Society of Cardiovascular Patient Care 2013 Updates: Guidelines and Troponin Turn-Around-Time Documentation Requirements



SOCIETY OF CARDIOVASCULAR PATIENT CARE



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There are no disclosures

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Overview of Acute Coronary Syndromes (ACS)

Discuss the new myocardial infarction (MI) and S-T Elevation MI (STEMI) guidelines

Discuss the Society of Cardiovascular Patient Care (SCPC) Troponin Turn-around-Time (TTAT) documentation requirements for accreditation

SCPC Site Survey Findings

What we know

Coronary Heart Disease is the #1 disease in the United States



Get the Facts



16.3 million people over age 20 in the U.S. have some form of **coronary heart disease**

Cardiovascular disease is the leading hospital discharge diagnostic group (DRG 390 - 459)

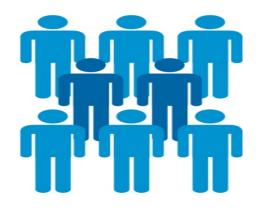


Get the Facts

5-8 million patients present to the Emergency Department (ED) annually for chest pain



to



Mission & Values:

SCPC was developed to share best practices that improve outcomes of patients with suspected or acute cardiovascular disease through innovative cross-disciplinary processes. *In short, to bring science to the bedside...*



Collaboration

SCPC shares with its facilities the goal of early diagnosis of myocardial infarction (MI) and improvement in patient outcomes through education, accreditation and process improvement.

Through the *process of accreditation* we help break down barriers and facilitate communication to achieve successful continuum of care.



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Accreditation and Certification



We create communities of excellence that bring science to the bedside.

Accreditation Partnerships



We create communities of excellence that bring science to the bedside.



Early Heart Attack Care





Heart attacks have beginnings

EHAC (Early Heart Attack Care) shifts the focus from *treating to preventing a heart attack*.

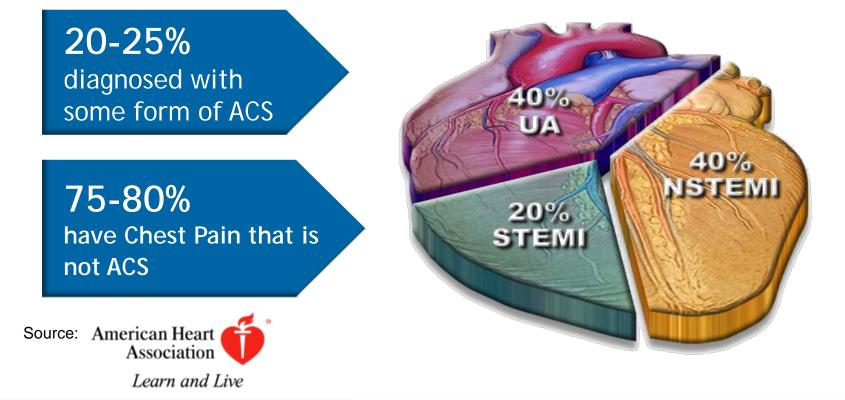
Adults ignore or deny symptoms or complex co-morbidities lead to confusion

- Mild chest pain
- Fatigue
- Shortness of breath
- Stuttering chest discomfort

Which places them in grave danger of heart muscle damage or death

Acute Coronary Syndrome (ACS)

ACS comprises three conditions: ST-elevation Myocardial Infarction (MI or STEMI); Non-ST-elevation MI (NSTEMI) and Unstable Angina (UA)



