STS General Thoracic Surgery Database Version 2.2 Update May 2014

Fields with questions, clarifications, examples are in red: 620, 730, 760, 770, 790, 800, 820, 840, 930, 1140, 1250, 1300, 1320, 1420, 1480, 1500, 1610, 1710, 1810, 1830, 1860, 1910, 1970, 1990, 2020, 2040, 2090, 2150, 2230, 2240, 2340, 2300

GENERAL INFORMATION

<u>July 2013</u> - The CPT codes provided on the DCF are for reference & information. The General Thoracic Surgery Database is a clinical database and for its purposes, the correct **clinical category** should be captured, not the CPT code. The CPT code a billing coder may use may not be correct for the purposes of a clinical database.

<u>October 2013</u> – Non Analyzed Procedures (NAP) - Can isolated NAPs or those performed with another NAP be submitted for harvest? If yes, what happens to these procedures? *These cases would not be analyzed or reported by DCRI in your harvest report even if submitted with your file. Your DQR would give you the # of procedures that fall into this category.*

October 2013 - Some participants keep track of their NAP cases & use them for internal queries and to track cases. Some surgeons want to track all cases and some programs want to track volumes and resource utilization. The GTSD does not track any of this information.

December 2013 - Addendums may be added to the medical record within 30 days to be consistent with CMS.

<u>April 2014</u> – INCLUSION: Would a thymectomy done during a CABG and MAZE be captured in the Thoracic database? It was an incidental finding when opening the sternum for the CABG. *No, it would be included in the ACSD as a "Other, non-cardiac-other." Sequence #5590.*

<u>April 2014</u> – When trying to establish which codes to use, keep the following in mind: *If it's done using a scope, use the Thoracoscopy codes. If it's an open procedure, use the thoracotomy codes.*

Version 2.2 was designed to focus on major, risk adjusted procedures. A noteworthy and time-saving difference to the new data specifications is that procedures which are not analyzed will no longer be mandatory to collect. For example, isolated non-analyzed procedures which are not associated with major procedures during the same anesthesia, such as bronchoscopy, are now optional.

For STS purposes, such data won't be analyzed and data collection for these procedures will be at the surgeon's discretion. For those continuing to collect procedure data on all cases, a short Data Collection Form is available for the non-analyzed procedures to save time.

- A Major Procedure Data Collection Form (DCF) should be initiated every time the patient enters the Operating Room for Major Procedure(s). Major procedures are analyzed, may be risk adjusted and are included in Harvest Reports.
- Fields that appear underlined and in blue on the DCF are required for Major procedure record inclusion. If any of these fields are

Missing data, the entire record will be excluded from the analysis.

- Procedures highlighted on the DCF, if performed as isolated procedures or with another highlighted procedure are not collected unless the Surgeon Participant chooses to track them. If collected, use the data set highlighted on the DCF or the Non-analyzed Procedure Data Set DCF. Sections and Fields that appear highlighted are suggested for these procedures.
- Highlighted procedures done in conjunction with major procedures should be included on the Major Procedure DCF.

Use the training manual to clarify field definitions and intent. Submit clinical questions to: FAQs. You will receive an email response and questions and answers will be posted below in red.

Training Manual Version 2.2 of the General Thoracic Surgery Database				
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification
10	RecordID	Operations Table Record Identifier		
20	RecordID	Procedures Table Record Identifier		
30	VendorID	Software Vendor's Identification		
40	SoftVrsn	Vendor's Software Version Number		
50	DataVrsn	Version Of STS Data Specification		
60	ParticID	Participant ID	Participant ID is a unique number assigned to each Database participant by the STS. A Database participant is defined as one entity that signs a Participation Agreement with the STS, submits one data file to the harvest, and gets back one report on their data. The participant ID must be entered into each record.	Each participant's data, if submitted to harvest, must be in one data file. If one participant keeps data in more than one file (e.g. at two sites), the participant must combine them back into one file for harvest submission. If two or more participants share single purchased software and enter cases into one database, the data must be extracted into two different files, one for each participant ID, with each record having the correct participant ID

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				number.	
70	PatID	Operations Table Patient Identifier	If a patient is admitted to the hospital more than once, each record for that patient will have the same value in this field. A record should be initiated for inpatient and outpatient thoracic procedures on every visit to the operating room (includes the Endoscopy Suite or Outpatient Surgical Center) whether planned or unplanned.	Once assigned to a patient, this number can never be changed or reused.	
80	PatID	Demographics Table Patient Identifier	If a patient is admitted to the hospital more than once, each record for that patient will have the same value in this field. A record should be initiated for inpatient and outpatient thoracic procedures on every visit to the operating room (includes the Endoscopy Suite or Outpatient Surgical Center) whether planned or unplanned.		
90	DemogDataVrsn	Demographics Table Data Version			
100	MedRecN	Medical Record #	Indicate the patient's medical record number at the hospital where surgery occurred. This field should be collected in compliance with state/local privacy laws.		
110	PatFName	Patient's First Name	Indicate the patient's first name documented in the medical record.		
120	PatMInit	Patient's Middle Initial	Middle name or initial as documented in medical record	Leave "blank" if no middle initial.	
130	PatLName	Patient's Last Name	Indicate the patient's last name documented in the medical record.		
140	SSN	Social Security Number	Unique patient identifier assigned by government	Although this is the Social Security Number in the USA, other countries may have a different National Patient Identifier Number. For example in Canada, this would be the Social Insurance Number. The Social Security Number is crucial to provide linkage for long term follow up and every attempt should be made to collect it.	

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150	STSTLink	STS Trial Link Number	The unique identification number assigned by the STS indicating the clinical trial in which this patient is participating. This field should be left blank if the patient is not participating in a clinical trial associated with the STS		
160	DOB	Date Of Birth	Indicate the patient's date of birth using 4-digit format for year.		
170	Age	Age At Time Of Surgery	This field is Required for Record Inclusion, because it is part of the risk models. If missing data, the entire record will be excluded from the analysis.	Calculated value based on DOB and surgery date	
180	PostalCode	Zip Code	Indicate the ZIP Code, outside the USA, this data may be known by other names such as Postal Code (needing 6 characters). Software should allow sites to collect at least up to 10 characters to allow for Zip+4 values. This field should be collected in compliance with state/local privacy laws.		
190	Gender	Gender	Indicate the patient's gender at birth as either male or female. This field is Required for Record Inclusion and is used in Risk Models. If missing data, the entire record will be excluded from the analysis.	Patients who have undergone gender reassignment surgery maintain the risk associated with their chromosomal gender.	
N\A	Race		The race fields are Required for Record Inclusion and included in Risk Models. If missing data, the entire record will be excluded from the analysis. The Census Bureau collects race data in accordance with guidelines provided by the U.S. Office of Management and Budget and these data are based on <u>self-</u> <u>identification</u> . The racial categories included in the census form generally reflect a social definition of race recognized in this country, and are not an attempt to define race biologically, anthropologically or genetically. In addition, it is recognized that the categories of the race item include racial and national origin or socio-cultural groups. People may choose to report more than one race to indicate their racial	Select all that apply. People who identify their origin (ETHNICITY) as Hispanic, Latino or Spanish may be of any race. Reference: <u>www.whitehouse.gov/omb/fedreg/1997</u> <u>standards.html</u>	

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200	RaceCaucasian	Race - Caucasian	mixture, such as "American Indian and White." In addition, it is recognized that the categories of the race item include both racial and national origin and socio- cultural groups. You may choose more than one race category. Indicate the patient's race, as reported by the patient or		
200	RaceCadeasian	Race - Caucasian	family, includes White.		
210	RaceBlack	Race - Black / African American	Indicate whether the patient's race, as determined by the patient or family, includes Black/African American.	This includes a person having origins in any of the black racial groups of Africa. Terms such as "Haitian" or "Negro" can be used in addition to "Black or African American." Definition source: Standards for Maintaining, Collecting, and Presenting Federal Data on Race and Ethnicity: The minimum categories for data on race and ethnicity for Federal statistics, program administrative reporting, and civil rights compliance reporting.	
220	RaceAsian	Race - Asian	Indicate whether the patient's race, as determined by the patient or family, includes Asian.	This includes a person having origins in any of the original peoples of the Far East, Southeast Asia, or the Indian subcontinent including, for example, Cambodia, China, India, Japan, Korea, Malaysia, Pakistan, the Philippine Islands, Thailand, and Vietnam. Definition source: Standards for Maintaining, Collecting, and Presenting Federal Data on Race and Ethnicity: The minimum categories for data on race and ethnicity for Federal statistics, program administrative reporting, and	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				civil rights compliance reporting.	
230	RaceNativeAm	Race - American Indian /	Indicate whether the patient's race, as determined by the	Includes all in North American native	
		Alaskan Native	patient or family, includes Native American.	peoples such as American	
				Indian/Alaskan Native, Inuit.	
240	RacNativePacific	Race - Native Hawaiian /	Indicate whether the patient's race, as determined by the	This includes a person having origins in	
		Pacific Islander	patient or family, includes Native Hawaiian/Pacific	any of the original peoples of Hawaii,	
			Islander.	Guam, Samoa, or other Pacific Islands.	
				Definition source: Standards for	
				Maintaining, Collecting, and Presenting	
				Federal Data on Race and Ethnicity:	
				The minimum categories for data on	
				race and ethnicity for Federal statistics,	
				program administrative reporting, and	
250		D OI		civil rights compliance reporting.	
250	RaceOther	Race Other	Indicate whether the patient's race, as determined by the		
			patient or family, includes any other race.		
260	RaceUnk	Race Unknown	Indicate if the patients race is unknown	Make every attempt to collect race since	
			-	it is part of Risk Models.	
270	Ethnicity	Hispanic Or Latino Ethnicity	Indicate if the patient is of Hispanic, Latino or Spanish	Hispanic, Latino or Spanish ethnicity	
			ethnicity as reported by the patient/family.	includes patient report of Cuban,	
				Mexican, Puerto Rican, South or	
				Central American, or other Spanish	
				culture or origin, regardless of race .	
				People who identify their origin as	
				Hispanic, Latino or Spanish may be of	
				any race.	
280	AdmissionStat	Admission Status	Indicate whether the procedure was an Inpatient or	Outpatient/Observation should be	
			Outpatient/Observation procedure.	selected if the operation was performed	
			This field is Required for Record Inclusion. If missing	as an ambulatory procedure or if it	
			data, the entire record will be excluded from the	included a period of overnight	
200			analysis.	observation.	
290	AdmitDt	Admission Date	Indicate the date of admission. For those patients who	For purposes of this data definition,	
			originally enter the hospital in an outpatient capacity,	Outpatient and Observation status are	
			the admit date is the date the patient's status changes to	the same. Enter INPATIENT admit	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
			inpatient.	date.	
300	PayorGov	Payor - Government Health Insurance	Indicate whether government insurance was used by the patient to pay for part or all of this admission.	Government insurance refers to patients who are covered by government- reimbursed care. This includes Medicare, Medicaid, Military Health Care (e.g. TriCare), State-Specific Plan, and Indian Health Service. CHIP (Children's Health Insurance Plan), High Risk Pools Local Government Health Insurance Plan (LGHIP), state or federal prisoners Do not code yes if Medicare or Medicaid was applied for during the hospital stay but is not paying for	
310	PayorGovMcare	Payor - Government Health Insurance - Medicare	All payor fields are "Select all that apply." If a patient has a Medicare HMO, code "Yes" to this field (PayorGovMcare) and PayorHMO re HMO.		

