

NQF 1952: Time to Intravenous Thrombolytic Therapy – 60 min Specifications – Measure Calculation

Percent of acute ischemic stroke patients receiving intravenous alteplase therapy during the hospital stay who have a time from hospital arrival to initiation of thrombolytic therapy (door-to-needle time) of 60 minutes or less		
Denominator		
Include:	Display Data Elements (bolded and underlined)	Dictionary Data Elements (in blue)
All patients with a final clinical diagnosis of ischemic stroke who received IV alteplase at my hospital	<u>Final Clinical Dx. of Stroke:</u> Ischemic Stroke AND <u>IV alteplase initiated at this hospital:</u> Yes	gs_stroketype = 2 [Ischemic Stroke] AND gs_ivthroinit = 1 [Yes]
Exclusions: (Always remove from denominator)		
<ul style="list-style-type: none"> Age < 18 years Stroke occurred after hospital arrival (in ED/Obs/inpatient) Patients whose date/time of ED arrival and/or date/time of IV alteplase administration is blank, unknown, or MM/DD/YYYY only. Patients with a negative calculated time difference Patients with a Date Last Known Well, but no time Last Known Well Patients that receive IV alteplase greater than 4.5 hours after Last Known Well Patients who received IV alteplase at an outside hospital or by EMS/Mobile Stroke Unit Clinical Trial 	<p><u>Age</u> < 18 OR</p> <p><u>Patient location when stroke symptoms discovered:</u> Stroke occurred after hospital arrival--In ED/Obs/Inpatient OR</p> <p><u>Arrival Date/Time</u> is blank, unknown, or MM/DD/YYYY only OR</p> <p><u>Date/time IV alteplase initiated</u> is blank, unknown, or MM/DD/YYYY only OR</p> <p><u>Date/time IV alteplase initiated</u> < <u>Arrival Date/Time</u> OR</p> <p><u>Date/Time Last Known Well:</u> Date included but time is blank or unknown, OR</p>	<p>gs_age < 18 OR</p> <p>gs_symptomlocation = 4 [Stroke occurred after hospital arrival--In ED/Obs/Inpatient] OR</p> <p>jc_arrdatetime is null OR</p> <p>jc_arrdatetime_precision = 0 [Unknown] OR = 3 [MM/DD/YYYY] OR</p> <p>jc_ivthrodttm) is null OR</p> <p>jc_ivthrodttm_precision = 0 [Unknown] OR = 3 [MM/DD/YYYY] OR</p> <p>jc_ivthrodttm < jc_arrdatetime OR</p> <p>gs_lastknownwell_precision = 3 [MM/DD/YYYY]</p>

	<p><u>Date/time IV alteplase initiated</u> - <u>Date/Time Last Known Well</u> > 4.5hours OR</p> <p><u>IV alteplase at an outside hospital or EMS / Mobile Stroke Unit</u>: Yes OR</p> <p><u>During this hospital stay, was the patient enrolled in a clinical trial in which patients with the same condition as the measure set were being studied</u>: Yes</p>	<p>OR</p> <p>(<u>gs_lastknownwell_precision</u>) = 5 [MM/DD/YYYY HH:MM] AND <u>jc_ivthrodtm</u> - <u>gs_lastknownwell</u> > 270) OR</p> <p><u>gs_ivtpaoutside</u> = 1 [Yes] OR <u>jc_clinical</u> = 1 [Yes]</p>
Exceptions: (Remove from denominator if present and numerator is not met)		
Patients who received IV alteplase greater than 60 minutes after arrival and have a documented Eligibility or Medical Reason for delay in treatment	<p><u>If IV alteplase was initiated greater than 60 minutes after arrival, documented Eligibility or Medical reason(s) for delay</u>: Yes AND</p> <p>(<u>Eligibility Reason</u>: is not blank OR <u>Medical Reason</u>: is not blank)</p>	<p><u>ivtpadelay</u> = 1 [Yes] AND</p> <p>(<u>ivtpadelay_er</u> != null OR <u>ivtpadelay_mr</u> != null)</p>
Numerator		
Patients who receive IV alteplase at my hospital within 60 minutes after arrival	<p><u>Date/time IV alteplase initiated</u> - <u>Arrival Date/Time</u>: ≤ 60 minutes</p>	<u>jc_ivthrodtm</u> - <u>jc_arrdatetime</u> ≤ 60 minutes

Patient ID: _____		Bold Question = Required	
DEMOGRAPHICS <i>Demographics Tab</i>			
Gender	<input type="radio"/> Male <input type="radio"/> Female <input type="radio"/> Unknown		
Date of Birth: ____/____/____	Age: _____		
Zip Code: _____ - _____	<input type="checkbox"/> Homeless		
Health Insurance Status	<input type="checkbox"/> Medicare <input type="checkbox"/> Medicaid <input type="checkbox"/> Private/VA/Champus/Other Insurance <input type="checkbox"/> Self Pay / No Insurance <input type="checkbox"/> ND		
RACE AND ETHNICITY			
Race (Select all that apply):	<input type="checkbox"/> White <input type="checkbox"/> UTD <input type="checkbox"/> American Indian/Alaska Native <input type="checkbox"/> Asian		<input type="checkbox"/> Black or African American <input type="checkbox"/> Native Hawaiian or Pacific Islander
	[if Asian selected] <input type="checkbox"/> Asian Indian <input type="checkbox"/> Chinese <input type="checkbox"/> Filipino <input type="checkbox"/> Japanese <input type="checkbox"/> Korean <input type="checkbox"/> Vietnamese <input type="checkbox"/> Other Asian		[if native Hawaiian or pacific islander selected] <input type="checkbox"/> Native Hawaiian <input type="checkbox"/> Guamanian or Chamorro <input type="checkbox"/> Samoan <input type="checkbox"/> Other Pacific Islander
Hispanic Ethnicity:	<input type="radio"/> Yes <input type="radio"/> No/UTD		
If Yes,	<input type="checkbox"/> Mexican, Mexican American, Chicano/a <input type="checkbox"/> Puerto Rican <input type="checkbox"/> Cuban <input type="checkbox"/> Another Hispanic, Latino or Spanish Origin		
ADMIN <i>Admin Tab</i>			
Final clinical diagnosis related to stroke	<input type="radio"/> Ischemic Stroke <input type="radio"/> Intracerebral Hemorrhage <input type="radio"/> Transient Ischemic Attack (<24 hours) <input type="radio"/> Subarachnoid Hemorrhage <input type="radio"/> Stroke not otherwise specified <input type="radio"/> <input type="radio"/> No stroke related diagnosis <input type="radio"/> <input type="radio"/> Elective Carotid Intervention only		
If not Stroke Related Diagnosis:	<input type="radio"/> Migraine <input type="radio"/> Electrolyte or metabolic imbalance <input type="radio"/> Seizure <input type="radio"/> Functional disorder <input type="radio"/> Delirium <input type="radio"/> Other <input type="radio"/> <input type="radio"/> Uncertain		
Was the Stroke etiology documented in the patient medical record:		<input type="radio"/> Yes <input type="radio"/> No	
Select documented stroke etiology (select all that apply):	<input type="radio"/> 1: Large-artery atherosclerosis (e.g., carotid or basilar stenosis) <input type="radio"/> 2: Cardioembolism (e.g., atrial fibrillation/flutter, prosthetic heart valve, recent MI) <input type="radio"/> 3: Small-vessel occlusion (e.g., subcortical or brain stem lacunar infarction <1.5 cm) <input type="radio"/> 4: Stroke of other determined etiology (e.g., dissection, vasculopathy, hypercoagulable or hematologic disorders. <input type="radio"/> Dissection <input type="radio"/> Hypercoagulability <input type="radio"/> Other <input type="radio"/> 5: Cryptogenic stroke (stroke of undetermined etiology) <input type="radio"/> Multiple potential etiologies identified <input type="radio"/> Stroke of undetermined etiology <input type="radio"/> Unspecified		
When is the earliest documentation of comfort measures only?	<input type="radio"/> Day 0 or 1 <input type="radio"/> Day 2 or after <input type="radio"/> Timing unclear <input type="radio"/> Not Documented/UTD		
Arrival Date/Time: ____/____/____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown	Admit Date:	____/____/____

