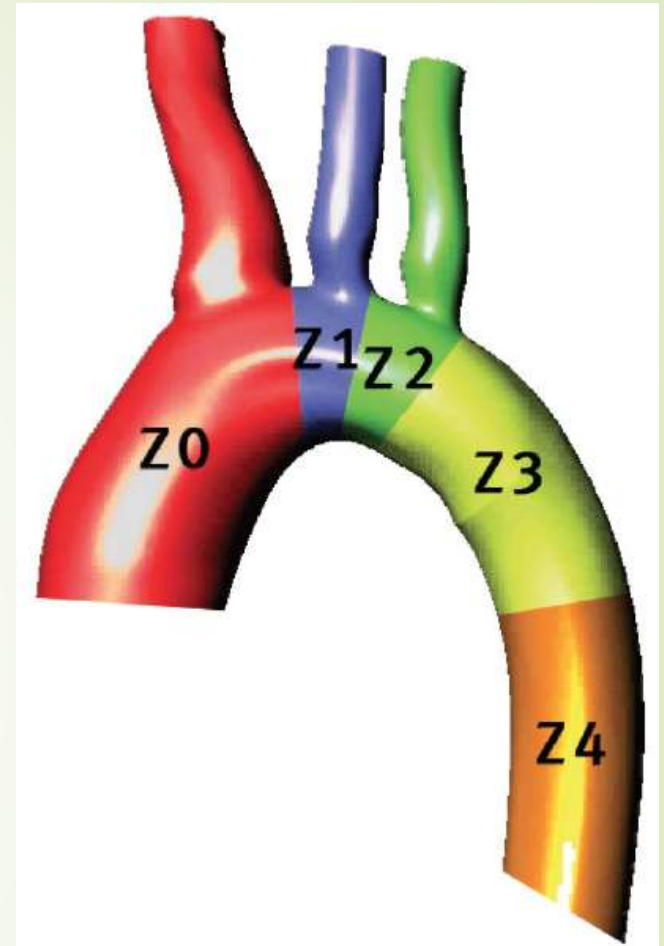
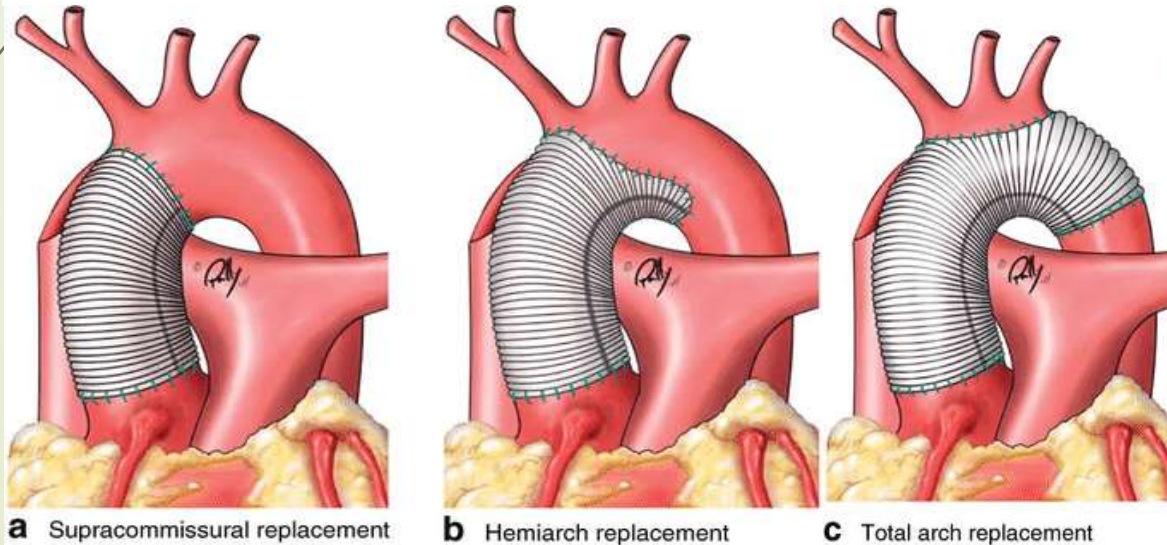
Long Name: Open Arch Procedure - Distal Site

Short Name: ArchDiscSite

Definition: Indicate the distal site

**Intent/Clarification:**

The intent of this is to define the level of the distal (far from the heart) anastomosis. Ascending aorta implies the ascending was resected with a clamp on the distal ascending aorta. Hemiarch means a single anastomosis was done somewhere in the ascending or proximal arch without separate grafts to the head vessels. Zone 1 means the innominate was reconnected with a graft between the innominate and left common carotid takeoffs. Zone 2 means the innominate and carotid were reconnected with a graft sewn to between the left common carotid and the left subclavian takeoffs. Zone three means the innominate, carotid and the left subclavian were reconnected with the graft being sewn beyond the left subclavian takeoff. Zone 4 means the graft was sewn to the mid descending thoracic aorta.