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320	MedicareFFS	Medicare Fee For Service	 Indicate whether the patient is a Medicare Fee For Service (FFS) patient, Medicare FFS=Medicare Part B There are four parts to Medicare: Medicare Part A, Hospital Insurance; Medicare Part B, Medical Insurance; Medicare Part C (Medicare Advantage), which was formerly known as <i>Medicare</i> + <i>Choice;</i> and Medicare Part D, prescription drug coverage You cannot assume if a patient has part A that they have Part B 	The Social Security Website at <u>www.socialsecurity.gov</u> has a list explaining what the letters behind the Medicare claim # stand for. Those letters do not tell you whether they have Part B/Fee for service. It is the relationship of the cardholder to the Medicare/SSN #. For example, B stands for "Aged wife, 62 or older". The A would stand for "Primary claimant=the wage earner". D1 is for an "Aged widower, age 60 or over". This is used for PQRS Check with your hospital billing department if you are unsure whether the patient is considered Medicare Part B. Even if not using the registry for PQRS, CMS will be tracking outcomes for value based purchasing.	
331	MHICNumber PayorGovMcaid	Medicare Health Insurance Claim Number Payor - Government Health Insurance - Medicaid	Indicate the Medicare Health Insurance Claim (MHIC) number of the primary beneficiary. The Indicate whether the government insurance used by the patient to pay for part or all of this admission included Medicaid	The MHIC # consists of the Social Security number and an alpha-numeric identifier (usually one digit but maybe two digits) It is the number found on a patient's Medicare card. This field should be collected in compliance with state/local privacy laws.	
350	PayorGovMil	Payor - Government Health Insurance - Military Health	Indicate whether the government insurance used by the patient to pay for part or all of this admission included	Examples include: TriCare, Champus, Department of Defense, and	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
		Care	Military Health Care.	Department of Veterans Affairs.	
360	PayorGovState	Payor - Government Health Insurance - State-Specific Plan	Indicate whether the government insurance used by the patient to pay for part or all of this admission included State-Specific Plan.		
370	PayorGovIHS	Payor - Government Health Insurance - Indian Health Service	Indicate whether the government insurance used by the patient to pay for part or all of this admission included Indian Health Service.		
380	PayorCom	Payor - Commercial Health Insurance	Indicate whether commercial insurance was used by the patient to pay for part or all of this admission.	Commercial insurance refers to all indemnity (fee-for-service) carriers and Preferred Provider Organizations (PPOs), (e.g., Blue Cross and Blue Shield). Workman's compensation is considered commercial insurance.	
390	PayorHMO	Payor - Health Maintenance Organization	Indicate whether Health Maintenance Organization (HMO) insurance was used by the patient to pay for part or all of this admission.	HMO refers to a Health Maintenance Organization characterized by coverage that provides health care services for members on a pre-paid basis.	
400	PayorNonUS	Payor - Non-U.S. Insurance	Indicate whether any non-U.S. insurance was used by the patient to pay for part or all of		
410	PayorNS	Payor - None / Self	Indicate whether no insurance was used by the patient to pay for this admission.	None refers to individuals with no or limited health insurance; thus, the individual is the payor regardless of ability to pay. Only mark "None" when "self" or "none" is denoted as the first insurance in the medical record.	
420	Surgeon	Surgeon's Name	Indicate the Surgeon's name. This field must have controlled data entry where a user selects the surgeon name from a user list. This will remove variation in spelling, abbreviations and punctuation within the field. Note: Surgeon name is encrypted in the analysis database. Punctuation, abbreviations and spacing differences cannot be corrected at the warehouse.	If two surgeons participate in the procedure and both surgeons are participating in the Database, the surgeon of record for the database is the physician under whom the patient is admitted or the physician responsible for the care of the patient. If this is not evident from the operative	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				dictation, communication with the	
430	SurgNPI	Surgeon's National Provider Identifier	Indicate the individual-level National Provider Identifier (NPI) of the surgeon performing the procedure. This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis	The NPI is a unique identification number for health care providers. Health care providers will use the NPIs in the administrative and financial transactions adopted under HIPAA. The NPI is a 10-position, intelligence-free numeric identifier (10-digit number) Meaning that the numbers do not carry other information about healthcare providers, such as the state in which they live or their medical specialty. NPI look up link: https://nppes.cms.bbs.gov/NPPES/NPIP	
440	TIN	Taxpayer Identification Number	Indicate the TIN used by the physician office for billing purposes for this patient's procedure. There may be individual, hospital and medical group practice TINs, so be sure to enter the correct one.	If the physician is part of a medical group practice, use the name and taxpayer identification number of the medical group.	
450	HospName	Hospital Name	Indicate the full name of the facility where the procedure was performed.		
460	HospZIP	Hospital Postal Code	Indicate the ZIP Code of the hospital. Outside the USA, these data may be known by other names such as Postal Code (needing 6 characters).	Software should allow sites to collect up to 10 characters to allow for Zip+4 values. This field should be collected in compliance with state/local privacy laws.	
470	HospStat	Hospital State	Indicate the abbreviation of the state or province in which the hospital is located.		
480	HospNPI	Hospital National Provider Identifier	Indicate the hospital's National Provider Identifier (NPI).	This number, assigned by the Center for Medicare and Medicaid Services (CMS), is used to uniquely identify facilities for Medicare billing purposes.	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification
				This is different from the surgeon NPI.
490	HeightCm	Height In Centimeters	Height and weight is extremely important for the accurate interpretation of PFTs, body surface area and risk calculations. Ft-in = cm 4'10'' = 147 4'11'' = 149 5'0'' = 152 5'1'' = 155 5'2'' = 157 5'3'' = 160 5'4'' = 163 5'5'' = 168 5'7'' = 170 5'8'' = 173 5'9'' = 175 5'10'' = 178 5'11'' = 180 6'0'' = 183 6'1'' = 185 6'2'' = 188 6'3'' = 190 6'4'' = 193 6'5'' = 198 6'7'' = 200	
500	WeightKg	Weight In Kilograms	Height and weight is extremely important for the accurate interpretation of PFTs, body surface area and risk calculations. To convert pounds to kilograms, divide # of lbs by $2.2.(1 \text{ kg} = 2.2 \text{ lbs})$	
510	WtLoss3Kg	Weight Loss In Past Three Months	This is a significant indicator of the patients overall health within the last few months. Unintentional weight loss may be an indicator of underlying pathology. If the	,

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			amount of weight loss is not documented or it is unclear how much has occurred in the 3 month window leave this field blank.			
520	Hypertn	Hypertension	The History & Physical form will list the patients past medical history and also will list the current medications. This definition comes from: ACCF/AHA Key Elements and Data Definitions for Measuring the Clinical Management and Outcomes of Patients with Acute Coronary Syndromes and Coronary Artery Disease A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Data Standards 2010 (Writing Committee to Develop Acute Coronary Syndromes and Coronary Artery Disease Clinical Data Standards)	Diagnosis of hypertension should not be based on a single elevated blood pressure reading, rather a diagnosis of hypertension, documented by one of the following: a) Documented history of hypertension diagnosed and treated with medication, diet and/or exercise b) Prior documentation of blood pressure >140 mmHg systolic or 90 mmHg diastolic for patients without diabetes or chronic kidney disease, or prior documentation of blood pressure >130 mmHg systolic or 80 mmHg diastolic on at least 2 occasions for patients with diabetes or chronic kidney disease. c) Currently on pharmacologic therapy to control hypertension.		
530	Steroid	Steroids	Corticosteroids (or steroids) have been developed for their anti-inflammatory and immunomodulatory effects. Patients on steroids who present for surgery may be at increased risk of complications because of the adrenal suppression caused by steroid therapy. This often poses the greatest risk and deserves particular attention. Examples of oral and intravenous steroid medications include prednisone, hydrocortisone, dexamethasone, and methylprednisolone.	DO NOT include topical creams or inhalers that are steroidal in form. DO NOT include a one or two time dose of systemic treatment, or a pre- operative/pre-cath protocol.		
540	CHF	Congestive Heart Failure	Congestive heart failure occurs when the heart is unable to pump blood effectively throughout the body. The term congestive is used because lung congestion causes some of the main symptoms of heart failure. ACCF/AHA Key Elements and Data Definitions for	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		