Case Record Form

Active Form Groups: Stroke, STK (StrokeCM), Comprehensive, Coverdell

Updated April 2019

Not Admitted:	<input type="radio"/> Yes, not admitted <input type="radio"/> No, patient admitted as in patient	Reason Not Admitted:	<input type="radio"/> Transferred from your ED to another acute care hospital <input type="radio"/> Discharged directly from ED to home or other location that is not an acute care hospital <input type="radio"/> Left from ED AMA <input type="radio"/> Died in ED <input type="radio"/> Discharged from observation status without an inpatient admission <input type="radio"/> other
If patient transferred from your ED to another hospital, specify hospital name	[Select hospital name from picker list] <input type="checkbox"/> Hospital not on list <input type="checkbox"/> Hospital not documented		
Select reason(s) for why patient transferred	<input type="checkbox"/> Evaluation for IV alteplase up to 4.5 hours <input type="checkbox"/> Post Management of IV alteplase (e.g. Drip and Ship) <input type="checkbox"/> Evaluation for Endovascular thrombectomy <input type="checkbox"/> Advanced stroke care (e.g., Neurocritical care, surgical or other time critical therapy) <input type="checkbox"/> Patient/family request <input type="checkbox"/> Other advanced care (not stroke related) <input type="checkbox"/> Not documented		
Discharge Date:	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only		
Documented reason for delay in transfer to referral facility?	<input type="radio"/> Yes <input type="radio"/> No		
Specific reason for delay documented in transfer patient (check all that apply):	<input type="checkbox"/> Social/religious <input type="checkbox"/> Initial refusal <input type="checkbox"/> Care team unable to determine eligibility <input type="checkbox"/> Management of concomitant emergent/acute conditions such as cardiopulmonary arrest, respiratory failure (requiring intubation) <input type="checkbox"/> Investigational or experimental protocol for reperfusion <input type="checkbox"/> Delay in stroke diagnosis * <input type="checkbox"/> In-hospital time delay * <input type="checkbox"/> Equipment-related delay * <input type="checkbox"/> Need for additional imaging* <input type="checkbox"/> Catheter lab not available* <input type="checkbox"/> Other *		
For patients discharged on or after 04/01/2011: What was the patient's discharge disposition on the day of discharge?	<input type="checkbox"/> 1 – Home <input type="checkbox"/> 2 – Hospice – Home <input type="checkbox"/> 3 – Hospice – Health Care Facility <input type="checkbox"/> 4 – Acute Care Facility <input type="checkbox"/> 5 – Other Health Care Facility <input type="checkbox"/> 6 – Expired <input type="checkbox"/> 7 – Left Against medical Advise / AMA <input type="checkbox"/> 8 – Not Documented or Unable to Determine (UTD)		
If Other Health Care Facility	<input type="radio"/> Inpatient Rehabilitation Facility (IRF) <input type="radio"/> Skilled Nursing Facility (SNF) <input type="radio"/> Intermediate Care facility (ICF) <input type="radio"/> Other <input type="radio"/> Long Term Care Hospital (LTCH)		
DIAGNOSIS CODE		<i>Clinical Codes Tab</i>	
ICD-9CM or ICD-10-CM Principal Diagnosis Code ICD-9CM or ICD-10-CM Other Diagnosis Codes ICD-9-CM or ICD-10-PCS Principal Procedure Code ICD-9-CM or ICD-10-PCS Other Procedure Codes ICD-9-CM Discharge Diagnosis Related to Stroke ICD-10-CM Discharge Diagnosis Related to Stroke No Stroke or TIA Related ICD-9-CM Code Present No Stroke or TIA Related ICD-10-CM Code Present	_____ _____ _____ <input type="checkbox"/> _____ <input type="checkbox"/> _____		

ARRIVAL AND ADMISSION INFORMATION		Admission Tab
During this hospital stay, was the patient enrolled in a clinical trial in which patients with the same condition as the measure set were being studied (i.e. STK,VTE)?		<input type="radio"/> Yes <input type="radio"/> No
Was this patient admitted for the sole purpose of performance of elective carotid intervention?		<input type="radio"/> Yes <input type="radio"/> No
Patient location when stroke symptoms discovered	<input type="radio"/> Not in a healthcare setting <input type="radio"/> Outpatient healthcare setting <input type="radio"/> Another acute care facility <input type="radio"/> Stroke occurred after hospital arrival (in ED/Obs/inpatient) <input type="radio"/> Chronic health care facility <input type="radio"/> ND or Cannot be determined	
How patient arrived at your hospital	<input type="radio"/> EMS from home/scene <input type="radio"/> Mobile Stroke Unit <input type="radio"/> Private Transportation/Taxi/Other from home/scene <input type="radio"/> Transfer from another hospital <input type="radio"/> ND or Unknown	
Referring hospital discharge Date/ Time	___/___/___:___ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown	
If transferred from another hospital, specify hospital name	[Select hospital name from picker list] <input type="checkbox"/> Hospital not on list <input type="checkbox"/> Hospital not documented	
Referring hospital arrival date/ time	___/___/___:___ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown	
If patient transferred to your hospital, select transfer reason(s)	<input type="checkbox"/> Evaluation for IV alteplase up to 4.5 hours <input type="checkbox"/> Post Management of IV alteplase (e.g. Drip and Ship) <input type="checkbox"/> Evaluation for Endovascular thrombectomy <input type="checkbox"/> Advanced stroke care (e.g., Neurocritical care, surgical or other time critical therapy) <input type="checkbox"/> Patient/family request <input type="checkbox"/> Other advanced care (not stroke related) <input type="checkbox"/> Not documented	
Was the patient an ED patient at the facility?	<input type="radio"/> Yes <input type="radio"/> No	
Was the patient a direct admission to the hospital?	<input type="radio"/> Yes <input type="radio"/> No	
Where patient first received care at your hospital	<input type="radio"/> Emergency Department / Urgent Care <input type="radio"/> Direct Admit, not through ED <input type="radio"/> Imaging suite <input type="radio"/> ND or Cannot be determined	
Advanced Notification by EMS or MSU?	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> N/A	
Where was the patient cared for and by whom? Check all that apply.	<input type="checkbox"/> Neuro Admit <input type="checkbox"/> Other Service Admission <input type="checkbox"/> Stroke Consult <input type="checkbox"/> No Stroke Consult <input type="checkbox"/> In Stroke Unit <input type="checkbox"/> Not in Stroke Unit	
Physician / Provider NPI:		
MEDICAL HISTORY		
Previously known medical hx of:	<input type="checkbox"/> None <input type="checkbox"/> Drugs/Alcohol <input type="checkbox"/> Previous Stroke <input type="checkbox"/> Atrial Fib/Flutter <input type="checkbox"/> Dyslipidemia <input type="checkbox"/> Ischemic stroke <input type="checkbox"/> CAD/Prior MI <input type="checkbox"/> Family History of Stroke <input type="checkbox"/> ICH <input type="checkbox"/> Carotid Stenosis <input type="checkbox"/> HF <input type="checkbox"/> SAH <input type="checkbox"/> Current Pregnancy (up to 6 weeks post partum) <input type="checkbox"/> HRT <input type="checkbox"/> Not Specified <input type="checkbox"/> DVT/PE <input type="checkbox"/> Hypertension <input type="checkbox"/> Previous TIA <input type="checkbox"/> Depression <input type="checkbox"/> Migraine <input type="checkbox"/> Prosthetic Heart Valve <input type="checkbox"/> Diabetes Mellitus <input type="checkbox"/> Obesity/Overweight <input type="checkbox"/> PVD <input type="checkbox"/> <input type="checkbox"/> Renal insufficiency – chronic <input type="checkbox"/> <input type="checkbox"/> Sickle Cell <input type="checkbox"/> <input type="checkbox"/> Sleep Apnea <input type="checkbox"/> <input type="checkbox"/> Smoker	
Ambulatory status prior to current event	<input type="radio"/> Able to ambulate independently (no help from another person) w/ or w/o device <input type="radio"/> With assistance (from person) <input type="radio"/> Unable to ambulate <input type="radio"/> ND	