## Intervention

Planned Staged Hybrid:  Yes  No

*PlanStagHybrid (4970)*

Open Arch Procedure:  Yes  No (If Yes ↓)

*ArchProc (4975)*

Distal Technique:  Open  Clamped

*ArchDisTech (4980)*

Distal Site:  Ascending Aorta  Hemiarch  Zone 1  Zone 2  Zone 3  Zone 4

*ArchDiscSite (4985)*

Distal Extention:  Elephant trunk  Frozen Elephant trunk  No

*ArchDisExt (4990)*

Arch Branch Reimplantation:  Yes  No (If Yes ↓)

*ArchBranReimp (4995)*

Innominate:  Yes  No

*ArchBranInnom (5000)*

Right Subclavian:  Yes  No

*ArchBranRSub (5001)*

Right Common Carotid:  Yes  No

*ArchBranRComm (5002)*

Left Common Carotid:  Yes  No

*ArchBranLComm (5005)*

Left Subclavian:  Yes  No

*ArchBranLSub (5010)*

Left Vertebral:  Yes  No

*ArchBranLVert (5011)*

Other:  Yes  No

*ArchBranOth (5012)*

**SEQ. #: 4990**

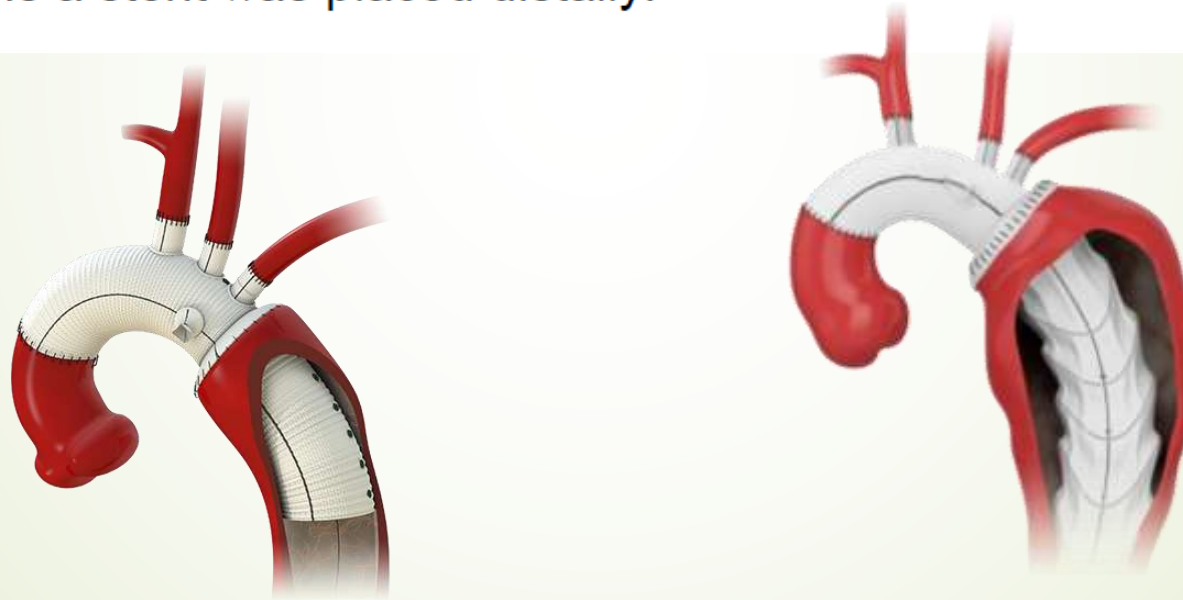
**Long Name:** Open Arch Procedure - Distal Extention

**Short Name:** ArchDisExt

**Definition:** Indicate distal extension type

**Intent/Clarification:**

The intent of the question is to define whether graft was left that extended (distally) beyond the arch anastomosis. An elephant trunk is a soft graft, while a frozen elephant trunk means a stent was placed distally.