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
			Measuring the Clinical Management and Outcomes of Patients with Acute Coronary Syndromes and Coronary Artery Disease	retention; or the description of rales, jugular venous distension, pulmonary edema on physical exam, or pulmonary edema on chest x-ray. A low ejection	
			A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Data Standards 2010 (Writing Committee to Develop Acute Coronary Syndromes and Coronary Artery Disease Clinical Data	fraction without clinical evidence of heart failure does not qualify as heart failure.	
			Standards)		
550	CAD	Coronary Artery Disease	Coronary artery disease is a type of atherosclerosis in which plaque builds up inside the arteries that carry blood to the heart. As the artery walls thicken, the passageway for blood narrows. Sometimes platelets gather at the narrowing, forming a clot that decreases or prevents blood flow to the region of the heart supplied by the artery. May include documentation of angina, myocardial infarction (MI), CABG, PCI*, or sudden cardiac death with no known cause. *Percutaneous Coronary Intervention (PCI) includes angioplasty and coronary artery stenting.	Documented blockage ≥ 50% of one or more coronary arteries or documentation of CAD in H&P.	
560	PVD	Peripheral Vascular Disease	This refers to diseases of blood vessels outside the heart and brain. It is often a narrowing of vessels that carry blood to the legs, arms, stomach or kidneys. Peripheral arterial disease excludes disease in the carotic or cerebral vascular arteries or thoracic aneurysms.	 Peripheral arterial disease can include any of the following: claudication either with exertion or rest; amputation for arterial vascular insufficiency; aorto-iliac occlusive disease reconstruction vascular reconstruction, bypass surgery, or percutaneous intervention to the extremities (excluding dialysis fistulas and vein stripping) peripheral angioplasty or stent 	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				 documented documented abdominal (below the diaphragm) aortic aneurysm with or without repair positive non-invasive or invasive testing documented ankle brachial index ≤ 0.9, angiography, ultrasound, MRI or CT imaging of > 50% stenosis in any peripheral artery. 	
570	PriorCTS	Prior Cardiothoracic Surgery	Prior cardiothoracic surgery causes scar tissue to form and may increase difficulty and or risk in subsequent procedures.	Capture open and minimally invasive procedures	
580	PreopChemoCur	Preoperative Chemo - Current Malignancy	Chemotherapy treatment for the current malignancy ONLY. Does not include chemo for prior thoracic or other malignancies.	Do not include methotrexate given for arthritis	
590	PreopChemoCurWhen	Preoperative Chemo - Current Malignancy - When	Indicate whether chemotherapy was given within 6 months or greater than 6 months prior to procedure		
600	PreopXRT	Preoperative Thoracic Radiation Therapy	Radiation to the chest area for any disease process at any time prior to current surgery. If answered yes, answer the next field.	Radiation therapy causes changes to the tissues which may increase difficulty and or risk in subsequent surgeries.	
610	PreopXRTDisWhen	Preoperative Thoracic Radiation Therapy - Disease And When Treated	If yes, select Same Disease, ≤ 6 months; Same Disease, > 6 months; Unrelated Disease, ≤ 6 months, or related Disease, > 6 months.	If patient did not receive preoperative radiation therapy as indicated by a "Yes" in PreopXRT, there should be no option to answer.	
620	CerebroHx	Cerebrovascular History	Select No CVD history; Any reversible event; or Any irreversible event. If a history of previous cerebrovascular disease exists, it should be noted whether the patient's symptoms were or reversible (i.e. transient ischemic attack) or whether the deficit is permanent (i.e. stroke).	CVD may be documented by any one of the following: CVA (symptoms > 24 hrs after onset, presumed to be from vascular etiology) TIA (recovery within 24 hrs) Non-invasive carotid test with > 79% diameter occlusion Documented "severe" or "critical" carotid stenosis. Prior carotid surgery or stenting or prior cerebral aneurysm clipping or coil. Does not include	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				neurological disease processes such as	
				metabolic or ischemic encephalopathy.	
What if a transient neuro event lasts more than 24 hours but resolves? Is this coded as reversible or irreversible?		asts more than 24 hours but ble or irreversible?	Use the 24 hour timeframe- if symptoms resolve within 2 persist for more than 24 hours, code as irreversible. This version.	24 hours, code as reversible. If symptoms will be reviewed for the next data	
630	PulmHypertn	Pulmonary Hypertension	High blood pressure in the arteries that supply the lungs is called pulmonary hypertension (PHT). The blood vessels that supply the lungs constrict and their walls thicken, so they cannot carry as much blood. This information may be found on a preoperative cardiac catheterization or echocardiogram. If the value is not known or documented, the data sheet should be marked accordingly.		
640	Diabetes	Diabetes	Diabetes is a condition whereby the body is not able to regulate levels of glucose (a sugar) in the blood, resulting in too much glucose being present in the blood. The American Diabetes Association criteria include documentation of the following: 1. A1c >6.5%; or 2. Fasting plasma glucose >126 mg/dl (7.0 mmol/l); or 3. Two-hour plasma glucose >200 mg/dl (11.1 mmol/l) during an oral glucose tolerance test; or 4. In a patient with classic symptoms of hyperglycemia or hyperglycemic crisis, a random plasma glucose >200 mg/dl (11.1 mmol/l)	Capture the presence and/or history of diabetes mellitus, regardless of duration of disease or need for anti-diabetic agents diagnosed prior to surgical intervention.	
650	DiabCtrl	Diabetes Control			
660	Dialysis	Currently On Dialysis	This measure is related to hemodialysis, peritoneal dialysis or CRRT. Does not include ultrafiltration.		
670	CreatMeasured	Creatinine Level Measured	Creatinine, urea and urate all increase as the ability of the kidneys to filter fluid within the body declines. Creatinine is a marker for kidney function.		
680	CreatLst	Last Creatinine Level	The value used should be most recent one prior to entering the operating room.		
690	HemoglobinMeasured	Hemoglobin Level Measured	Hemoglobin is the protein molecule in red blood cells that carries oxygen from the lungs to the body's tissues		

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			and returns carbon dioxide from the tissues to the lungs. The iron contained in hemoglobin is responsible for the red color of blood.		
700	HemoglobinLst	Last Hemoglobin Level	The value used should be the most recent one prior to entering the operating room.		
710	COPD	COPD	Chronic Obstructive Pulmonary Disease (COPD) is a preventable and treatable lung disease with some significant extrapulmonary effects. It is characterized by airflow limitation that is not fully reversible, usually progressive and associated with an abnormal inflammatory response in lung tissue. Diagnosis is confirmed and severity is graded using pulmonary function testing (PFT). Bronchitis and emphysema are considered COPD, asthma is not.		
720	InterstitialFib	Interstitial Fibrosis	Interstitial lung disease, or ILD, is a common term that includes more than 200 chronic lung disorders Interstitial lung diseases are named after the tissue between the air sacs of the lungs called the interstitium. This tissue can be affected by fibrosis (scarring) and lead to respiratory insufficiency.		
730	CigSmoking	Cigarette Smoking	This field applies to cigarettes only. Select: Never Smoked; Past Smoker (stopped > 1 month prior to operation); or Current Smoker. "Current smoker" should be selected if the patient stopped smoking < than 1 month prior to surgical procedure. This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.		
How do you code smoking status if there is conflicting documentation in the chart?		s if there is conflicting	Code yes to smoking if any provider documents it in the rof pack years documented.	record and capture the highest number	
740	PackYearKnown	Pack Years Known or can be estimated			
750	PackYear	Pack-Years Of Cigarette Use	Multiply the average number of packs per day by the		

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			number of years the patient smoked.			
760	PFT	Pulmonary Function Tests Performed	Note that per the definition, PFTs are part of the NQF measure set and are required before any major anatomic lung resection. This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
April 2014	The following <u>NQF meas</u> Segmentectomy).	ure specifications apply to 045	8 PFTs before Major Anatomic Lung Resection (Pneumonec	tomy, Lobectomy, or Formal		
	S.6. Numerator Details					
	Number of patients unde Database, Version 2.2, se	ergoing major anatomic lung re equence number 760) is marke	esection who undergo at least one pulmonary function test; ed as "Yes"	PFT (STS General Thoracic Surgery		
	S.9. Denominator Detail	S				
	 5.9. Denominator Details 1. Primary procedure is one of the following CPT codes: Removal of lung, total pneumonectomy; (32440) Removal of lung, sleeve (carinal) pneumonectomy (32442) Removal of lung, single lobe (lobectomy) (32480) Removal of lung, two lobes (bilobectomy) (32482) Removal of lung, two lobes (bilobectomy) (32482) Removal of lung, single segment (segmentectomy) (32484) Removal of lung, sleeve lobectomy (32486) Removal of lung, sleeve lobectomy (32488) Resection of apical lung tumor (e.g., Pancoast tumor), including chest wall resection, without chest wall reconstruction(s) (32503) Resection of apical lung tumor (e.g., Pancoast tumor), including chest wall resection, with chest wall reconstruction (32504) Thoracoscopy, surgical; with lobectomy (32663) Thoracoscopy with removal of a single lung segment (segmentectomy) (32669) Thoracoscopy with removal of a single lung segment (segmentectomy) (32670) Thoracoscopy with removal of lung, pneumonectomy (32671) Non-missing data on whether or not PFT was done Status of Operation (Status - STS General Thoracic Surgery Database, Version 2.2, sequence number 1420) is marked as "Elective" 					

	Training Manual Version 2.2 of the General Thoracic Surgery Database				
SeqNo	ShortName	LongName	DataFieldIntent FieldNameClarification		
	S.11. Denominator Exc	lusion Details			
	Pulmonary function tes	ts performed (PFT - STS General	Thoracic Surgery Database, Version 2.2, sequence number 760) is marked as "No" and reason PF		
	not performed (PFT No	tPerReas – STS GTSD, Version 2.	2, sequence number 770) is marked "tracheostomy or ventilator," "patient unable to perform," or		
	"urgent or emergent st	atus."			
770	PFTNotPerReas	PFT Not Performed Reason	There are acceptable reasons not to perform PFTs.		
			These will be included in the NQF exclusions:		
			Not Major Lung Resection		
			Never smoked, no lung disease		
			Patient unable to perform		
			Tracheostomy or ventilator dependent		
			Urgent or emergent procedure		
April		A wedge is not a major anaton	nic resection, even if therapeutic, but is a major (analyzed case) if it is therapeutic. The PFT field		
2014		770 should be answered "Not	a major lung resection" for therapeutic (analyzed) wedges.		
780	FEV	Forced Expiratory Volume Tex	st This field is Required for Record Inclusion. If missing		
		Performed	data, the entire record will be excluded from the		
			analysis.		
790	FEVPred	FEV1 Predicted	This field is Required for Record Inclusion. If missing		
			data, the entire record will be excluded from the		
			analysis.		
Should	the FEV1 be pre or post	bronchodilator?	The value should ideally be taken from a good quality spirometry exam prior to surgery. Choose		
What tir	meframe is acceptable p	reop?	the highest value reported for % predicted, whether pre or post bronchodilator. Do not use values		
			obtained more than 6 months prior to surgery.		
ADDEN	IDUM October 2013		Do not use values obtained more than 12 months prior to surgery.		
800	DLCO	DLCO Test Performed	The diffusing capacity (DLCO) is a test of the integrity		
			of the alveolar-capillary surface area for gas transfer.		
Should	DLCO be corrected for	volume or hemoglobin?	Use the uncorrected/unadjusted DLCO. Do not use values obtained more than 6 months prior to		
What tin	meframe is acceptable p	reop?	surgery.		
ADDEN	DUM October 2013		Do not use values obtained more than 12 months prior to surgery.		
810	DLCOPred	DLCO Predicted	The diffusing capacity (DLCO) may be reduced, <80%		
			predicted, in disorders such as emphysema, pulmonary		
			fibrosis, obstructive lung disease, pulmonary embolism,		
			pulmonary hypertension and anemia. DLCO>120% of		
			predicted may be seen in normal lungs, asthma,		
			pulmonary nemorrhage, polycythemia, and left to right		
			intracardiac snunt.		