Pre-stroke Modified Rankin Score	<input type="radio"/> 0 – No symptoms at all <input type="radio"/> 1 – No significant disability; despite symptoms; able to carry out all usual duties and activities <input type="radio"/> 2 – Slight disability; unable to perform all previous activities, but able to look after own affairs without assistance <input type="radio"/> 3 – Moderate disability; requiring some help, but able to walk without assistance <input type="radio"/> 4 – Moderately severe disability; unable to walk without assistance and unable to attend to own bodily needs without assistance <input type="radio"/> 5 – Severe disability; bedridden, incontinent, and requiring constant nursing care and attention <input type="radio"/> 6 – Dead <input type="radio"/> Unknown/ ND
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DIAGNOSIS & EVALUATION

Symptom Duration if diagnosis of Transient Ischemic Attack (less than 24 hours)	<input type="radio"/> Less than 10 minutes <input type="radio"/> 10 – 59 minutes <input type="radio"/> > = 60 minutes <input type="radio"/> ND
Had stroke symptoms resolved at time of presentation?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> ND
Initial NIH Stroke Scale	<input type="radio"/> Yes <input type="radio"/> No/ND
If yes:	<input type="radio"/> Actual <input type="radio"/> Estimate from record <input type="radio"/> ND
Total Score:	_____ (refer to web program for questions)
^What is the first NIHSS score obtained prior to or after hospital arrival?	_____ <input type="checkbox"/> UTD
^Is there documentation that an initial NIHSS score was done at this hospital	<input type="radio"/> Yes <input type="radio"/> No
^What is the date and time that the NIHSS score was first performed at this hospital?	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
NIHSS score obtained from transferring facility:	_____ <input type="radio"/> ND
Initial exam findings (Select all that apply)	<input type="checkbox"/> Weakness/Paresis <input type="checkbox"/> Altered Level of Consciousness <input type="checkbox"/> Disturbance <input type="checkbox"/> Aphasia/Language <input type="checkbox"/> Other neurological signs/symptoms <input type="checkbox"/> No neurological signs/symptoms <input type="checkbox"/> ND
Ambulatory status on admission	<input type="radio"/> Able to ambulate independently (no help from another person) w/ or w/o device <input type="radio"/> With assistance (from person) <input type="radio"/> Unable to ambulate <input type="radio"/> ND

HEMORRHAGIC STROKE SCALES

^First Glasgow Coma Scale (GCS)	Eye _____	Verbal _____	<input type="checkbox"/> Intubated	Motor _____	Total GCS _____ <input type="checkbox"/> ND
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SUBARACHNOID HEMORRHAGE (SAH)

^Is there documentation any time during the hospital stay that the hemorrhage was non-aneurysmal or due to head trauma?	<input type="radio"/> Yes <input type="radio"/> No
^Was an initial Hunt and Hess scale done at this hospital?	<input type="radio"/> Yes <input type="radio"/> No
^If yes, Hunt and Hess score:	_____
^What is the date and time that the Hunt and Hess Scale was first performed at this hospital?	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^WFNS SAH Grading Scale	_____

INTRACEREBRAL HEMORRHAGE (ICH)

^Was an initial ICH score done at this hospital?	<input type="radio"/> Yes <input type="radio"/> No
^If yes, ICH score:	_____
^What is the date and time that the ICH score was first performed at this hospital?	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^^FUNC Score (ICH)	_____