**SEQ. #: 4995**

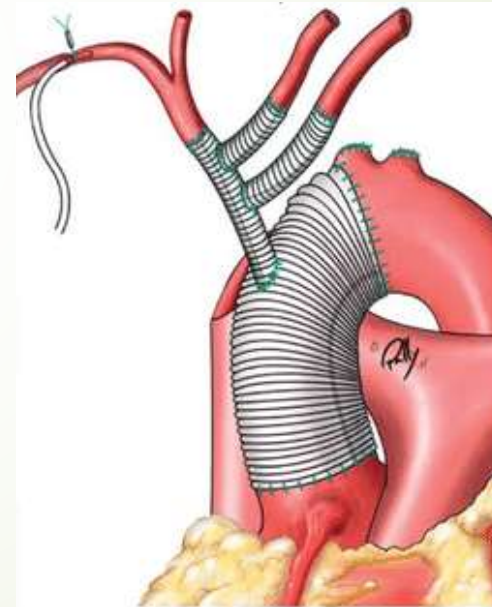
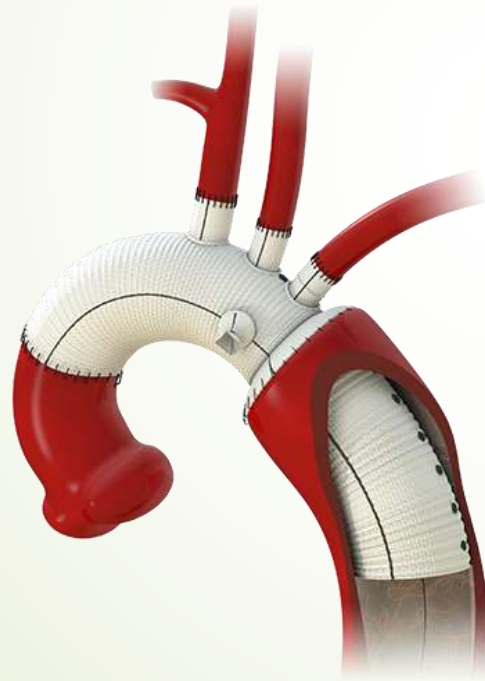
**Long Name:** Open Arch Procedure - Arch Branch Reimplantation

**Short Name:** ArchBranReimp

**Definition:** Indicate whether arch branch reimplantation was performed

**Intent/Clarification:**

The intent of this is to define the end branches that were sewn to the graft.



**d** Trifurcated graft

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## Intervention

Planned Staged Hybrid:  Yes  No

PlanStagHybrid (4970)

Open Arch Procedure:  Yes  No (If Yes ↓)

ArchProc (4975)

Distal Technique:  Open  Clamped

ArchDisTech (4980)

Distal Site:  Ascending Aorta  Hemiarch  Zone 1  Zone 2  Zone 3  Zone 4

ArchDiscSite (4985)

Distal Extention:  Elephant trunk  Frozen Elephant trunk  No

ArchDisExt (4990)

Arch Branch Reimplantation:  Yes  No (If Yes ↓)

ArchBranReimp (4995)

Innominate:  Yes  No

ArchBranInnom (5000)

Right Subclavian:  Yes  No

ArchBranRSub (5001)

Right Common Carotid:  Yes  No

ArchBranRComm (5002)

Left Common Carotid:  Yes  No

ArchBranLComm (5005)

Left Subclavian:  Yes  No

ArchBranLSub (5010)