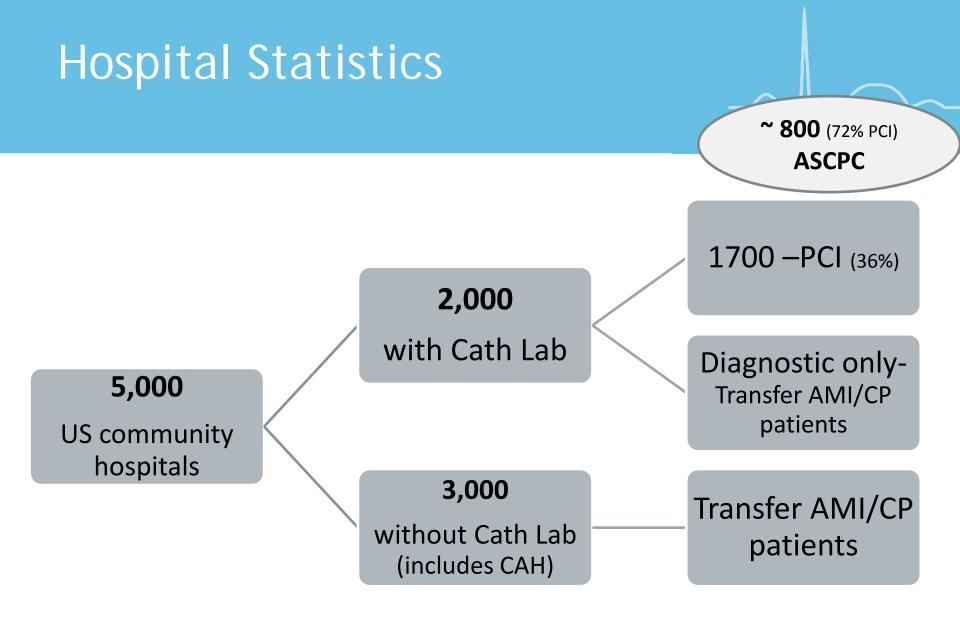
Definitions for Treating MI -Reperfusion

•<u>Percutaneous Coronary Intervention (PCI)</u> – most frequently used invasive method of treating the narrowing, or stenosis, of coronary arteries; performed in cardiac catheterization facilities (cath lab) at acute care hospitals

•<u>Primary PCI (PPCI)</u>- also known as (aka) emergency angioplasty, is a lifesaving intervention performed during a heart attack (acute ST-segment elevation myocardial infarction aka: STEMI)

•<u>Non-primary PCI</u> - aka: elective angioplasty, scheduled intervention to relieve the narrowing of the artery; goal of preventing a heart attack from occurring in the future

Key Point: All laboratorians should be very familiar with the protocols and diagnostic capabilities (cath lab, PPCI, lytics, transfer) to address acute cardiac events.



Source: American Hospital Association & ACC/NCDR/Cath-PCI

Estimated In-hospital Mortality by Door-to-Reperfusion Times

TIME (minutes)

CM Gibson

15

30

60

90

120

180

240



	2.9 (2.8-3.1)
	3.0 (2.9-3.2)
	3.5 (3.4-3.6)
	4.3 (4.24.4)
7	5.6 (5.4-5.7)
	8.4 (8.2-8.7)
	10.3 (10.0-10.7)

There is no *floor* to the *mortality reduction* that can be achieved by *reducing time to treatment* **and** Each 30 min. of delay translates into a 7.5% increase in relative risk of 1-yr mortality.

*Adjusted for age, sex, race, findings on presentation, medical history, procedural characteristics, angiographic findings, and hospital factors

Any delay in D2B time associated with increased in-hospital mortality Rathore SS, et al. *BMJ* 2009; 339:b1807. Yale University School of Medicine; ACC-NCDR

New Updated MI Guidelines

Third Universal Definition of Myocardial Infarction

Kristian Thygesen, Joseph S. Alpert, Allan S. Jaffe, Maarten L. Simoons, Bernard R. Chaitman and Harvey D. White

Circulation. published online August 24, 2012

•Key Points:

•Optimal precision, as described by coefficient of variation (CV) at the 99th percentile URL for each assay, should be defined as <10%

•Assays with CV >20% at the 99th percentile URL should not be used

•Blood samples for the measurement of cTn should be drawn on first assessment and repeated 3- 6 h later

Consensus Document

Journal of the American College of Cardiology © 2012 by the American College of Cardiology Foundation Published by Elsevier Inc.