MEDICATION PRIOR TO ADMISSION

No medications prior to admission	<input type="checkbox"/>
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Antiplatelet or Anticoagulant Medication(s):		<input type="checkbox"/> Yes <input type="checkbox"/> No/ND	
<input type="checkbox"/> Antiplatelet Medication <input type="radio"/> aspirin <input type="radio"/> aspirin/dipyridamole (Aggrenox) <input type="radio"/> clopidogrel (Plavix) <input type="radio"/> prasugrel (Effient) <input type="radio"/> ticagrelor (Brilinta) <input type="radio"/> ticlopidine (Ticlid) <input type="radio"/> Other Antiplatelet		<input type="checkbox"/> Anticoagulant Medication <input type="radio"/> apixaban (Eliquis) <input type="radio"/> argatroban <input type="radio"/> dabigatran (Pradaxa) <input type="radio"/> desirudin (Iprivask) <input type="radio"/> endoxaban (Savaysa) <input type="radio"/> fondaparinux (Arixtra) <input type="radio"/> full dose LMW heparin <input type="radio"/> lepirudin (Refludan) <input type="radio"/> rivaroxaban (Xarelto) <input type="radio"/> unfractionated heparin IV <input type="radio"/> warfarin (Coumadin) <input type="radio"/> other Anticoagulant	
Antihypertensive	<input type="radio"/> Yes	<input type="radio"/> No/ND	
Cholesterol-Reducer	<input type="radio"/> Yes	<input type="radio"/> No/ND	
Diabetic medication	<input type="radio"/> Yes	<input type="radio"/> No/ND	
Antidepressant medication	<input type="radio"/> Yes	<input type="radio"/> No/ND	
SYMPTOM TIMELINE		Hospitalization Tab	
Date/Time Patient last known to be well?		<input type="checkbox"/> Time of Discovery same as Last Known well	Date/Time of discovery of stroke symptoms?
____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown			____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
Comments:			
BRAIN IMAGING			
Brain imaging completed at your hospital for this episode of care?	<input type="radio"/> Yes <input type="checkbox"/> CT <input type="checkbox"/> MRI <input type="radio"/> No/ND <input type="checkbox"/> ONC	Date/Time Brain Imaging First Initiated at your hospital:	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
Interpretation of first brain image after symptom onset, done at any facility:		<input type="radio"/> Acute Hemorrhage <input type="radio"/> No Acute Hemorrhage <input type="radio"/> Not Available	
Was acute Vascular or perfusion imaging (e.g. CTA, MRA, DSA) performed at your hospital?	<input type="radio"/> Yes <input type="radio"/> No	Date/Time 1 st vessel or perfusion imaging initiated at your hospital:	____/____/____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
If yes, type of vascular imaging (select all that apply)	<input type="checkbox"/> CTA <input type="checkbox"/> MR Perfusion <input type="checkbox"/> CT Perfusion <input type="checkbox"/> DSA (catheter angiography) <input type="checkbox"/> MRA <input type="checkbox"/> Image type not documented		
Was a target lesion (large vessel occlusion) visualized?	<input type="radio"/> Yes <input type="radio"/> No		
If yes, select site of large vessel occlusion (select all that apply):	<input type="checkbox"/> ICA <input type="checkbox"/> Intracranial ICA <input type="checkbox"/> Cervical ICA <input type="checkbox"/> Other/UTD	<input type="checkbox"/> MCA <input type="checkbox"/> M1 <input type="checkbox"/> M2 <input type="checkbox"/> Other/UTD	<input type="checkbox"/> Basilar <input type="checkbox"/> Other cerebral artery branch <input type="checkbox"/> Vertebral Artery
ADDITIONAL TIME TRACKER			
Date/Time Stroke Team Activated:	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time Stroke Team Arrived:	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown
Date/Time of ED Physician Assessment:	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time Neurosurgical services consult:	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown

Date/Time Brain Imaging Ordered: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time Brain Imaging Interpreted: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown
Date/Time IV alteplase Ordered: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A		
Date/Time Lab Tests Ordered: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time lab Tests Completed: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown
Date/Time ECG Ordered: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time ECG Completed: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown
Date/Time Chest X-ray Ordered: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown <input type="radio"/> N/A	Date/Time Chest X-ray Completed: ____/____/____ ____:____	Select one option <input type="radio"/> MM/DD/YYYY HH:MM <input type="radio"/> MM/DD/YYYY <input type="radio"/> Unknown
Additional Comments:			

IV THROMBOLYTIC THERAPY

IV alteplase initiated at this hospital?	<input type="radio"/> Yes <input type="radio"/> No	Date/Time IV alteplase initiated:	____/____/____ ____:____
Documented exclusions (Contraindications or Warnings) for not initiating IV thrombolytic in the 0-3hr treatment window?	<input type="radio"/> Yes <input type="radio"/> No		
Documented Contraindications or Warnings for not initiating IV thrombolytic in the 3-4.5hr treatment window?	<input type="radio"/> Yes <input type="radio"/> No		

SHOW ALL

If yes, documented exclusions for 0 -3-hour treatment window or 3 – 4.5 treatment window, select reason for exclusion.

For discharges on or after 1 April 2016

Exclusion Criteria (contraindications) 0-3 hr treatment window. Select all that apply:

- C1: Elevated blood pressure (systolic > 185 mm Hg or diastolic > 110 mm Hg) despite treatment
- C2: Recent intracranial or spinal surgery or significant head trauma, or prior stroke in previous 3 months
- C3: History of previous intracranial hemorrhage, intracranial neoplasm, arteriovenous malformation, or aneurysm
- C4: Active internal bleeding
- C5: Acute bleeding diathesis (low platelet count, increased PTT, INR >= 1.7 or use of NOAC)
- C6: Symptoms suggest subarachnoid hemorrhage
- C7: CT demonstrates multi-lobar infarction (hypodensity >1/3 cerebral hemisphere)
- C8: Arterial puncture at non-compressible site in previous 7 days
- C9: Blood glucose concentration <50 mg/dL (2.7 mmol/L)

Relative Exclusion Criteria (Warnings) 0-3 hr treatment window. Select all that apply:

- W1: Care-team unable to determine eligibility
- W2: IV or IA thrombolysis/thrombectomy at an outside hospital prior to arrival
- W3: Life expectancy < 1 year or severe co-morbid illness or CMO on admission
- W4: Pregnancy
- W5: Patient/family refusal
- W6: Rapid improvement
- W7: Stroke severity too mild

- W8: Recent acute myocardial infarction (within previous 3 months)
- W9: Seizure at onset with postictal residual neurological impairments
- W10: Major surgery or serious trauma within previous 14 days
- W11: Recent gastrointestinal or urinary tract hemorrhage (within previous 21 days)

Exclusion Criteria (contraindications) 3-4.5 hr treatment window. Select all that apply:

- C1: Elevated blood pressure (systolic > 185 mm Hg or diastolic > 110 mm Hg) despite treatment
- C2: Recent intracranial or spinal surgery or significant head trauma, or prior stroke in previous 3 months
- C3: History of previous intracranial hemorrhage, intracranial neoplasm, arteriovenous malformation, or aneurysm
- C4: Active internal bleeding
- C5: Acute bleeding diathesis (low platelet count, increased PTT, INR ≥ 1.7 or use of NOAC)
- C6: Symptoms suggest subarachnoid hemorrhage
- C7: CT demonstrates multi-lobar infarction (hypodensity >1/3 cerebral hemisphere)
- C8: Arterial puncture at non-compressible site in previous 7 days
- C9: Blood glucose concentration <50 mg/dL (2.7 mmol/L)

Relative Exclusion Criteria (Warnings) 3-4.5 hr treatment window. Select all that apply:

- W1: Care-team unable to determine eligibility
- W2: IV or IA thrombolysis/thrombectomy at an outside hospital prior to arrival
- W3: Life expectancy < 1 year or severe co-morbid illness or CMO on admission
- W4: Pregnancy
- W5: Patient/family refusal
- W6: Rapid improvement
- W7: Stroke severity too mild
- W8: Recent acute myocardial infarction (within previous 3 months)
- W9: Seizure at onset with postictal residual neurological impairments
- W10: Major surgery or serious trauma within previous 14 days
- W11: Recent gastrointestinal or urinary tract hemorrhage (within previous 21 days) Additional Relative Exclusion Criteria 3-4.5 hr treatment window. Select all that apply:
 - AW1: Age > 80
 - AW2: History of both diabetes and prior ischemic stroke
 - AW3: Taking an oral anticoagulant regardless of INR
 - AW4: Severe Stroke (NIHSS > 25)

Other Reasons (Hospital-related or other factors) 0-3-hour treatment window.

- Delay in Patient Arrival
- In-hospital Time Delay
- Delay in Stroke diagnosis
- No IV access
- Advanced Age
- Stroke too severe
- Other – requires specific reason to be entered in the PMT when this option is selected.