Training Manual Version 2.2 of the General Thoracic Surgery Database				
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification
820	Zubrod	Zubrod Score	Select: Normal activity, no symptoms; Symptoms, fully ambulatory; Symptoms, in bed <= 50% of time; Symptoms, in bed > 50% but < 100% of time; Bedridden, or Moribund.	This score is used in risk calculation therefore it is important not to "under code".
			Code the most severe Zubrod score within two weeks of surgery. For example: a patient enters hospital with Zubrod score of 1 but after 1 week in the hospital, the Zubrod score changes to 3, then the score entered on the data sheet should be "3". A new data collection form generated for a subsequent surgery may have a different Zubrod score than the previous data collection form (previous surgery). This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis	
There is	confusion about how and	when to capture the Zubrod	Clarification and Examples	
score. Since it is part of the risk model it is important to be consistent and accurate when coding this element.			Use the score that most accurately represents the patient' patient is ambulatory at the time of admission but deterio bedridden, capture the bedridden status. Conversely, if th stabilized and ambulatory just prior to surgery, capture an	s status at the time of surgery. If a rates in the hospital, becoming e patient comes in bedridden, but is nbulatory.
830	LungCancer	Lung Cancer	If Lung Cancer documented, and resection performed, complete both Clinical Staging (ClinStageLungT, ClinStageLungN, and ClinStageLungM) AND Pathological Staging (PathStageLungT, PathStageLungN, and PathStageLungM). This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.	
840	ClinStagDoneLung	Clinical Staging Performed For Lung Cancer	Clinical staging is based on evidence gathered before primary treatment. Diagnostic and/or radiologic tests are	

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
			performed to determine the type and extent of the cancer		
			and used to guide treatment decisions.		
May	Patient (RR) had a VATS	S wedge for lung cancer in Octob	er 2013. Now with If it is a new primary, it has to be stag	ged.	
2014	recurrence and is having	a completion lobectomy. Can I u	use the Clinical If it's mets, then no need to stage.		
	Staging Methods that we	re done prior to the first surgery,	or must they be		
	since the first surgery and	d up to the present procedure?			
850	ClinStagLungBronc	Clinical Staging Method - Lung	Bronchoscopy is a procedure in which a cylindrical		
		- Bronchoscopy	fiberoptic scope is inserted into the airways. This scope		
			contains allows the visual examination of the lower		
			airways. During a bronchoscopy, a physician can		
			visually examine the lower airways, including the		
			larynx, trachea, bronchi, and bronchioles. The procedure		
			is used to examine the mucosal surface of the airways		
			for abnormalities that might be associated with a variety		
			of lung diseases. Its use includes the visualization of		
			airway obstructions such as a tumor, or the collection of		
			specimens for the diagnosis of cancer originating in the		
			bronchi of the lungs (bronchogenic cancer). It can also		
			be used to collect specimens for culture to diagnose		
			infectious diseases such as tuberculosis. The type of		
			specimens collected can include sputum (composed of		
			saliva and discharges from the respiratory passages),		
			tissue samples from the bronchi or bronchioles, or cells		
			collected from washing the lining of the bronchi or		
			bronchioles. The instrument used in bronchoscopy, a		
			bronchoscope, is a slender cylindrical instrument		
			containing a light and an eyepiece. There are two types		
			of bronchoscopes, a rigid tube that is sometimes referred		
			to as an open-tube or ventilating bronchoscope, and a		
			more flexible fiber optic tube. This tube contains four		
			smaller passages—two for light to pass through, one for		
			seeing through and one that can accommodate medical		
			instruments that may be used for biopsy or suctioning,		
			or that medication can be passed through.		
860	ClinStagLungEBUS	Clinical Staging Method - Lung	EBUS is an invasive procedure in which physicians use		

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
		- EBUS	ultrasound devices inside the airways and the lung for			
			exploration of the structures of airway walls, the			
			surrounding mediastinum, and the lungs.			
870	ClinStagLungEUS	Clinical Staging Method - Lung	A procedure that combines endoscopy and ultrasound to			
		- EUS	obtain images and information about the digestive tract			
			and the surrounding tissue and organs. In EUS a small			
			ultrasound transducer is installed on the tip of the			
			endoscope allowing the transducer to get close to the			
			organs inside the body so the resultant ultrasound			
			images are often more accurate and detailed than ones			
			obtained by traditional ultrasound.			
880	ClinStagLungMedia	Clinical Staging Method - Lung	Mediastinoscopy is a procedure that enables			
		- Mediastinoscopy/Chamberlain	visualization of the contents of the mediastinum, usually			
			for the purpose of obtaining a biopsy. Mediastinoscopy			
			is often used for staging of lymph nodes of lung cancer			
			or for diagnosing other conditions affecting structures in			
			the mediastinum such as sarcoidosis or lymphoma.			
			Mediastinoscopy involves making an incision			
			approximately 1 cm above the suprasternal notch of the			
			sternum, or breast bone. Dissection is carried out down			
			to the pretracheal space and down to the carina. A scope			
			(mediastinoscope) is then advanced into the created			
			tunnel which provides a view of the mediastinum. The			
			scope may provide direct visualization or may be			
			attached to a video monitor.			
			The Chamberlain procedure is used to biopsy lymph			
			nodes in the center of the chest, or to biopsy a mass in			
			the center of the chest. The Chamberlain procedure			
			differs from a cervical mediastinoscopy by the location			
			of the incision, and the location of the lymph nodes or			
			mass to be biopsied.			
			The Chamberlain procedure is used to biopsy lymph			
			nodes or masses in the aorto-pulmonary window on the			
			left side of the chest, or nodes in the hilar areas of the			
			lung. (In contrast, the cervical mediastinoscopy			

Training Manual				
		Version 2.2 d	of the General Thoracic Surgery Database	
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification
Seque	Shortwanie		procedure is used to biopsy nodes or masses to the front or side of the trachea, or windpipe.) The aorto-pulmonary window is the area in the center of the chest bound by the aorta superiorly, and the pulmonary artery inferiorly. This area contains lymph nodes that filter lymph coming from the left lung, especially the left upper lobe. If a lung cancer is present in the left lung, the Chamberlain procedure is useful for staging the cancer (determining the extent of spread.) The hilar areas of the lung (the hilum) are the areas of the lung where the pulmonary	
890	ClinStagLungPET	Clinical Staging Method - Lung - PET or PET/CT	artery and vein (the blood supply) join the lung. Positron emission tomography, also called PET imaging or a PET scan, is a type of nuclear medicine imaging. Nuclear medicine or radionuclide imaging procedures are noninvasive and, with the exception of intravenous injections, are usually painless medical tests that help diagnose medical conditions. These imaging scans use radioactive materials called radiopharmaceuticals or radiotracers.	
900	ClinStagLungCT	Clinical Staging Method - Lung - CT	Computed tomography (CT) scan, also called computerized axial tomography (CAT) scan, is used to create cross-sectional images of structures in the body. In this procedure, x-rays are taken from many different angles and processed through a computer to produce a three-dimensional (3-D) image called a tomogram.	
910	ClinStagLungVATS	Clinical Staging Method - Lung - VATS	Video-assisted thoracoscopic surgery (VATS) is a minimally invasive surgical technique used to diagnose and treat problems in the chest. During this surgery, a tiny camera (thoracoscope) and surgical instruments are inserted in the chest through small incisions. The thoracoscope transmits images of the inside of the chest onto a video monitor, guiding the surgeon performing the procedure. Video-assisted thoracoscopic surgery (VATS) can be used for many purposes, ranging from a	

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
			biopsy to removal of tumors or entire lobes from the lung.		
920	ClinStagLungLap	Clinical Staging Method - Lung - Laparoscopy	Laparoscopy is a minimally invasive procedure used as a diagnostic tool and surgical procedure that is performed to examine the abdominal and pelvic organs, or the thorax, head, or neck. Tissue samples can also be collected for biopsy using laparoscopy and malignancies treated when it is combined with other therapies. Laparoscopy can also be used for some cardiac and vascular procedures.		
921	ClinStagLungOth	Clinical Staging Method - Lung-Other	Indicate if any other method/technology was used for clinical staging.		
930	ClinStageLungT	Lung CA Tumor size - T	Choose from the list, if more than one tumor is present, choose the largest. American Joint Committee on Cancer 2010		
How are staging?	small nodules reported o	on lung CT addressed for	If there is no biopsy, the PET CT is negative, nodules are < 5 mm and the surgeon/oncologist chooses not to address these, do not consider them when staging. 40% of people over the age of 50 have small lung nodules which are not cancerous.		
940	LCInvAdjStr	Lung Cancer- Invasion of Adjacent Structures	Based on preop testing, does tumor appear to invade adjacent structures? American Joint Committee on Cancer 2010	Choose all that apply	
950	ClinStageLungTInvPl	Clinical Staging Lung Tumor Invasive Pleura	This refers to <u>visceral pleura only</u> . If the tumor invades the parietal pleura, code as invading the chest wall (next field)		
960	ClinStageLungTInvCW	Clinical Staging Lung Tumor Invasive Chest Wall	Code tumors that invade the <u>parietal pleura</u> as invading the chest wall.		
970	ClinStageLungTInvDia	Clinical Staging Lung Tumor Invasive Diaphragm			
980	ClinStageLungTInvPN	Clinical Staging Lung Tumor Invasive Phrenic Nerve			
990	ClinStageLungTInvPer	Clinical Staging Lung Tumor Invasive Pericardium			
1000	ClinStageLungTInvMB	Clinical Staging Lung Tumor Invasive Main Bronchus			
1010	ClinStageLungTInvOb	Clinical Staging Lung Tumor			

	Training Manual Version 2.2 of the General Thoracic Surgery Database				
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
		Obstructive			
1020	ClinStageLungTInvNod	Clinical Staging Lung Tumor			
		Invasive Nodule(s)			
1030	ClinStageLungTInvMed	Clinical Staging Lung Tumor			
		Invasive Mediastinum			
1040	ClinStageLungTInvHt	Clinical Staging Lung Tumor			
		Invasive Heart			
1050	ClinStageLungTInvGrV	Clinical Staging Lung Tumor			
	es	Invasion Great Vessels			
1060	ClinStageLungTInvTr	Clinical Staging Lung Tumor			
		Invasion Trachea			
1070	ClinStageLungTInvRLN	Clinical Staging Lung Tumor			
		Invasive Recurrent Laryngeal			
		Nerve			
1080	ClinStageLungTInvEo	Clinical Staging Lung Tumor			
		Invasive Esophagus			
1090	ClinStageLungTInvVB	Clinical Staging Lung Tumor			
		Invasive Vertebral Body			
1100	ClinStageLungTInvC	Clinical Staging Lung Tumor			
		Invasive Carina			
1110	ClinStageLungTInvNDL	Clinical Staging Lung Tumor			
		Invasive Nodule(s) Diff Lobe			
1120	ClinStageLungN	Lung Cancer Nodes - N	Code nodal involvement if any. Ipsilateral = same side		
			as tumor, contralateral= opposite side		
1130	ClinStageLungM	Lung Cancer Metastasis - M	Metastasis or metastatic disease (sometimes abbreviated		
			mets), is the spread of cancer from one organ to another		
			non-adjacent organ or part.		
1140	EsophCancer	Esophageal Cancer	If Esophageal Cancer documented, "Yes", and resection		
			performed, complete both Clinical Staging		
			(ClinStageEsophT, ClinStageEsophN,		
			ClinStageEsophM, ClinStageEsophH, and		
			ClinStageEsophG) AND Pathological Staging		
			(PathStageEsophT, PathStageEsophN,		
			PathStageEsophM, PathStageEsophH, and		
			PathStageEsophG).		