Vol. 60, No. 23, 2012 ISSN 0735-1097/\$36.00 http://dx.doi.org/10.1016/j.jacc.2012.08.969

EXPERT CONSENSUS DOCUMENT

ACCF 2012 Expert Consensus Document on Practical Clinical Considerations in the Interpretation of Troponin Elevations

A Report of the American College of Cardiology Foundation Task Force on Clinical Expert Consensus Documents

Developed in Collaboration With the American Association for Clinical Chemistry, American College of Chest Physicians, American College of Emergency Physicians, American Heart Association, and Society for Cardiovascular Angiography and Interventions



New Updated MI Guidelines – Follow-up Article

How to Use High-Sensitivity Cardiac Troponins in Acute Cardiac Care Kristian Thygesen et al European Heart Journal doi:10.1093/eurheart/ehs154 PDF online 2012

Summary Regarding Use of hsCardiac Troponin in Clinical Routine:

- Use 99th%ile concentration of the reference population as the cTn URL
- The diagnosis of acute myocardial necrosis requires a significant change with serial testing...a minimum change of >20% in follow-up testing is required
- Additional testing of other early markers of acute myocardial necrosis, such as myoglobin or creatine kinase MB is no longer needed
- Blood sampling in patients with suspicion of AMI should be performed on admission and 3 h later...repeated 6 h after admission in patients of whom the 3 h values are unchanged but...clinical suspicion of AMI is still high

New Updated STEMI Guidelines

2013 ACCF/AHA Guideline for the Management of ST-Elevation Myocardial Infarction

Patrick T. O'Gara et al

Circulation. published online December 17, 2012

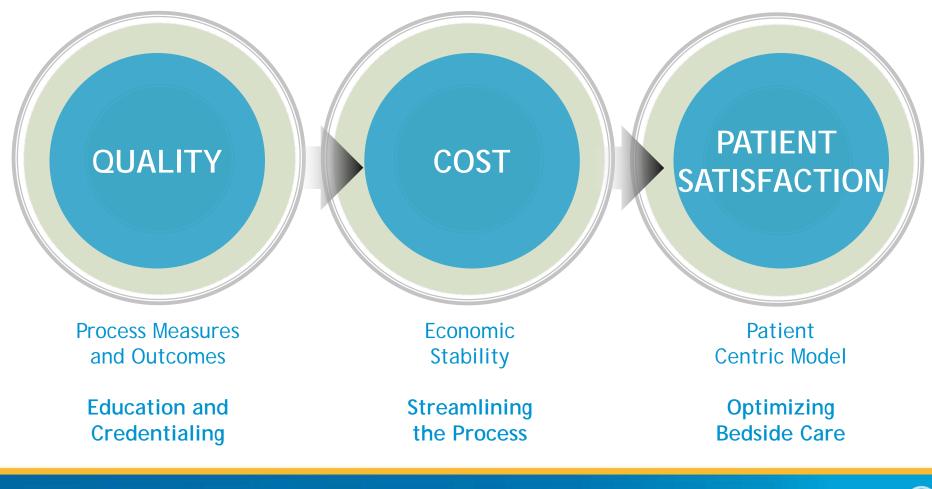
•Key Points:

•Major and comprehensive revision of the prior 2004 Guideline

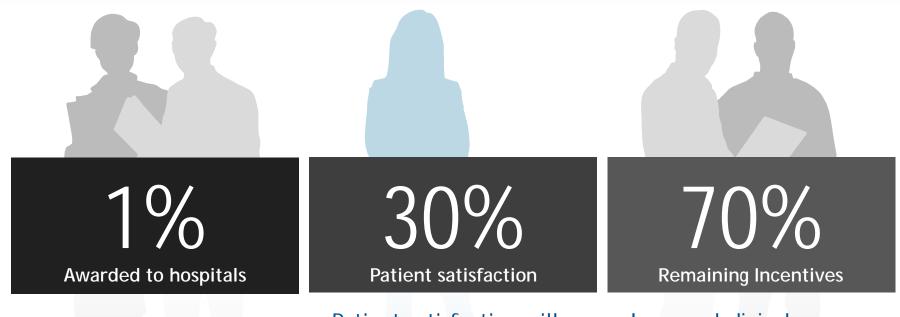
•Concept and terminology changes: "Door to Balloon (Needle)" replaced with "first medical contact (FMC) to device" time