Other Reasons (Hospital-related or other factors) 3-4.5-hour treatment window.

- Delay in Patient Arrival
- In-hospital Time Delay
- Delay in Stroke diagnosis
- No IV access
- Other – requires specific reason to be entered in the PMT when this option is selected

If IV alteplase was initiated greater than 60 minutes after hospital arrival, were Eligibility or Medical reason(s) documented as the cause for delay:	○ Yes	○ No
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If IV alteplase was initiated greater than 45 minutes after hospital arrival, were Eligibility or Medical reason(s) documented as the cause for delay:	○ Yes	○ No
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If IV alteplase was initiated greater than 30 minutes after hospital arrival, were Eligibility or Medical reason(s) documented as the cause for delay:	○ Yes	○ No
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Eligibility Reason(s):	<input type="checkbox"/> Social/Religious <input type="checkbox"/> Initial refusal
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	<input type="checkbox"/> Care-team unable to determine eligibility <input type="checkbox"/> Specify eligibility reason: _____
Medical Reason(s):	<input type="checkbox"/> Hypertension requiring aggressive control with IV medications <input type="checkbox"/> Further diagnostic evaluation to confirm stroke for patients with hypoglycemia (blood glucose < 50), seizures, or major metabolic disorders <input type="checkbox"/> Management of concomitant emergent/acute conditions such as cardiopulmonary arrest, respiratory failure (requiring intubation) <input type="checkbox"/> Investigational or experimental protocol for thrombolysis <input type="checkbox"/> Specify medical reason: _____
Hospital Related or Other Reason(s):	<input type="checkbox"/> Delay in stroke diagnosis <input type="checkbox"/> In-hospital time delay <input type="checkbox"/> Equipment-related delay <input type="checkbox"/> Other _____
IV alteplase at an outside hospital or Mobile Stroke Unit?	<input type="radio"/> Yes <input type="radio"/> No
Investigational or experimental protocol for thrombolysis?	<input type="radio"/> Yes <input type="radio"/> No If yes, specify _____
Additional Comments Related to Thrombolytics:	

ENDOVASCULAR THERAPY

Is there documentation of LVO in the medical record?	<input type="radio"/> Yes <input type="radio"/> No
Is there documentation in the medical record that the patient is eligible for MER therapy or a mechanical thrombectomy procedure?	<input type="radio"/> Yes <input type="radio"/> No
Catheter-based stroke treatment at this hospital?	<input type="radio"/> Yes <input type="radio"/> No
IA alteplase or MER Initiation Date/Time	____/____/____ ____:____ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
Catheter-based stroke treatment at outside hospital?	<input type="radio"/> Yes <input type="radio"/> No

Note, if your hospital is collecting data for the Comprehensive Stroke Center and/or Mechanical Endovascular Reperfusion measure set, please ensure you complete additional data entry on the Advanced Stroke Care.

COMPLICATIONS

Complications of Reperfusion Therapy (Thrombolytic or MER)	<input type="checkbox"/> Symptomatic Intracranial hemorrhage <36 hours <input type="checkbox"/> Life threatening, serious systemic hemorrhage <36 hours <input type="checkbox"/> UTD	<input type="checkbox"/> Other serious complications <input type="checkbox"/> No serious complications
If bleeding complications occur in patient after IV alteplase:	<input type="checkbox"/> Symptomatic hemorrhage detected prior to patient transfer <input type="checkbox"/> Symptomatic hemorrhage detected only after patient transfer	<input type="checkbox"/> Unable to determine <input type="checkbox"/> N/A

OTHER IN-HOSPITAL TREATMENT AND SCREENING

Dysphagia Screening			
Patient NPO throughout the entire hospital stay?	<input type="radio"/> Yes <input type="radio"/> No		
Was patient screened for dysphagia prior to any oral intake including water or medications?	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC		
If yes, Dysphagia screening results:	<input type="radio"/> Pass <input type="radio"/> Fail <input type="radio"/> ND		
Treatment for Hospital-Acquired Pneumonia	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> NC		

VTE Interventions	<input type="checkbox"/> 1- Low dose unfractionated heparin (LDUH)	<input type="checkbox"/> 7- Venous foot pumps (VFP)
	<input type="checkbox"/> 2- Low molecular weight heparin (LMWH)	<input type="checkbox"/> 8-Oral Factor Xa Inhibitor
	<input type="checkbox"/> 3- Intermittent pneumatic compression devices (IPC)	<input type="checkbox"/> 9- Aspirin
	<input type="checkbox"/> 4- Graduated compression stockings (GCS)	<input type="checkbox"/> A- None of the above or ND
	<input type="checkbox"/> 5- Factor Xa Inhibitor	
	<input type="checkbox"/> 6- Warfarin	
What date was the initial VTE prophylaxis administered after hospital admission?		____/____/____ <input type="checkbox"/> Unknown
Is there physician/APN/PA or pharmacist documentation why VTE prophylaxis was not administered at hospital admission?		<input type="radio"/> Yes <input type="radio"/> No
For discharges on or after 01/01/2013: Is there physician/APN/PA documentation why Oral Factor Xa Inhibitor was administered for VTE prophylaxis?		<input type="radio"/> Yes <input type="radio"/> No
Other Therapeutic Anticoagulation	<input type="checkbox"/> apixaban (Eliquis) <input type="checkbox"/> argatroba <input type="checkbox"/> dabigatran (Pradaxa)	<input type="checkbox"/> desirrudin (Iprivask) <input type="checkbox"/> endoxaban (Savaysa) <input type="checkbox"/> lepirudin (Refludan) <input type="checkbox"/> rivaroxaban (Xarelto) <input type="checkbox"/> unfractionated heparin IV <input type="checkbox"/> other anticoagulant
Was DVT or PE documented?		<input type="radio"/> Yes <input type="radio"/> No/ND
Was antithrombotic therapy administered by the end of hospital day 2?		<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC
If yes, select all that apply		<input type="checkbox"/> Antiplatelet <input type="checkbox"/> Anticoagulant
Is there documentation by a physician/advanced practice nurse/physician assistant (physician/APN,PA) or pharmacist in the medical record of a reason for not administering antithrombotic therapy by end of hospital day 2?		<input type="radio"/> Yes <input type="radio"/> No
Was patient treated for a urinary tract infection (UTI) during this admission?		<input type="radio"/> Yes <input type="radio"/> No
If patient was treated for a UTI, did the patient have a Foley catheter during this admission?		<input type="checkbox"/> Yes, patient had catheter in place on arrival <input type="checkbox"/> Yes, but only after admission <input type="checkbox"/> No <input type="checkbox"/> Unable to determine
MEASUREMENTS (first measurement upon presentation to your hospital)		
Total Chol: _____ mg/dl	Triglycerides: _____ mg/dl	HDL: _____ mg/dl
		LDL: _____ mg/dl
		<input type="checkbox"/> Lipids: NC <input type="checkbox"/> Lipids: ND
A ₁ C: _____ % A ₁ C <input type="checkbox"/> ND	Blood Glucose (required if patient received IV alteplase): <input type="checkbox"/> ND _____ mg/dl <input type="checkbox"/> Too Low <input type="checkbox"/> Too High	
Serum Creatine: _____ <input type="checkbox"/> ND	^What is the first platelet count obtained prior to or after hospital arrival? _____	
INR: _____ <input type="checkbox"/> ND <input type="checkbox"/> NC		
^Is there documentation in the medical record that the INR value performed closest to hospital arrival was greater than 1.4?		<input type="radio"/> Yes <input type="radio"/> No
Vital Signs:	Heart Rate (beats per minute): _____ bpm ^What is the first blood pressure obtained prior to or after hospital arrival? (required if patient received IV alteplase) _____/_____ <input type="checkbox"/> Vital signs UTD	
Height: _____	<input type="radio"/> in <input type="radio"/> cm <input type="radio"/> ND	
Weight: _____	<input type="radio"/> lbs <input type="radio"/> kg <input type="radio"/> ND	
Waist Circumference: _____	<input type="radio"/> in <input type="radio"/> cm <input type="radio"/> ND	
BMI: _____	<input type="checkbox"/> ND	
CATHETER-BASED/ENDOVASCULAR STROKE TREATMENT		
<i>Advanced stroke Care Tab</i>		
^Is there documentation that the route of alteplase administration was intra-arterial (IA)?		<input type="radio"/> Yes <input type="radio"/> No
^Is there documentation that IA thrombolytic therapy was initiated at this hospital?		<input type="radio"/> Yes <input type="radio"/> No