	Training Manual Version 2.2 of the General Thoracic Surgery Database				
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
		~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	This field is Required for Record Inclusion. If missing		
			data, the entire record will be excluded from the		
			analysis.		
Are ther	e guidelines (measuremer	nts) for coding esophageal tumor	Use these measurements in cm from incisors:		
location	?		Upper third = 17-25 cm		
			Middle third = $26-34$ cm		
			Lower third = $35-42$ cm		
1150	ClinStagDoneEsoph	Clinical Staging Performed For	Clinical staging is the Pre-Treatment estimate of cancer.	Choose all that apply	
		Esophageal Cancer	Indicate whether clinical staging was performed and if		
			so choose the method(s)		
1160	ClinStagEsophBronc	Clinical Staging Method -	Bronchoscopy is a procedure in which a cylindrical		
		Esophageal - Bronchoscopy	fiberoptic scope is inserted into the airways. This scope		
			contains allows the visual examination of the lower		
			airways. During a bronchoscopy, a physician can		
			visually examine the lower airways, including the		
			larynx, trachea, bronchi, and bronchioles. The procedure		
			is used to examine the mucosal surface of the airways		
			for abnormalities that might be associated with a variety		
			of lung diseases. Its use includes the visualization of		
			airway obstructions such as a tumor, or the collection of		
			specimens for the diagnosis of cancer originating in the		
			bronchi of the lungs (bronchogenic cancer). It can also		
			be used to collect specimens for culture to diagnose		
			infectious diseases such as tuberculosis. The type of		
			specimens collected can include sputum (composed of		
			saliva and discharges from the respiratory passages),		
			tissue samples from the bronchi or bronchioles, or cells		
			collected from washing the lining of the bronchi or		
			bronchioles. The instrument used in bronchoscopy, a		
			bronchoscope, is a slender cylindrical instrument		
			containing a light and an eyepiece. There are two types		
			of bronchoscopes, a rigid tube that is sometimes referred		
			to as an open-tube or ventilating bronchoscope, and a		
			more flexible fiber optic tube. This tube contains four		
			smaller passages—two for light to pass through, one for		

	Training Manual Version 2.2 of the General Thoracic Surgery Database							
SeqNo	SeqNo ShortName LongName DataFieldIntent Field							
			seeing through and one that can accommodate medical					
			instruments that may be used for biopsy or suctioning,					
			or that medication can be passed through.					
1170	ClinStagEsophEBUS	Clinical Staging Method -	EBUS is an invasive procedure in which physicians use					
		Esophageal - EBUS	ultrasound devices inside the airways and the lung for					
			exploration of the structures of airway walls, the					
			surrounding mediastinum, and the lungs.					
1180	ClinStagEsophEUS	Clinical Staging Method -	A procedure that combines endoscopy and ultrasound to					
		Esophageal - EUS	obtain images and information about the digestive tract					
			and the surrounding tissue and organs. In EUS a small					
			ultrasound transducer is installed on the tip of the					
			endoscope allowing the transducer to get close to the					
			organs inside the body so the resultant ultrasound					
			images are often more accurate and detailed than ones					
1100	ClinStagEconhMadia	Clinical Staging Mathod	Madiagtinggeony is a procedure that anghles					
1190	Christagesophiviedia	Esophagoal	visualization of the contents of the mediastinum usually					
		Esopliageal - Mediastinoscopy/Chamberlain	for the purpose of obtaining a biopsy. Mediastinoscopy					
		Wiedlastinoscopy/Chamberlain	is often used for staging of lymph podes or for					
			diagnosing other conditions affecting structures in the					
			mediastinum such as sarcoidosis or lymphoma					
			Mediastinoscopy involves making an incision					
			approximately 1 cm above the suprasternal notch of the					
			sternum, or breast bone. Dissection is carried out down					
			to the pretracheal space and down to the carina. A scope					
			(mediastinoscope) is then advanced into the created					
			tunnel which provides a view of the mediastinum. The					
			scope may provide direct visualization or may be					
			attached to a video monitor.					
			The Chamberlain procedure is used to biopsy lymph					
			nodes in the center of the chest, or to biopsy a mass in					
			the center of the chest. The Chamberlain procedure					
			differs from a cervical mediastinoscopy by the location					
			of the incision, and the location of the lymph nodes or					

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
			mass to be biopsied. The Chamberlain procedure is used to biopsy lymph nodes or masses in the aorto-pulmonary window on the left side of the chest, or nodes in the hilar areas of the lung. (In contrast, the cervical mediastinoscopy procedure is used to biopsy nodes or masses to the front or side of the trachea, or windpipe.) The aorto- pulmonary window is the area in the center of the chest bound by the aorta superiorly, and the pulmonary artery inferiorly.		
1200	ClinStagEsophPET	Clinical Staging Method - Esophageal - PET or PET/CT	Positron emission tomography, also called PET imaging or a PET scan, is a type of nuclear medicine imaging. Nuclear medicine or radionuclide imaging procedures are noninvasive and, with the exception of intravenous injections, are usually painless medical tests that help diagnose medical conditions. These imaging scans use radioactive materials called radiopharmaceuticals or radiotracers.		
1210	ClinStagEsophCT	Clinical Staging Method - Esophageal - CT	Computed tomography (CT) scan, also called computerized axial tomography (CAT) scan, is used to create cross-sectional images of structures in the body. In this procedure, x-rays are taken from many different angles and processed through a computer to produce a three-dimensional (3-D) image called a tomogram.		
1220	ClinStagEsophVATS	Clinical Staging Method - Esophageal - VATS	Video-assisted thoracoscopic surgery (VATS) is a minimally invasive surgical technique used to diagnose and treat problems in the chest. During this surgery, a tiny camera (thoracoscope) and surgical instruments are inserted in the chest through small incisions. The thoracoscope transmits images of the inside of the chest onto a video monitor, guiding the surgeon performing the procedure. Video-assisted thoracoscopic surgery (VATS) can be used for many purposes, ranging from a biopsy to removal of tumors.		
1230	ClinStagEsophEGD	Clinical Staging Method -	Esophagogastroduodenoscopy (EGD) is an examination		

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
		Esophageal - EGD	of the lining of the esophagus, stomach, and upper			
			duodenum with a small camera (flexible endoscope)			
1.0.10	~ ~ ~ ~ ~ ~		which is inserted down the throat.			
1240	ClinStagEsophLap	Clinical Staging Method -	Laparoscopy is a minimally invasive procedure used as			
		Esophageal - Laparoscopy	a diagnostic tool and surgical procedure that is			
			performed to examine the abdominal and pelvic organs.			
			language in the second			
			combined with other therapies			
1241	ClinStagEsophOth	Clinical Staging Method -	Indicate if any other method/technology was used for			
1211	ChildrengEsophioth	Esophageal - Other	clinical staging.			
1250	ClinStageEsophT	Esophageal Cancer Tumor - T	Record T based on EUS if done, if not done estimate T			
			based on CT or PET/CT. No esophageal thickening=			
			T1. Choose T2 if esophageal thickening is present.			
			If thickening noted on CT scan, code as T2. If stricture is	s noted on endoscopy or barium swallow		
	1		or the patient is experiencing dysphagia, code as T3.			
1260	ClinStageEsophN	Esophageal Cancer Nodes - N	Indicate nodal status. Nodes > 1cm on CT or PET/CT or			
			EUS are considered positive. All positive PET nodes are			
			considered positive. Count biopsy positive nodes.			
1270	<u>Clin Stopp Econh</u> M	Eachageal Cancer Metastasia	Choose NX II nodes cannot be assessed.			
1270	ChristageEsophivi	Esophageal Cancer Metastasis -	metastasis or metastatic disease (sometimes abbreviated			
		111	non-adjacent organ or part			
1300	CategoryPrim	Category Of Disease - Primary	Choose the primary diagnosis or reason for the	May 2014- Input should be based upon		
1500	CutogoryTillin	Category of Disease Triniary	procedure. This field is Required for Record Inclusion.	the final pathology report. If you		
			If missing data, the entire record will be excluded from	entered a Category of Disease before		
			the analysis.	final path, then you need to change it		
				based on the final pathology. Example,		
				if you start with a diagnosis of		
				"abnormal radiological finding", a		
				wedge resection is done and cancer is		
				found, the diagnosis should be changed		
				to cancer based upon the pathology		
1010				report.		
1310	CategoryPrimOth	Category Of Disease - Primary	Capture unlisted primary diagnosis here after carefully			

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
		- Other Specify	reviewing choices above.			
1311	CategoryPrimOthICD	Category Of Disease - Primary - Other ICD	The intent is to track category of disease codes for possible inclusion in next version and /or for internal analysis.			
1320	CategorySecond	Category Of Disease - Secondary	Secondary diagnosis is captured here	June 2013- The Secondary diagnosis can be left blank. As long as a primary diagnosis is selected, the record will be accepted as complete without having a secondary indicated.		
1330	CategorySecondOth	Category Of Disease - Secondary - Other Specify	Capture unlisted secondary diagnosis here after carefully reviewing choices above.			
1331	CategorySecondOthICD	Category Of Disease - Secondary - Other ICD	The intent is to track category of disease codes for possible inclusion in next version and /or for internal analysis.			
1340	SurgDt	Date Of Surgery	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
1350	OREntryT	OR Entry Time	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
1360	ORExitT	OR Exit Time	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.	Even if the thoracic surgeon was present for only part of the case, code the entire time.		
1370	AnesthStartT	Anesthesia Start Time	This is the start of anesthetic management, placing lines, induction of anesthesia.			
1380	AnesthEndT	Anesthesia End Time	If the patient is extubated in the OR, indicate time of extubation otherwise use OR exit time as anesthesia end time.			
1390	ProcStartT	Procedure Start Time	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
1400	ProcEndT	Procedure End Time	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
1410	MultiDay	Multi-Day Operation	These are cases that continue through midnight.			