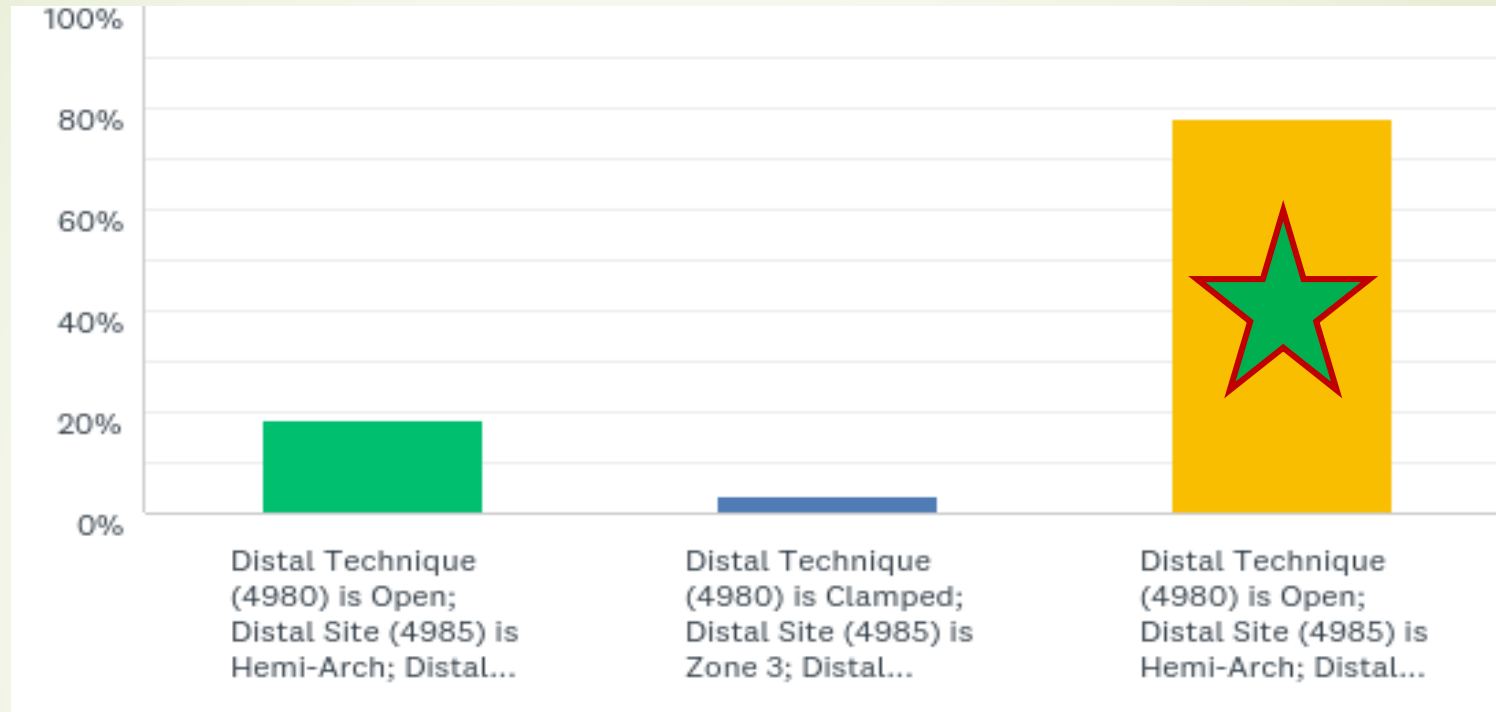
Left Vertebral:  Yes  No

ArchBranLVert (5011)

Other:  Yes  No

ArchBranOth (5012)

# #12: What sort of Open Arch Procedure (4975) did this patient undergo?



Answer Choices
Distal Technique (4980) is Open; Distal Site (4985) is Hemi-Arch; Distal Extension (4990) is Elephant Trunk; Arch Branch Re-implantation (4995) is Yes.
Distal Technique (4980) is Clamped; Distal Site (4985) is Zone 3; Distal Extension (4990) is Frozen Elephant Trunk; Arch Branch Re-implantation (4995) is No.
Distal Technique (4980) is Open; Distal Site (4985) is Hemi-Arch; Distal Extension is No; Arch Branch Re-implantation (4995) is No.

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## #13: How would you abstract the woven graft implanted during the procedure?

### Answer Choices:

- **Device Inserted (5440) is Yes; Location (5450) is Zone 1; Delivery Method (5455) is Open.**
- **Device Inserted (5440) is Yes; Location (5450) is Zone A; Delivery Method (5455) is Open.**
- **Device Inserted (5440) is Yes; Location (5450) is Zone B; Delivery Method (5455) is Open.**
- **Sorry, but I'm "Zoned Out"!!!!!!**

### Points to Consider:

- **Was a Device Inserted?**
- **Location of Device**
- **Delivery Method**



**SEQ. #: 5440**

**Long Name:** Aorta Device Inserted

**Short Name:** ADevIns

**Definition:** Indicate whether one or more devices were inserted into the aorta.

**Intent/Clarification:**

This will include all synthetic prosthetics inserted. This may include Dacron, PTFE, homografts, autografts, stents, and stentgrafts. Some aortic interventions may not require prosthetic materials or device implants such as primary repair of a pseudoaneurysm. This will be indicated as “No.”

**SEQ. #: 5450**

**Long Name:** Aorta Device - Location #01

**Short Name:** ADevLoc01

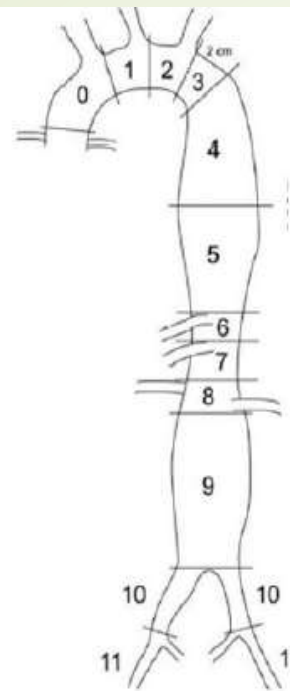
**Definition:** Indicate the location within the aorta where device #01 was inserted.