^What is the date and time that IA thrombolytic therapy was initiated for this patient at this hospital?	____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^Is there documentation in the medical record that the first endovascular treatment procedure was initiated greater than 8 hours after arrival at this hospital?		<input type="radio"/> Yes <input type="radio"/> No
^Is there documentation of skin puncture at this hospital to access the arterial site selected for endovascular treatment of a cerebral artery occlusion?		<input type="radio"/> Yes <input type="radio"/> No
^What is the date and time of skin puncture at this hospital to access the arterial site selected for endovascular treatment of a cerebral artery occlusion?	____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^Did the patient receive intravenous (IV) alteplase at this hospital or a transferring hospital prior to receiving intra-arterial (IA) alteplase or mechanical reperfusion therapy at this hospital?		<input type="radio"/> Yes <input type="radio"/> No
^^Was a mechanical endovascular reperfusion procedure attempted during this episode of care (at this hospital)?		<input type="radio"/> Yes <input type="radio"/> No
^Was a mechanical thrombectomy procedure attempted but unsuccessful or aborted before removal of the LVO?		<input type="radio"/> Yes <input type="radio"/> No
^^Are reasons for not performing mechanical endovascular reperfusion therapy documented?		<input type="radio"/> Yes <input type="radio"/> No
^^Reasons for not performing mechanical endovascular reperfusion therapy (select all that apply):	<input type="checkbox"/> Significant pre-stroke disability (pre-stroke mRS > 1) <input type="checkbox"/> No evidence of proximal occlusion <input type="checkbox"/> NIHSS <6 <input type="checkbox"/> Brain imaging not favorable/hemorrhage transformation (ASPECTS score <6) <input type="checkbox"/> Groin puncture could not be initiated within 6 hours of symptom onset <input type="checkbox"/> Anatomical reason - unfavorable vascular anatomy that limits access to the occluded artery <input type="checkbox"/> Patient/family refusal <input type="checkbox"/> MER performed at outside hospital <input type="checkbox"/> Allergy to contrast material <input type="checkbox"/> Equipment-related delay * <input type="checkbox"/> No endovascular specialist available * <input type="checkbox"/> Delay in stroke diagnosis * <input type="checkbox"/> Vascular imaging not performed * <input type="checkbox"/> Advanced Age * <input type="checkbox"/> Other * * These reasons do not exclude from measure population	
^If MER treatment at this hospital, type of treatment:	<input type="checkbox"/> Retrievable stent <input type="checkbox"/> Other mechanical clot retrieval device beside stent retrieval <input type="checkbox"/> Clot suction device <input type="checkbox"/> Intracranial angioplasty, with or without permanent stent <input type="checkbox"/> Cervical carotid angioplasty, with or without permanent stent <input type="checkbox"/> Other	
^Is there documentation in the medical record of the first pass of a mechanical reperfusion device to remove a clot occluding a cerebral artery at this hospital?		<input type="radio"/> Yes <input type="radio"/> No
^What is the date and time of the first pass of a clot retrieval device at this hospital?	____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^^Is a cause(s) for delay in performing mechanical endovascular reperfusion therapy documented?		<input type="radio"/> Yes <input type="radio"/> No
^^Reasons for delay (select all that apply):	<input type="checkbox"/> Social/religious <input type="checkbox"/> Initial refusal <input type="checkbox"/> Care-team unable to determine eligibility <input type="checkbox"/> Management of concurrent emergent/acute conditions such as cardiopulmonary arrest, respiratory failure (requiring intubation) <input type="checkbox"/> Investigational or experimental protocol for thrombolysis <input type="checkbox"/> Delay in stroke diagnosis * <input type="checkbox"/> In-hospital time delay * <input type="checkbox"/> Equipment-related delay * <input type="checkbox"/> Need for additional imaging * <input type="checkbox"/> Catheter lab not available * <input type="checkbox"/> Other *	
^What is the location of the clot in the cerebral circulation?	<input type="radio"/> Proximal cerebral occlusion <input type="radio"/> Distal cerebral occlusion <input type="radio"/> Neither proximal or distal, OR unable to determine (UTD) from the medical record documentation	