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
1420	Status	Status	This indicates the clinical status of the patient at the time of surgery. Emergent- procedure must be performed as soon as possible, within 24 hours. Urgent-not emergent or elective but necessary within the same hospital stay. Elective- The patient is stable in the days or weeks prior to surgery. Palliative- The procedure is being done to provide			
Can you give urgent vs. emergent clinical examples?			comfort or reliefEmergent status is coded for cases that require immediate intervention to prevent life threatening deterioration or death such as (but not limited to) esophageal perforation, severe hemorrhage or massive hemoptysis.			
			be discharged. Examples of urgent cases would include decortication for empyema. Cases that are performed during the same hospitalization considered urgent. A medical patient with an incidental	oronchopleural fistula, pneumothorax or for convenience would not be CXR finding who undergoes a diagnostic		
			bronchoscopy or mediastinoscopy prior to discharge wor elective.	ald have the procedure status coded as		
1430	Reop	Reoperation	Did the patient have a previous operation in the same cavity or organ that affects this operative field?			
Decemb 2013	er If a patient is returned unit after the initial su location (ICU, Regula new operation, thus re	to the operating room from the p rgery, before they are sent to a pa r Floor Bed, etc.), is the second s quiring a separate STS data colle	boost anesthesia care atient disposition surgery considered a ection forms?For this purpose, PACU would = in Yes, fill out a $2^{nd}$ form.	termediate care.		
1440	Robotic	Robotic Technology Assisted	Was robotic technology used for any part of the procedure?			
1450	IntraopPRBC	Intraoperative Packed Red Blood Cells				
1460	IntraopPRBCNum	Intraoperative Packed Red Blood Cells - Number				
1470	ASA	ASA Classification	ASA Classification is determined by the anesthesiologist of the procedure based on the patient's condition. This is a standard risk scale for patients			

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName		DataFieldIntent	FieldNameClarification	
			undergoing anesthe This field is Require data, the entire reco analysis.	sia. ed for Record Inclusion. If missing rd will be excluded from the		
1480	Proc	Procedure	Check ALL the procedures that were performed. Complete Primary to indicate Primary procedure. The General Thoracic Surgery Database requires a separate data collection form for every OR or procedural area visit for major general thoracic procedure(s).			
June 2013	June Procedure done was 1)Left Thoractomy with upper lobectomy 2)Medialstinal Lymph 2013 node dissection. The only procedue code that I can find is 38746 which is Thoracic Lymphadenectomy, regional, including mediastinal and peritracheal nodes. There were no Peritacheal nodes removed, but there were mediastinal. Is it appropriated to use this Code (38746) for this procedue?					
Oct.	Is an Incisionless Transo	ral Fundoplication procedure list	ed in the Thoracic	Code "other" & write in the procedure.		
2013	Database Version 2.2, ald I list that procedure?	ong with a CPT Code? I cannot f	find it. How would			
Oct. 2013	Pt has: 1. Redo Right Th 2. Right chest parti 3. Pleurodisis Code 32225 will be used Plurodisis? Code 32650 not thoracotomy.	for the Partial Decort. How do v appears to be used only if done v	we code the with thoracoscopy,	Code both. But, 2430 Decortication, pulmonary, par	tial (32225) will be the primary procedure.	
1490	ProcOth	Procedure Unlisted - Specify				
1491	ProcOthCPT	Procedure Unlisted - CPT				
1500 Oct. 2013	Primary What STS procedure cod fistual? A Thoracoplasty major BP fistula even the	Primary Procedure le should I use for a VATS with o was not done. Can I use 32815- bugh the procedure was done tho	closure of BP - open closure of rascopically??	Yes, code 2860 Open closure of major bi	ronchial fistula (32815)	
1510	LungResect	Lung Resection Performed				
1520	Laterality	Laterality				
1530	PatDisp	Patient Disposition	The intent is to capt OR/PACU recovery Select ICU; Interme	ure the level of care following / period. ediate Care Unit; Regular Floor Bed;		

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			Not Applicable (Expired in OR). This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis. ICU level of care counts as ICU day- ex. PACU used for ICU overflow Do not include PACU stay unless patient was kept beyond the recovery phase as described above.			
1532	ICUVisitInit	Initial Visit To ICU	All ICU days should be included on first procedure			
1533	ICUVisitInitDays	Initial ICU Visit Days				
1534	ICUVisitAdd	Additional Visit To ICU				
1535	ICUVisitAddDays	Additional Visit To ICU Days				
1540	PathStageLungT	Pathologic Staging - Lung Cancer - T	Use the final pathology report to code the lung tumor T descriptor.	If Lung Cancer documented, "Yes", and lung resection performed, complete both Clinical Staging (ClinStageLungT, ClinStageLungN, and ClinStageLungM) AND Pathological Staging (PathStageLungT, PathStageLungN, and PathStageLungM). Reference: Goldstraw P, Crowley J et al. The IASLC Lung Cancer Staging Project: proposals for the revision of the TNM stage groupings in the forthcoming (seventh) edition of the TNM Classification of malignant tumours. J Thorac Oncol 2007; 2 (8): 706-714.		
1550	PathStageLungN	Pathologic Staging - Lung Cancer - N	Use the final pathology report to code the lung tumor node(s)			
1560	PathStageLungM	Pathologic Staging - Lung Cancer - M	Use the final pathology report to code the lung tumor M descriptor.			
1570	LungCANodes	Lung Cancer - Number of Nodes	Enter the number of nodes sampled/harvested during this procedure.			
1580	LungCAPathMarg	Lung Cancer - Pathology	Use the final pathology report to code whether the			

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName		DataFieldIntent	FieldNameClarification	
		Margins	surgical margins we	ere positive.		
1590	PathStageEsophT	Pathologic Staging - Esophageal Cancer - T	Use the final pathol descriptor for the pr	ogy report to code the appropriate T rimary tumor.	If Esophageal Cancer documented, "Yes" and resection performed, complete both Clinical Staging AND Pathological Staging.	
1600	PathStageEsophN	Pathologic Staging - Esophageal Cancer - N	Use the final pathol node(s)	ogy report to code assessment of the		
1610	PathStageEsophM	Pathologic Staging - Esophageal Cancer - M	Use the final pathol	ogy report to code metastatic status.		
June	Our patient had an esoph	agectomy and her final path was	'small cell	It should be listed in the pathology re	eport (H1 if Squamous, H2 if Adeno).	
2013	carcinoma'. On the Data ( Squamous or Adeno histo	Collection Form it only allows yo plogy. Please advise how to enter	ou to enter either this.	You should reread it & it it's not then in your files why it was left blank for	re, leave blank. You may wish to notate r your reference.	
1620	PathStageEsophH	Pathologic Staging - Esophageal Cancer - H	Use the final pathol	ogy report to code histologic type.		
1630	PathStageEsophG	Pathologic Staging - Esophageal Cancer - G	Use the final pathol differentiation. If a differentiation.	ogy report to code the grade of range is reported choose the worst		
1640	EsophCANodes	Esophageal Cancer - Number of Nodes	Enter the number of this procedure.	f nodes sampled/harvested during		
1650	EsophCAPathMarg	Esophageal Cancer - Pathology Margins	Use the final pathol surgical margins we	ogy report to code whether the ere positive.		
1710	POEvents	Postoperative Events Occurred	This field is meant postoperative comp developed due to th recording a Data Co occurred anytime du stay or until 30 days This does not includ operation or were p fibrillation.	to capture any instance of lications listed below that the patient be operation for which you are collection Sheet. These need to have uring the patient's entire hospital s post-op if they were discharged. de events that occur during the resent preoperatively, such as atrial		
May 2014	When does the postopera Is it when the patient leav	tive period begin for recording P ves the OR suite (OR Exit Time)	O events? or is it the	Post op begins when the patient leave	es the OR.	
	Procedure End Time (pat	ient is still in the OR)?				
1720	ReturnOR	Unexpected Return To The OR	The intent of this figure to the Operating Ro	eld is to ask if the patient went back oom for any type of <u>unexpected</u>	Do not include scheduled trips, such as surveillance bronchs or additional OR	

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			surgery during their stay at the hospital.	trips to assess original surgery.		
1730	ReturnORRsn	Reason for Unexpected Return to the OR	Choose the primary reason the patient was taken back to the OR.			
1750	AirLeak5	Air Leak Greater Than Five Days				
1760	Atelectasis	Atelectasis Requiring Bronchoscopy				
1770	CPIEff	Post-op-Pleural Effusion Requiring Drainage	Include only effusions requiring drainage with thoracentesis or chest tube. Do not code medically managed effusions.			
1780	Pneumonia	Pneumonia	Three of the five criteria must be met. (fever, leukocytosis, CXR with infiltrate, positive sputum culture, antibiotic treatment) Note: atelectasis and effusions do not necessarily indicate pneumonia, and neither does a single positive sputum culture without the other criteria/clinical findings documented.			
1790	ARDS	Adult Respiratory Distress Syndrome	Diagnosis of ARDS in medical record, or documentation of all criteria.			
1800	RespFail	Respiratory Failure	Inadequate gas exchange resulting in hypoxia and or hypercarbia.			
1810	Bronchopleural	Bronchopleural Fistula	Indicate if the patient experienced a complete or partial dehiscence of the bronchial stump documented by bronchoscopy or other operative intervention in the post-operative period.			
1820	PE	Pulmonary Embolus	Documented evidence in the chart by a high probability VQ scan, pulmonary angiogram or CT scan of the chest.			
1830	Pneumo	Pneumothorax	Only code a pneumothorax that required reinsertion of a chest tube. Do not code pneumothorax mentioned on CXR but not treated.			
Oct. 2013	Patient went home with H The definition for Post-op reinsertion." This patient count as a post op event of	Heimlich Valve due to "small api p events states: "pneumothorax r t never had his removed, they jus or not? Thank you.	ical pneumothorax." Code as air leak $> 5$ days, not pneumotho equiring chest tube st left it in. Does this	orax.		
1840	Vent	Initial Vent Support >48 Hours	The length of initial ventilatory support should be noted once the patient has the endotracheal tube removed after			