**Intent/Clarification:**

Zone 0 is the Ascending Aorta and includes letter A-C. Verify exact location with CV Surgeon. Aortic Root (letter A) is below sinotubular junction.



**Location :**

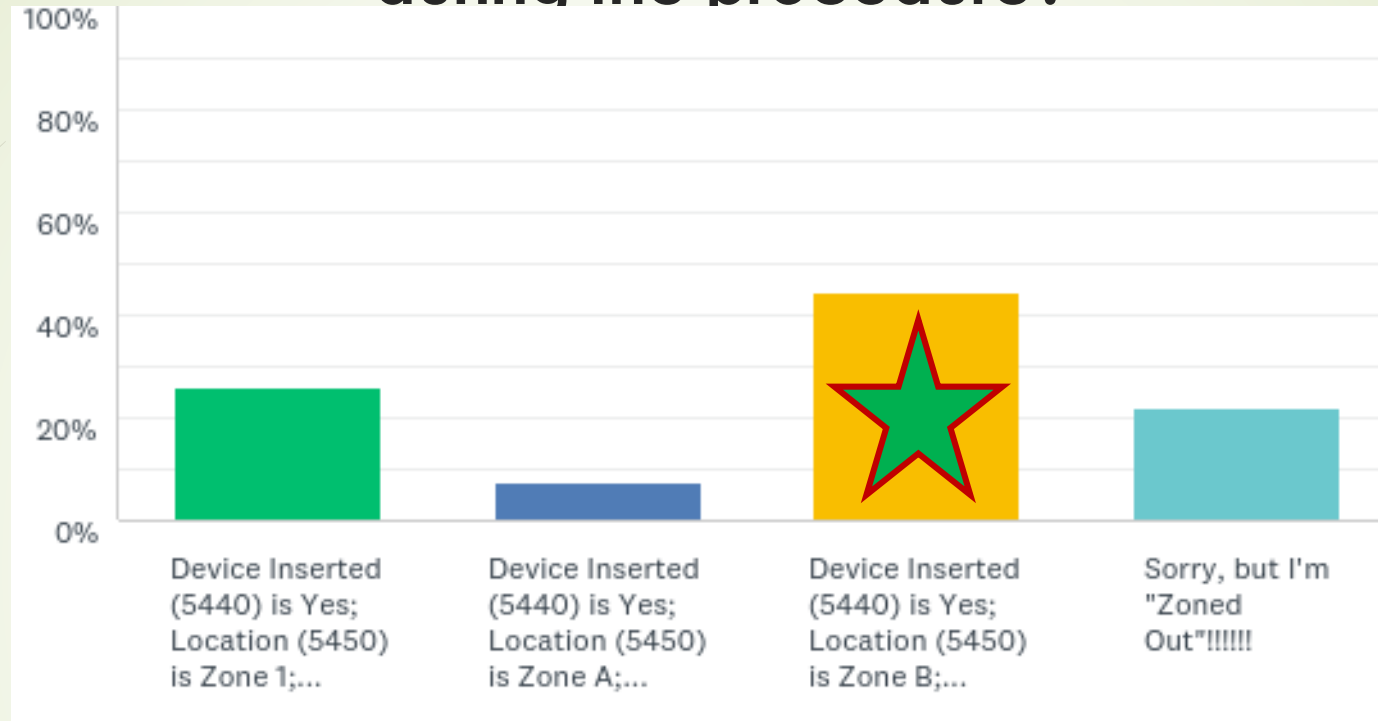


- X. No additional devices inserted (only for locations 2 – 15)
- A. Below sinotubular junction
- B. Sinotubular junction to mid ascending
- C. Mid ascending to distal ascending
- D. Zone 1 (between innominate and left carotid)
- E. Zone 2 (between left carotid and left subclavian)
- F. Zone 3 (first 2 cm. distal to left subclavian)
- G. Zone 4 (end of zone 3 to mid descending aorta ~ T6)
- H. Zone 5 (mid descending aorta to celiac)
- I. Zone 6 (celiac to superior mesenteric)
- J. Zone 7 (superior mesenteric to renals)
- K. Zone 8 (renal to infra-renal abdominal aorta)
- L. Zone 9 (infrarenal abdominal aorta)
- M. Zone 10 (common iliac)
- N. Zone 11 (external iliacs)