^What cerebral artery is occluded?		<input type="radio"/> Anterior cerebral artery (ACA) <input type="radio"/> A1 ACA <input type="radio"/> Anterior communicating artery <input type="radio"/> Internal carotid artery (ICA) <input type="radio"/> ICA terminus (T-lesion; T occlusion) <input type="radio"/> Middle cerebral artery (MCA) <input type="radio"/> M1 MCA <input type="radio"/> M2 MCA <input type="radio"/> M3/M4 MCA <input type="radio"/> Vertebral artery (VA) <input type="radio"/> Basilar artery (BA) <input type="radio"/> Posterior cerebral artery (PCA) <input type="radio"/> Other cerebral artery branch/segment <input type="radio"/> The clinical location of the primary occluded vessel was not documented, OR unable to determine (UTD) from the medical record documentation.	
^Thrombolysis in Cerebral Infarction (TICI) Post-Treatment Reperfusion Grade		<input type="radio"/> Grade 0 <input type="radio"/> Grade 1 <input type="radio"/> Grade 2a <input type="radio"/> Grade 2b <input type="radio"/> Grade 3 <input type="radio"/> ND	
^Is there a documented TICI reperfusion grade post-treatment?	<input type="radio"/> 1 - A TICI reperfusion grade greater than or equal to (>=) 2B was documented posttreatment	<input type="radio"/> 2 - A TICI reperfusion grade less than (<) 2B was documented post-treatment	<input type="radio"/> 3 - A TICI reperfusion grade was not done post-treatment, OR Unable to determine (UTD) from the medical record documentation
^What was the date and time that a TICI 2B/3 was first documented during the mechanical thrombectomy procedure?		____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
COMPLICATIONS			
^Was there a positive finding on brain imaging of parenchymal hematoma, SAH, and/or IVH following IV or IA alteplase, or mechanical endovascular reperfusion therapy initiation?		<input type="radio"/> Yes <input type="radio"/> No	
^Date/Time of positive brain image :		____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^Results of positive brain image		<input type="checkbox"/> PH2 (Parenchymal Hematoma Type 2) <input type="checkbox"/> IVH (Intraventricular Hemorrhage) <input type="checkbox"/> SAH (Subarachnoid Hemorrhage) <input type="checkbox"/> RIH (Remote site of intraparenchymal hemorrhage outside the area of infarction) <input type="checkbox"/> Other positive finding not listed above <input type="checkbox"/> Not documented	
^What is the last NIHSS score documented prior to initiation of alteplase at this hospital?		_____	
This score obtained from:		<input type="radio"/> Baseline NIHSS <input type="radio"/> Subsequent NIHSS	
^What is the highest NIHSS score documented within 36 hours following initiation of IV alteplase?		_____	
^What is the last NIHSS score documented prior to initiation of IA alteplase or MER at this hospital?		_____	
This score obtained from:		<input type="radio"/> Baseline NIHSS <input type="radio"/> Subsequent NIHSS	
^What is the highest NIHSS score documented within 36 hours following IA alteplase or MER initiation?		_____	
^Is there documentation that a procoagulant reversal agent was initiated at this hospital?		<input type="radio"/> Yes <input type="radio"/> No	
^Date/Time procoagulant initiated		____/____/____ ____:____	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
^Is there documentation by a physician/APN/PA or pharmacist in the medical record of a reason for not administering a procoagulant reversal agent?		<input type="radio"/> Yes <input type="radio"/> No	

	Dosage 1. _____ 2. _____ 3. _____ 4. _____	Frequency 1. _____ 2. _____ 3. _____ 4. _____	Dosage 1. _____ 2. _____ 3. _____ 4. _____	Frequency 1. _____ 2. _____ 3. _____ 4. _____
	If NC, documented contraindications	<input type="checkbox"/> Allergy to or complications r/t antithrombotic <input type="checkbox"/> Patient/Family refused <input type="checkbox"/> Risk for bleeding or discontinued due to bleeding	<input type="checkbox"/> Serious side effect to medication <input type="checkbox"/> Terminal illness/Comfort Measures Only <input type="checkbox"/> Other	
Other Antithrombotic(s)	Prescribed? If yes,	<input type="radio"/> Yes <input type="radio"/> No/ND		
	Medication: <input type="checkbox"/> Desirudin (Iprivask) <input type="checkbox"/> Ticagrelor (Brilinta) <input type="checkbox"/> Prasugrel (Effient) *contraindicated in stroke and TIA <input type="checkbox"/> Other	Dosage 1. _____ 2. _____ 3. _____ 4. _____	Frequency 1. _____ 2. _____ 3. _____ 4. _____	
Persistent or Paroxysmal Atrial Fibrillation/Flutter		<input type="radio"/> Yes <input type="radio"/> No		
If atrial fib/flutter or history of PAF documented, was patient discharged on anticoagulation?		<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC		
If NC, documented reasons for no anticoagulation	<input type="checkbox"/> Allergy to or complication r/t warfarin or heparins <input type="checkbox"/> Mental status <input type="checkbox"/> Patient refused <input type="checkbox"/> Risk for bleeding or discontinued due to bleeding	<input type="checkbox"/> Risk for falls <input type="checkbox"/> Serious side effect to medication <input type="checkbox"/> Terminal illness/Comfort Measures Only		
Anti-hypertensive Tx (Select all that apply)	<input type="checkbox"/> None prescribed/ND <input type="checkbox"/> Other anti-hypertensive med <input type="checkbox"/> Ace Inhibitors <input type="checkbox"/> Beta Blockers	<input type="checkbox"/> None - Contraindicated <input type="checkbox"/> Diuretics <input type="checkbox"/> ARB <input type="checkbox"/> CA++ Channel Blockers		
Cholesterol-Reducing Tx (Select all that apply)	<input type="checkbox"/> None prescribed/ND <input type="checkbox"/> None – contraindicated <input type="checkbox"/> Statin <input type="checkbox"/> Fibrate	<input type="checkbox"/> Niacin <input type="checkbox"/> Absorption Inhibitor <input type="checkbox"/> PCSK 9 inhibitor <input type="checkbox"/> Other med		
Statin Medication:	<input type="checkbox"/> Amlodipine + Atorvastatin (Caduet) <input type="checkbox"/> Atorvastatin (Lipitor) <input type="checkbox"/> Ezetimibe + Simvastatin (Vytorin) <input type="checkbox"/> Fluvastatin (Lescol) <input type="checkbox"/> Fluvastatin XL (Lescol XL) <input type="checkbox"/> Lovastatin (Altoprev) <input type="checkbox"/> Lovastatin (Mevacor) <input type="checkbox"/> Lovastatin + Niacin (Advicor) <input type="checkbox"/> Pitavastatin (Livalo) <input type="checkbox"/> Pravastatin (Pravachol) <input type="checkbox"/> Rosuvastatin (Crestor) <input type="checkbox"/> Simvastatin (Zocor) <input type="checkbox"/> Simvastatin + Niacin (Simcor)	Statin Total Daily Dose:		
Documented Reason for Not Prescribing Guideline Recommended Dose?	<input type="checkbox"/> Intolerant to moderate (>75yr) or high (<=75yr) intensity statin <input type="checkbox"/> No evidence of atherosclerosis (cerebral, coronary, or peripheral vascular disease)	<input type="checkbox"/> Other documented reason <input type="checkbox"/> Unknown/ND		
Documented reason for not prescribing a statin medication at discharge?	<input type="radio"/> Yes <input type="radio"/> No			
New Diagnosis of Diabetes?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> ND			