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			the operative procedure. For patients that are re-			
			intubated in the operating room at the conclusion of the			
			operation, this should still be considered initial			
			ventilator support and not re-intubation.			
1850	Reintube	Reintubate	Do not include reintubation for a planned postop			
			bronchoscopy or other planned procedure. For patients			
			that are re-intubated in the operating room at the			
			conclusion of the operation, this should still be			
			considered initial ventilator support and not re-			
1960	Tao ah	Tuo ah a a starray	Intubation.	New DCE antional		
1800	Trach	Tracheostomy	Do not include changing out a tracheostomy tube that	New DCF optional		
			intraoperatively during the initial operation			
			Do not include prophylactic mini tracheostomy performe	d on the day of surgery		
1870	OtherPul	Other Pulmonary Event	Pulmonary events not listed that extend the length of	Example: BiPan		
1070			stay or impact the patient's outcome.			
1880	AtrialArryth	Atrial Arrhythmia Requiring	This field is intended to capture new onset of atrial	Patients with an episode of Afib that		
		Treatment	arrhythmias that requires treatment. Treatment may	does not require treatment do not meet		
			include medications to slow the heart rate, increase the	this definition.		
			blood pressure, or any anti-coagulation administered for			
			embolic prophylaxis. This does not include those			
			patients with a preoperative history of atrial arrhythmias.			
1890	VentArryth	Ventricular Arrhythmia				
		Requiring Treatment				
1900	MI	Myocardial Infarct				
1910	DVT	DVT Requiring Treatment				
May	Patient JS was being foll	owed for a DVT, confirmed by D	Doppler post op, but Correct, it's pre-existing.			
2014	this was chronic and sho	wed "no significant interval chan	ge" from pre-op.			
	DC Summary states patie	ent "did not require full anticoagu	ilation," although			
	sne did receive 6 1/2 day	s worth of sq heparin q8 hrs. I an	n unsure whether or			
	not to consider this DVT	a Post Operative Event. I'm incli	ined to say not,			
1020	Since it was present pre-0	p. Other Cardiovacewler Evert	Condinuoscular avants not listed that autond the longth of	Example: Deriverdial offusion		
1920	OtherCV	Other Cardiovascular Event	cardiovascular events not listed that extend the length of	Example: Pericardial effusion,		
1020	GastricOutlet	Gastric Outlet Obstruction	stay of affected the patient's outcome.			
1930	GastricOutiet					

	Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
1940	Ileus	Ileus				
1950	AnastoMed	Anastomosis Requiring Medical Treatment Only	Placement of a drain under image guidance (CT scan or ultrasound) is considered medical treatment of an anastomotic leak.			
1970	DilationEsoph	Dilation Of The Esophagus		December 2013 - This includes the entire 30-day post-op period.		
1980	OtherGI	Any Other GI Event	Gastrointestinal events not listed that extended the length of stay or affected the patient's outcome.			
1990	PostopPRBC	Postoperative Packed Red Blood Cells				
April 2014	Pt has a thoracic procedu of foreign body). 10 day of right brachial artery ps which is NOT a thoracic non-thoracic surgery, the Should those blood proc database for the initial pr products NOT be capture a second procedure? The manual/specs to clarify.	re(esophagoscopy with removal s later the pt has a surgery(repair seudoaneurysm with fasciotomy) procedure. 4 days AFTER the pt has blood products. ducts be captured on the thoracic ocedure? Or should the blood d because they were given after are is nothing in the training	This is a non-analyzed procedure so no post-op events ne	ed to be answered.		
2000	PostopPRBCUnits	Postoperative Packed Red Blood Cells - Units				
2010	UTI	Urinary Tract Infection	Positive urine culture and treatment required. Do not code based on urinalysis results.			
2020	UrinRetent	Urinary Retention				
May 2014	Do straight cath's count h place?	ere or only full foley recatherizat	tions that are left in Yes, straight cath is a catheterization			
2030	DischFoley	Discharged With Foley Catheter				
2040	Empyema SurgSiteInfect	Empyema Requiring Treatment	Empyema refers to an infected pleural space requiring additional antibiotic coverage or placement of additional chest tubes/drains.	Empyema may be confirmed by thoracentesis drainage of cloudy fluid or frank pus. Fluid analysis, if performed, would likely reveal leukocytosis, pH < 7.2, glucose < 60, elevated LDH, elevated protein and positive cultures. A superficial surgical site infection		

SeqNo         ShortName         LongName         DataFieldIntent           documented infection of areas opened or manipulated during the procedure. It can involve tissue related to the Unfection         Unfection	FieldNameClarification
documented infection of areas opened or manipulated (SSI) m during the procedure. It can involve tissue related to the Infection	r leiu vanie Ciai in cation
primary or secondary surgical incision(s). It may be superficial-involving skin and subcutaneous tissue and muscle Organ space infection-involving body cavity, such as empyema or mediastinitis.	<ul> <li>) must meet the following criteria:</li> <li>ction occurs within 30 days after the ative procedure</li> <li>lves only skin and subcutaneous</li> <li>e of the incision</li> <li>ent has at least 1 of the following:</li> <li>urulent drainage from the superficial incision</li> <li>ganisms isolated from an tically obtained culture of fluid or</li> <li>e from the superficial incision least 1 of the following signs or ptoms of infection: pain or</li> <li>erness, localized swelling, redness, eat, and superficial incision is berately opened by surgeon and is are positive or not cultured. A are-negative finding does not meet criterion.</li> <li>agnosis of superficial incisional by the surgeon or attending ician.</li> <li>to report a stitch abscess (minimal mmation and discharge confined to points of suture penetration) as an extion.</li> <li>e incisional site infection involves stends into the fascial and muscle rs, report as a deep incisional SSI. sify infection that involves both rficial and deep incision sites as incisional SSI.</li> </ul>

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				A deep incisional SSI (DIP or DIS) must meet the following criterion: Infection occurs within 30 days after the operative procedure and involves deep soft tissues (e.g., fascial and muscle layers) of the incision and patient has at least 1 of the following: a. purulent drainage from the deep incision but not from the organ/space component of the surgical site b. a deep incision spontaneously dehisces or is deliberately opened by a surgeon and is culture-positive or not cultured when the patient has at least 1 of the following signs or symptoms: fever (>38°C), or localized pain or tenderness. A culture-negative finding does not meet this criterion. c. an abscess or other evidence of infection involving the deep incision is found on direct examination, during reoperation, or by histopathologic or radiologic examination d. diagnosis of a deep incisional SSI by a surgeon or attending physician. An "organ /space" surgical site infection would include empyema or mediastinitis. The diagnosis of organ space infection must meet the following criteria according to the CDC: Infection occurs within 30 days after the	
				reoperation, or by histopatholo radiologic examination d. diagnosis of a deep incisiona a surgeon or attending physicia An "organ /space" surgical site infection would include empye mediastinitis. The diagnosis of space infection must meet the f criteria according to the CDC: Infection occurs within 30 days operative procedure	

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo ShortName	LongName	DataFieldIntent	FieldNameClarification		
			and infection involves any part of the body, beyond the skin incision, fascia, or muscle layers, that is opened or manipulated during the operative procedure and patient has at least 1 of the following: a. purulent drainage from a drain that is placed through a stab wound into the organ/space b. organisms isolated from an aseptically obtained culture of fluid or tissue in the organ/space c. an abscess or other evidence of infection involving the organ/space that is found on direct examination, during reoperation, or by histopathologic or radiologic examination d. diagnosis of mediastinitis, an organ/space SSI by a surgeon or attending physician		
2070 Sepsis	Sepsis	Evidenced by positive blood cultures in the medical record.			
2080OtherInfect	Other Infection Requiring IV Antibiotics				
2090CentNeuroEvt	New Central Neurological Event				
Aprillif a patient has history of a stroke pre-operatively and has 2014 a stroke in the post-operative period. Would I mark yes to New Central Neurological Event?		Yes, it's a new event.			
2100 RecLarynParesis	Recurrent Laryngeal Nerve Paresis				
2110 Delirium	Delirium				
2120 OtherNeuro	Other Neurological Event		Example: Seizure		