<b>Delivery Method:</b>	1=Open 2= Endovascular			
<b>Outcome:</b>	1= Maldeployed 2= Deployed and removed 3= Successfully deployed			
<b>Model Number:</b>	Enter device model number			
<b>UDI:</b>	Enter unique device identifier (not serial number)			
<b>Location (Letter)</b>	<b>Delivery Method</b>	<b>Outcome</b>	<b>Model #</b>	<b>UDI</b>
<b>B</b>	<b>1</b>	<b>Leave Blank</b>		
ADevLoc02 (5475)	ADevDelMeth02 (5480)	ADevOut02 (5485)	ADevModel02 (5490)	ADevUDI02 (5495)

**Currently unclear about where an associated valve device will be recorded.**

# #13: How would you abstract the woven graft implanted during the procedure?



Answer Choices
Device Inserted (5440) is Yes; Location (5450) is Zone 1; Delivery Method (5455) is Open.
Device Inserted (5440) is Yes; Location (5450) is Zone A; Delivery Method (5455) is Open.
Device Inserted (5440) is Yes; Location (5450) is Zone B; Delivery Method (5455) is Open.
Sorry, but I'm "Zoned Out"!!!!!!

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## #14: How would you abstract the following Discharge (Section Q) fields?

### Answer Choices:

- **Date of Last Follow-up (7000) is Discharge Date; Verification of 30-day Status (7002) is Phone call to/from patient or family; Discharge/Mortality Status (7005) is Discharged alive, last known status is alive.**
- **Date of Last Follow-up (7000) is Mortality Date; Verification of 30-day Status (7002) is Phone call/letter to/from medical provider; Discharge/Mortality Status (7005) is Discharged alive, died after discharge.**
- **Date of Last Follow-up (7000) is Discharge Date; Verification of 30-day Status (7002) is Phone call/letter to/from medical provider; Discharge/Mortality Status (7005) is Discharged alive, died after discharge.**

### Points to Consider:

- **Date of Last Follow-up**
- **Verification of 30-day Status**
- **Discharge/Mortality Status**



**From:** Joseph E. Bavaria, MD [<mailto:jbavaria@sts.org>]  
**Sent:** Thursday, June 02, 2016 4:48 PM  
**To:** STS Data Managers  
**Subject:** Data Completeness Requirements for Star Ratings

Dear STS Data Manager & Participants,

As a participant in the STS National Database, you know the importance of good data. This importance is underscored by the contractual obligation of each participant to submit complete and accurate data to the Database. Data analysis, risk adjustment, measure development, and nationally benchmarked results all rely on these high data standards.

Although in-hospital mortality data are recorded with high completeness and fidelity, it has come to our attention that some programs often choose "unknown" as the response for 30-day status, which may impact the accuracy of operative mortality determinations. Therefore, in order to assure the highest level of accuracy when reporting operative mortality, the following data thresholds are being implemented to determine eligibility for a composite score (star rating):

1. Cases performed January 1–December 31, 2015 must have a **90% completeness threshold** for fields related to operative mortality status.
2. For all cases performed on or after January 1, 2016, the operative mortality fields must have a **95% completeness rate**.
3. For all cases performed on or after January 1, 2017, the operative mortality fields must have a **98% completeness rate**.

If the following fields are missing or coded as "unknown", then the record will be considered incomplete. Going forward, participants who do not meet the mortality-related data completeness thresholds for a particular harvest will not be eligible to receive a composite score (star rating).

**Adult Cardiac Surgery Database v2.81**

Discharge Status (5010)  
Status at 30 Days after Discharge (5015)  
Operative Death (5025)

**Congenital Heart Surgery Database v3.3**

Mortality Status at Hospital discharge (4230)  
Mortality Status at Database Discharge (4250)  
Status at 30 days after surgery (4300)

**General Thoracic Surgery Database v2.3**

Discharge Status (2200)  
Status at 30 days after surgery (2240)

**January, 2017:**  
**Operative Mortality fields > 98%**  
**completeness rate:**

- **Discharge Status**
- **Status at 30 days**
- **Operative Death**

**< 98% = ineligible to receive  
Composite Score (star rating)**

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**SEQ. #: 7000**

**Long Name:** Date of Last Follow-Up

**Short Name:** LFUDate

**Definition:** Indicate the date on which the last follow-up was made. If patient dies in the hospital, this value will be the same as the date of death. If no follow-up is made after patient is discharged, this value will be the same as the discharge date.