Basis for Diagnosis (Select all that apply)	<input type="checkbox"/> HbA1c <input type="checkbox"/> Oral Glucose Tolerance	<input type="checkbox"/> Fasting Blood Sugar <input type="checkbox"/> Test Other
Diabetic Tx (select all that apply)	<input type="checkbox"/> None prescribed/ND <input type="checkbox"/> None – Contraindicated <input type="checkbox"/> Other subcutaneous/injectable agents	<input type="checkbox"/> Insulin <input type="checkbox"/> Oral agents
Anti-Smoking Tx	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> NC	
Smoking Cessation Therapies Prescribed (select all that apply)	<input type="checkbox"/> Counseling <input type="checkbox"/> Over the Counter Nicotine Replacement Therapy <input type="checkbox"/> Prescription Medications <input type="checkbox"/> Other <input type="checkbox"/> Treatment not specified	
Was the patient prescribed any antidepressant class of medication at discharge?	<input type="radio"/> Yes, SSRI	<input type="radio"/> Yes, any other antidepressant class <input type="radio"/> No/ND
OTHER LIFESTYLE INTERVENTIONS		
Reducing weight and/or increasing activity recommendations	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC	
TLC Diet or Equivalent	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC	
Antihypertensive Diet	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC	
Was Diabetic Teaching Provided?	<input type="radio"/> Yes <input type="radio"/> No/ND <input type="radio"/> NC	
STROKE EDUCATION		
Patient and/or caregiver received education and/or resource materials regarding all the following:		
Check all as Yes: <input type="checkbox"/>		
Risk Factors for Stroke	<input type="radio"/> Yes <input type="radio"/> No	Stroke Warning Signs and Symptoms <input type="radio"/> Yes <input type="radio"/> No
How to Activate EMS for Stroke	<input type="radio"/> Yes <input type="radio"/> No	Need for Follow-Up After Discharge <input type="radio"/> Yes <input type="radio"/> No
Their Prescribed medications	<input type="radio"/> Yes <input type="radio"/> No	
STROKE REHABILITATION		
Patient assessed for and/or received rehabilitation services during this hospitalization?		<input type="radio"/> Yes <input type="radio"/> No
Check all rehab services that patient received or was assessed for:	<input type="checkbox"/> Patient received rehabilitation services during hospitalization <input type="checkbox"/> Patient transferred to rehabilitation facility <input type="checkbox"/> Patient referred to rehabilitation services following discharge <input type="checkbox"/> Patient ineligible to receive rehabilitation services because symptoms resolved <input type="checkbox"/> Patient ineligible to receive rehabilitation services due to impairment (i.e. poor prognosis, patient unable to tolerate rehabilitation therapeutic regimen)	
STROKE DIAGNOSTIC TESTS AND INTERVENTIONS		
Cardiac ultrasound/echocardiography <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned	Extended implantable cardiac rhythm monitoring <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned	Carotid imaging <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned
Hypercoagulability testing <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned	Carotid revascularization <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned	Extended surface cardiac rhythm monitoring > 7 days <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned

Intracranial vascular imaging <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned	Short-term cardiac rhythm monitoring <= 7 days <input type="radio"/> Performed during this admission or in the 3 months prior <input type="radio"/> Planned post discharge <input type="radio"/> Not performed or planned
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OPTIONAL FIELDS – Please do not enter any patient identifiers in this section **Optional Fields Tab**

Field 1	Field 2	Field 3	Field 4	Field 5
Field 6	Field 7	Field 8	Field 9	Field 10
Field 11		Field 12		
Field 13	___/___/___ :___	<input type="checkbox"/> MM/DD/YYYY <input type="checkbox"/> Unknown	Field 14	___/___/___ :___
Additional Comments:				

Administrative

PMT used concurrently or retrospectively or combination?	<input type="radio"/> Concurrently	<input type="radio"/> Retrospectively	<input type="radio"/> Combination
Was a stroke admission order set used in this patient?	<input type="radio"/> Yes	<input type="radio"/> No	
Was a stroke discharge checklist used in this patient?	<input type="radio"/> Yes	<input type="radio"/> No	
Patient adherence contract/compact used?	<input type="radio"/> Yes	<input type="radio"/> No	

Outpatient **Outpatient Tab**

Patient	
Encounter Date: ___/___/___	E/M Code: _____
What is the date/time the patient departed from the emergency department?	___/___/___ :___ <input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
For discharges on or after 07/01/2012: What was the patient's discharge code from the outpatient setting?	<input type="checkbox"/>

Core Measure Tab

CORE MEASURE TAB (many elements are auto-populated within the online PMT)						
Check if patient is part of a sample		<input type="checkbox"/>				
First Name		Last Name				
Race	<input type="radio"/> Black or African American	<input type="radio"/> American Indian or Alaska Native	<input type="radio"/> Asian	<input type="radio"/> White	<input type="radio"/> Native Hawaiian or Pacific Islander	<input type="radio"/> UTD
Zip Code		Homeless	<input type="checkbox"/>			
What is the patient's source of payment for this episode of care?				<input type="radio"/> Medicare	<input type="radio"/> Non-Medicare	
HIC Number						
History & Last Known Well						
Was there physician/APN/PA documentation of a diagnosis, signed ECG tracing, or a history of ANY atrial fibrillation/flutter in the medical record?				<input type="radio"/> Yes	<input type="radio"/> No	
Is there documentation that the patient was on a lipid-lowering medication prior to hospital arrival?				<input type="radio"/> Yes	<input type="radio"/> No	
Is there documentation that the date and time of last known well was witnessed or reported?				<input type="radio"/> Yes	<input type="radio"/> No	
What was the date and time at which the patient was last known to be well or at his or her baseline state of		___/___/___ :___	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown			
When is the earliest physician/APN/PA documentation of comfort measures only?		<input type="radio"/> Day 0 or 1	<input type="radio"/> Day 2 or after	<input type="radio"/> Timing unclear	<input type="radio"/> Not Documented/UTD	
Thrombolytics						

Is there documentation that IV alteplase therapy initiated at this hospital?	<input type="radio"/> Yes	<input type="radio"/> No
Is there documentation on the day of or day after hospital arrival of a reason for extending the initiation of IV thrombolytic to 3 to 4.5 hours of Time Last Known Well?	<input type="radio"/> Yes	<input type="radio"/> No
Did the patient receive IV or IA alteplase at this hospital or within 24 hours prior to arrival?	<input type="radio"/> Yes	<input type="radio"/> No
Is there documentation on the day of or day after hospital arrival of a reason for not initiating IV thrombolytic?	<input type="radio"/> Yes	<input type="radio"/> No
Early Antithrombotics		
Was antithrombotic therapy administered by the end of hospital day 2?	<input type="radio"/> Yes	<input type="radio"/> No
Labs		
Was the LDL-cholesterol (LDL-c) measured within the first 48 hours or 30 days prior to hospital arrival?	<input type="radio"/> Yes	<input type="radio"/> No
Was the patient's highest LDL-cholesterol (LDL-c) level greater than or equal to 100 mg/dL in the first 48 hours or within 30 days prior to hospital arrival?	<input type="radio"/> Yes	<input type="radio"/> No
Discharge Information		
Discharge Date/Time	___/___/___ __:___	<input type="checkbox"/> MM/DD/YYYY only <input type="checkbox"/> Unknown
Was antithrombotic therapy prescribed at hospital discharge?	<input type="radio"/> Yes	<input type="radio"/> No
Is there documentation by a physician/advanced practice nurse/physician assistant (physician/APN/PA) or pharmacist in the medical record of a reason for not prescribing antithrombotic therapy at hospital discharge?	<input type="radio"/> Yes	<input type="radio"/> No
Was anticoagulation therapy prescribed at hospital discharge?	<input type="radio"/> Yes	<input type="radio"/> No
Is there documentation by a physician/advanced practice nurse/physician assistant (physician/APN/PA) or pharmacist in the medical record of a reason for not prescribing anticoagulation therapy at hospital discharge?	<input type="radio"/> Yes	<input type="radio"/> No
Was a statin medication prescribed at discharge?	<input type="radio"/> Yes	<input type="radio"/> No
Stroke Core Measure Additional Comments:		
CSTK Additional Comments:		
END OF FORM		