StepNo         ShortName         LongName         DataFieldIntent         FieldNameClarification           2140         RenFailRIFLE         Renal Failure - RIFLE Criteria         Creatinine must rise to 3 times baseline or be ≥ 4.0 with an acute rise of at least 0.5 and/or a new requirement for dialysis.         The post operative creatinine will be used to evaluate renal function acute rise of at least 0.5 and/or a new requirement for dialysis initiated- Do Not Code as New Renal Failure.         The post operative creatinine must rise to 3 times baseline or be ≥ 4.0 with no dialysis initiated- Do Not Code as New Renal Failure.         The post operative creatinine must rise to 3 times baseline or be ≥ 4.0 with not dialysis initiated- Do Not Code as New Renal Failure.           8         View Provide Attribution called the RIFLE classification system. It is used to define grades of severity based on objective measurements.         The will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time.           8         Field NameClassifications of Loss and End-stage disease are beyond the current scope of follow-up.         Risk (R) - Increase in Serum creatinine level X 1.5 or decrease in GFR by 25%, or U O 4.0.5 mL&gh for 6 hours must reatinine level X 3.0, or serum creatinine level X 3.0, or oserum creatinine level X 3.0, or oserum creatinine level X 3.0, or serum creatinine level X 3.0, or serum	Version 2.2 of the General Thoracic Surgery Database						
2140       Renal Failure - RIFLE Criteria       Creatinine must rise to 3 times baseline or be ≥ 4.0 with an acute rise of at least 0.5 and/ or a new requirement for dialysis.       The post operative creatinine will be used to evaluate renal function according to the RIFLE criteria. The Acute Dialysis Quality Initiative, a multidisciplinary collaboration, defined a range of acute renal dysfunction called the RIFLE classification system. It is used to define grades of severity based on objective measurements.         STS will use the underlined serum creatinine values of an alyze post op =4.1       with no dialysis initiated- Do Not Code as New Renal Failure.         STS will use the underlined serum creatinine values of analyze post op renal function. GFR and urine output will not be included at this time.       STS will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time.         Renal Failure (T) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mLAgh for 12 hours       Failure (T) - Increase in serum creatinine level X 3.0 or serum creatinine level X 3.0 serum distored the serue of the server X 4 bours, or anuria for 12 hours         Failure (T) - Increase in serum creatinine level X 4 bours, or anuria for 12 hours       Failure (T) - Increase in Serum creatinine level X 3.0 servem	SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
2) 40 (kehrältKirLE 2) 40 (ke	2140		Densel Feilens DIFLE Original	$C_{\rm restriction} = 1 + 2 + 1 + 1 + 1 + 1 + 1 + 1 + 1 + 1 +$			
for dialysis. Example: pre op creatinine = 3.8 and the post op = 4.1 with no dialysis initiated- Do Not Code as New Renal Failure. Here RFLE classification system. It is used to evaluate renal dysfunction called the RFLE classification system. It is used to define grades of severity based on objective measurements. STS will use the underfined serum creatinine values to analyze post op renal failure eriteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in GFR by 25%, or U0 <0.5 mL/kg/h for 6 hours Injury (I) - Increase in GFR by 25%, or U0 <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level X 3.0, or serum creatinine level X 3.0, or serum creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%: 10 <0.3 mL/kg/h for 24 hours, or anuia for 12 hours Ends. (I) - Persistent ARF, complete loss of kidney function >4 weeks End-stage disease (L) - Loss of kidney function >4 weeks	2140	Kenfalkifle	Renai Failure - RIFLE Criteria	Creatinine must rise to 5 times baseline of $be \ge 4.0$ with on south rise of at least 0.5 and/or a new requirement	The post operative creatinine will be		
Example: pro op creatinine = 3.8 and the post op = 4.1 Example: pro op creatinine = 3.8 and the post op = 4.1 with no dialysis initiated- Do Not Code as New Renal Failure. Acute Dialysis Quality Initiative, a multidisciplinary collaboration, defined the RIFLE classification system. It is used to define grades of severity based on objective measurements. STS will use the underlined serum creatinine values to analyze post op renal failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level X 3.0, or serum creatine				for dialysis	according to the RIFLE criteria. The		
with no dialysis initiated- Do Not Code as New Renal Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Failure. Fai				Example: pre on creatinine $-3.8$ and the post on $-4.1$	Acute Dialysis Quality Initiative a		
Failure.       Failure.       Failure.       Failure.         Renal Failure.       Failure.       Failure.       Failure.         Renal Failure.       Failure.       Failure.       Failure.         Renal Failure.       Failure.       Failure.       Failure.         Failure.       Failure.       Failure.       Failure.       Failure.         Failure.       Failure.       Failure.       Failure.       Failure.       Failure.         Failure.       Failure.       Failure.       Failure. <td></td> <td></td> <td></td> <td>with no dialysis initiated. Do Not Code as New Renal</td> <td>multidisciplinary collaboration defined</td>				with no dialysis initiated. Do Not Code as New Renal	multidisciplinary collaboration defined		
the RIFLE classification system. It is used to define grades of severity based on objective measurements. STS will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time. Renal Failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 6 hours Injury (I) - Increase in serum creatinine level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks Endo-stage kidney disease (E) - Loss of kidney function >3 months				Failure	a range of acute renal dysfunction called		
used to define grades of severity based on objective measurements. STS will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time. Renal Failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 6 hours Injury (I) - Increase in serum creatinine level X 2.0 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level X 3.0, or serum creatinine level X 3.0, or serum creatinine level $\ge$ 4 mg/dL , with an acute rise of at least 0.5 or decrease in GFR by 75%; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					the RIFLE classification system. It is		
on objective measurements. STS will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time. Renal Failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 6 hours Injury (I) - Increase in serum creatinine level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level ≥4.0, or serum creatinine level ≥4.0, or decrease in GFR by 75%, IO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent AFF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					used to define grades of severity based		
STS will use the underlined serum creatinine values to analyze post op renal function. GFR and urine output will not be included at this time. Renal Failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 6 hours Injury (I) - Increase in serum creatinine level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level X 3.0, or serum creatinine level X 3.0, or serum creatinine level X 40, or decrease in GFR by 75%, iUO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					on objective measurements.		
creatinine values to analyze post op renal function. GFR and urine output will not be included at this time. Renal Failure criteria are highlighted. Classifications of Loss and End-stage disease are beyond the current scope of follow-up. Risk (R) - Increase in serum creatinine level X 1.5 or decrease in GFR by 25%, or UO <0.5 mL/kg/h for 6 hours Injury (I) - Increase in serum creatinine level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours Failure (F) - Increase in serum creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					STS will use the underlined serum		
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current scope of follow-up.         Risk (R) - Increase in serum creatinine         level X 1.5 or decrease in GFR by 25%,         or UO <0.5 mL/kg/h for 6 hours					and End-stage disease are beyond the		
Risk (ℝ) - Increase in serum creatinine         level X 1.5 or decrease in GFR by 25%,         or UO <0.5 mL/kg/h for 6 hours					current scope of follow-up.		
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Injury (I) - Increase in serum creatinine         level X 2.0 or decrease in GFR by 50%,         or UO <0.5 mL/kg/h for 12 hours					or UO <0.5 mL/kg/h for 6 hours		
level X 2.0 or decrease in GFR by 50%, or UO <0.5 mL/kg/h for 12 hours <b>Failure (F)</b> - Increase in serum creatinine level X 3.0, or serum creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%,; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours <b>Loss (L)</b> - Persistent ARF, complete loss of kidney function >4 weeks <b>End-stage kidney disease (E)</b> - Loss of kidney function >3 months					<b>Injury</b> ( <b>I</b> ) - Increase in serum creatinine		
or UO <0.5 mL/kg/h for 12 hours <b>Failure (F)</b> - Increase in serum creatinine level X 3.0, or serum creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%,; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours <b>Loss (L)</b> - Persistent ARF, complete loss of kidney function >4 weeks <b>End-stage kidney disease (E)</b> - Loss of kidney function >3 months					level X 2.0 or decrease in GFR by 50%,		
Failure (F) - Increase in serum creatinine level X 3.0, or serum creatinine level $\geq 4$ mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%,; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					or UO <0.5 mL/kg/h for 12 hours		
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creatinine level ≥4 mg/dL, with an acute rise of at least 0.5 or decrease in GFR by 75%,; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					creatinine level X 3.0, or serum $(11)$		
acute rise of at least 0.5 or decrease in GFR by 75%,; UO <0.3 mL/kg/h for 24 hours, or anuria for 12 hours <b>Loss (L)</b> - Persistent ARF, complete loss of kidney function >4 weeks <b>End-stage kidney disease (E)</b> - Loss of kidney function >3 months					creatinine level $\geq 4 \text{ mg/dL}$ , with an		
GFR by 75%,; UO <0.3 mL/kg/n for 24 hours, or anuria for 12 hours Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					acute rise of at least 0.5 or decrease in $CED$ has $750(-100) = 0.2$ and $4\pi$ for 24		
Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					UFK by $75\%$ ; UU <0.3 mL/kg/n for 24		
Loss (L) - Persistent ARF, complete loss of kidney function >4 weeks End-stage kidney disease (E) - Loss of kidney function >3 months					nours, or anuria for $12$ nours		
End-stage kidney disease (E) - Loss of kidney function >3 months					Loss (L) - Persistent AKF, complete		
kidney function >3 months					ioss of kidney function >4 weeks End-stage kidney disease (E) I ass of		
kidney function >5 months					kidney function >3 months		
Reterance					Reference.		

Training Manual Version 2.2 of the General Thoracic Surgery Database					
SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification	
				http://ccforum.com/content/8/4/R204	
2150	ChyloMed	Chylothorax Requiring Drainage/Medical Treatment Only	This indicates the use of non-operative measures to address a postoperative chylothorax and refers to cessation of oral intake, initiation of total parenteral nutrition, etc.	Chylothorax is identified by the milky appearance of pleural fluid, which, if analyzed would have triglyceride levels> 110 mg/dl	
2170	OtherSurg	Other events requiring OR with general anesthesia			
2180	UnexpectAdmitICU	Unexpected Admission To ICU	Did the patient have an unexpected admission or re- admission to the ICU in the post-operative period? If so, then the answer to this question would be "yes."		
2190	DischDt	Discharge Date	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.		
2200	MtDCStat	Discharge Status	This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.		
2210	DisLoctn	Discharge Location	If Alive, select Home; Extended Care/Transitional Care Unit/Rehab; Other Hospital; Nursing Home; Hospice; or Other.		
2230	Readm30Dis	Readmission within 30 days of Discharge	Do not include ER or clinic visits.	This is 30 days after discharge, not to be confused with 30 days after surgery, collected in the mortality field.	
Patients undergo outpatient mediastinoscopy or bronchoscopy for			Code the initial procedure, if your surgeon wishes to capture nonanalyzed procedures, on the		
staging and are then admitted for planned surgery a few weeks			short form, which has no field for readmission. Outpatient visits are not considered admissions		
later. Is this coded as a readmission?			so subsequent admission within 30 days would not be considered a readmission.		
Oct. My patient is from out of state with no PCP listed in her he 2013 Following her surgical procedure in my state, she returned way of following up to know if she was readmitted within discharge. The only represented are Xer (No. 11)			I home. I have no 1 30 days of L answer this	ient to find out whether she has been	
question?					
2240	Mt30Stat	Status 30 Days After Surgery	A process must be in place to determine patient's status at 30 days after surgery after discharge. There must be NO default to "Alive". Please see STS Database News articles on the STS Web site regarding how to collect this information.	Correction! This should be 30 days after surgery, not after discharge.	

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification		
			Documentation must be completed and/or have the ability to replicate obtaining status upon request of an auditor. Note: Data marked as "Unknown" will not show up as missing in the Data Quality Report, but will not be included in analyses. This field is Required for Record Inclusion. If missing data, the entire record will be excluded from the analysis.			
2250	MtDate	Date Of Death				
2270 2290	CTubeDis IVAntibioOrdered	Discharged With Chest Tube IV Antibiotics Ordered Within One Hour	Do not capture long term drainage devices. Indicate "Yes" if an order was documented, regardless if patient actually received any IV antibiotics. Applies to Inpatients only.	If vancomycin or a fluoroquinolone is given, the timeframe is extended to 2 hours since these must be infused at a slower rate.		
2300	IVAntibioGiven	IV Antibiotics Given Within One Hour	If "Not indicated for procedure" selected, appropriate documentation must be in medical record. Applies to Inpatients only.	If vancomycin or a fluoroquinolone is given, the timeframe is extended to 2 hours since these must be infused at a slower rate.		
June 2013	June If the patient is in ICU and already on multiple antibiotics do I code this as 2013 yes even though one is not specifically ordered for the surgery? Should these patients' antibiotics be stopped and the patient be given the prophylactic one instead or in addition?Code "no."That is a physician decision.					
2310	CephalAntiOrdered	Cephalosporin Antibiotic Ordered Prophylactic Antibiotics	Indicate "Yes" if an order was documented for a first or second-generation cephalosporin antibiotic, regardless if the patient actually received the antibiotic. If "Not indicated for procedure" or "Not indicated due to documented allergy; another appropriate antibiotic given" selected, appropriate documentation must be in medical record. Applies to Inpatients only. Indicate "Yes" if an order to discontinue was			
2520		Discontinuation Ordered	documented as per the definition. If "Not, due to documented infection" selected, substantiating documentation must be in medical record.			

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SeqNo	ShortName	LongName	DataFieldIntent	FieldNameClarification
			Applies to Inpatients only.	
2330	DVTProphylaxis	DVT Prophylaxis Measures	Applies to Inpatients only.	
2340	SmokCoun	Smoking Cessation Counseling	Choose yes, no, refused or patient is a nonsmoker.	July 2013: For the purpose of this sequence number, a "nonsmoker" can mean either someone who has never smoked or someone who is no longer smoking. If they are not a current smoker then they won't need smoking cessation counseling.