**Intent/Clarification:**

This is the date that is last documented in the chart or obtained by contacting the physician's office. Required date format: mm/dd/yyyy

**SEQ. #: 7001**

**Long Name:** Mort-30d Status

**Short Name:** Mt30Stat

**Definition:** Indicate whether the patient was alive or dead at 30 days post-surgery (whether in hospital or not).

**Intent/Clarification:**

- Alive
- Dead
- ~~Unknown~~

**SEQ. #: 7002**

**Long Name:** Mort-Op Death-Method Of Verification

**Short Name:** Mt30StatMeth

**Definition:** Indicate the primary method used to verify the patient's 30-day mortality status.

---

**Intent/Clarification:**

- Phone call to patient or family
- Letter from medical provider
- Evidence of life or death in medical record
- Office visit on or after 30 days after the date of surgery.
- Social Security Death Master File/NDI
- Other

**SEQ. #: 7005**

**Long Name:** Discharge / Mortality Status

**Short Name:** DischMortStat

**Definition:** Indicate the discharge and current vital status of the patient

**Intent/Clarification:**

- In hospital, alive
- Died in hospital
- Discharged alive, last known status is alive
- Discharged alive, died after discharge

"In hospital, alive" refers to patient's that are in the hospital at the 30 day mark that were never discharged. It is provided so sites do not get marked as missing on the required mortality fields for their composite scores/STAR ratings.

## Q. Discharge / Mortality

Date of Last Follow-up: \_\_\_/\_\_\_/\_\_\_ (mm/dd/yyyy)

**Mortality Date**

LFUDate (7000)

Status at 30 days After Surgery:  Alive  Dead  Unknown

Mt30Stat (7001)

Primary method used to verify 30-day status:

Phone call to patient or family

Office visit  $\geq$  30 days after procedure

Letter from medical provider

Social Security Death Master File /NDI

Medical record (evidence of life or death)

Other

Discharge/Mortality status:  In hospital, alive  Discharged alive, last known status = alive

DischMortStat (7005)

Died in hospital  Discharged alive, died after discharge

# #14: How would you abstract the following Discharge (Section Q) fields?



## Answer Choices

Date of Last Follow-up (7000) is Discharge Date; Verification of 30-day Status (7002) is Phone call to/from patient or family; Discharge/Mortality Status (7005) is Discharged alive, last known status is alive.

Date of Last Follow-up (7000) is Mortality Date; Verification of 30-day Status (7002) is Phone call/letter to/from medical provider; Discharge/Mortality Status (7005) is Discharged alive, died after discharge.

Date of Last Follow-up (7000) is Discharge Date; Verification of 30-day Status (7002) is Phone call/letter to/from medical provider; Discharge/Mortality Status (7005) is Discharged alive, died after discharge.



## #15: Finally, how would you abstract these Mortality (Section Q) fields?

### Answer Choices:

- Primary cause of Death (7122) is Cardiac; Operative Death (7124) is No; Discharge Death Location (7125) is Acute Rehabilitation.
- Primary cause of Death (7122) is Cardiac; Operative Death (7124) is Yes; Discharge Death Location (7125) is Acute Rehabilitation.
- Primary cause of Death (7122) is Cardiac; Operative Death (7124) is Yes; Discharge Death Location (7125) is Extended Care Facility.

### Points to Consider:

- Primary Cause of Death
- Was this an Operative Death?
- Discharge Death Location

**SEQ. #: 7122**

**Long Name:** Mort-Prim Cause

**Short Name:** MtCause

**Definition:** Indicate the PRIMARY cause of death, i.e., the first significant abnormal event which ultimately led to death.

**Intent/Clarification:** If the patient died due to multiple organ system failure, select the system that either was the initiator of the Multisystem Organ Failure (MSOF) or the primary cause of the patient's demise.

- Cardiac
- Neurologic
- Renal
- Vascular
- Infection
- Pulmonary
- Unknown
- Other

**SEQ. #: 7124**

**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.

SEQ. #: 7125

Long Name: Post Discharge Death Location

Short Name: PostDisDthLoc

Definition: Indicate the location where the patient died after being discharged from the original hospitalization.

Intent/Clarification:

- **Home** (or, temporarily, at the home of a relative)
- **Extended Care Facility/Transitional Care Unit (TCU)** (Code LTAC as Extended Care/Transitional Care Unit/Rehab. Do not count as part of acute care stay.
- **Hospice**
- **Acute Rehabilitation** (Ultimate plan for patient to return home after a short-stay)
- **Hospital, During Readmission**
- **Other**
- **Unknown**

(If Discharge/Mortality Status = "Died in hospital" or "Discharged alive, died after discharge" ↓)

Mortality - Date \_\_\_/\_\_\_/\_\_\_\_\_ (mm/dd/yyyy)

MtDate (7121)

Primary Cause of Death (select only one)

MtCause (7122)

Cardiac  Neurologic  Renal  Vascular  Infection  Pulmonary  Unknown  Other

(If Discharge/Mortality Status = "Died in hospital" ↓)

In-Hospital death location:  OR During Initial Surgery  OR during reoperation  In Hospital (Other than OR)

InHospDthLoc (7123)

(If Discharge/Mortality Status = "Discharged alive, died after discharge" )

Operative Death:  Yes  No

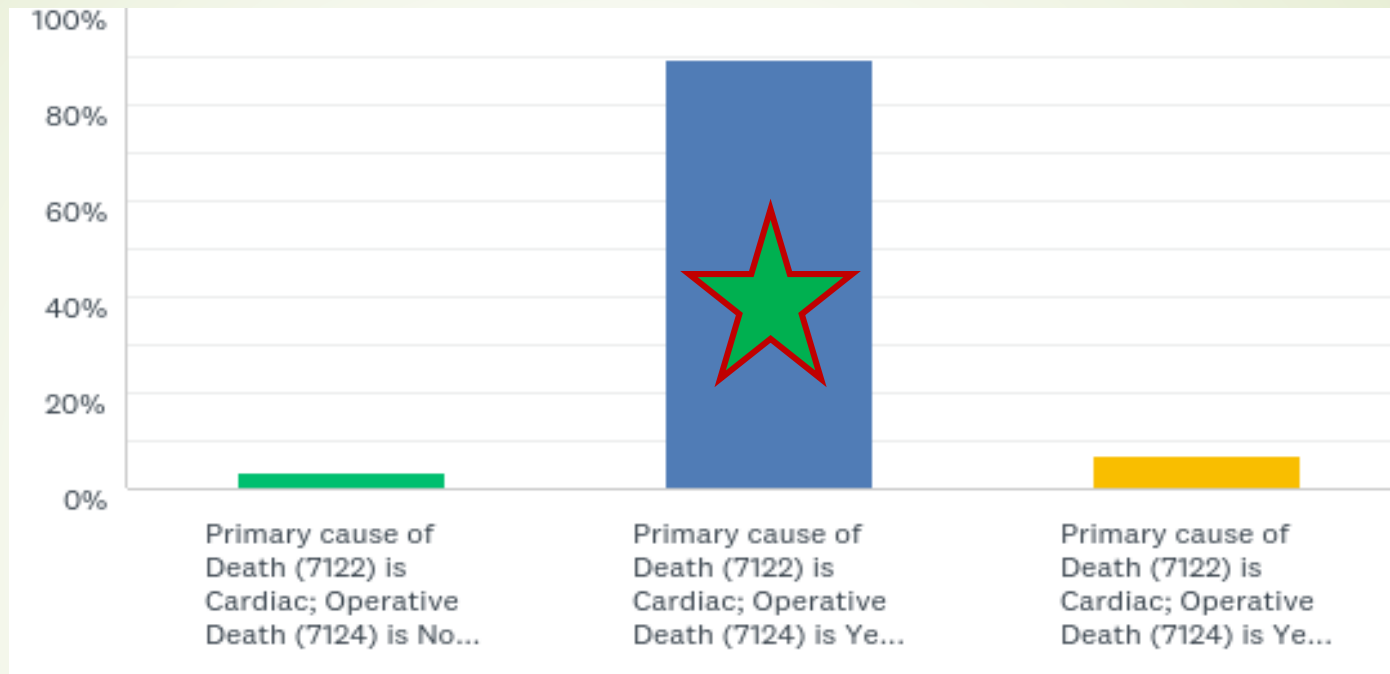
MtOpD (7124)

Post Discharge death location:

PostDisDthLoc (7125)

Home  Extended Care Facility  Hospice  Acute Rehabilitation  Hospital during readmission  Other  Unknown

# #15: Finally, how would you abstract these Mortality (Section Q) fields?



Answer Choices
Primary cause of Death (7122) is Cardiac; Operative Death (7124) is No; Discharge Death Location (7125) is Acute Rehabilitation.
Primary cause of Death (7122) is Cardiac; Operative Death (7124) is Yes; Discharge Death Location (7125) is Acute Rehabilitation.
Primary cause of Death (7122) is Cardiac; Operative Death (7124) is Yes; Discharge Death Location (7125) is Extended Care Facility.

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