System goals of EMS-FMC-to-device = 90 minutes or less
For transfers goals of EMS-FMC-to-device = 120 minutes or less
and D1D2R = 90 minutes
For transfers goals is "Door in-Door out" = 30 minutes or less
Fibrinolytic therapy goal = 30 minutes

Healthcare Today



Value Based Purchasing



1% of Medicare payments to hospitals will be withheld during FY 2013 and awarded to hospitals that meet a set of quality performance measures. Patient satisfaction will determine 30% of the incentive payments while..

Improved clinical outcomes will determine the remaining **70% of the incentive payments**.

Value Based Purchasing

Two educational articles on VBP: Free CME/CE

CMS Value-based Purchasing Targets Complications, Readmissions by Jean Moody-Williams, Medscape: Article #763832 (thru 05/29/13)

Value Based Purchasing: Excellent care Boosts the Bottom Line by Charles F. Bombard, Nurse.com- CE663 (thru 08/10/2015)

Observation Units and the Lab



What is Observation Services? Medicare Definition

A well-defined set of specific , clinically appropriate services

- commonly ordered for patients who present to the emergency department (ED)
- require a significant period of treatment or monitoring

Ongoing short term treatment, assessment, and reassessment decision for further treatment to...

- inpatient
- discharge

Medicare policy manual rev. 137 12-30-10

Observation Services

In 2003 national survey:

Emergency Department Observation Units (EDOUs):

- -19% of US hospitals
- -12% planning a unit
- A 2007 subsequent survey: -
 - EDOU increased to 36%
 - > 1/2 managed by ED MD's

Ross et al. Critical Pathways, 2012 The State of the ART: Emergency Room Observation Units.

ACS in Observation = Laboratory Impact

BENCHMARKS:

- Average length of stay (LOS) in a <u>dedicated</u> OBS ~ 15 hours
- ~ 70-80% are discharged / inpatient admit rate ~20%
- Less than 1% of patients staying longer than 48 hours
- ... observation protocols have been shown to decrease unnecessary resource utilization and cost to 50% to 70% of routine inpatient care costs
- Accelerated Diagnostic Protocols (ADP) for serial cardiac biomarkers can help achieve benchmarks

Readmissions = Laboratory Impact





Healthcare Stats: Readmission

- Hospitals readmit nearly 1 in 5 Medicare patients within one month of discharge (cost = \$17 billion /yr)
- National average for readmissions ~19%
- CMS effort to curb readmissions for three conditions:
 - heart attack, heart failure, pneumonia
 - HF: #1 cause for admission over age 65 and readmissions
- Penalty/fines assessments:
 - 1% October 2012
 - 2% October 2013
 - 3% in 2014

Laboratory Role and Readmissions

April 2013 Clinical Laboratory News: Volume 39, Number 4

The Race to Reduce Readmissions: Can Lab Tests Help Predict

Who Will Return to the Hospital?

- Key Points:
 - Simple test combinations used as "risk predictors"
 - Laboratory tests can prevent early discharges leading to increased readmissions
 - Lab based readmission calculators:
 - CORE Readmission Risk Calculator Yale Medical School
 - Intermountain Risk Score Intermountain Health

Laboratory Role Overall: Clinical Support and Expertise



Accountability

Changing Perspectives of Turn-Around-Time Tracking: Healthcare Implications

Changing Perspectives of TAT Tracking: Healthcare Implications

Recent studies and research support the movement towards the following:

- Assessing the "whole process" (ie: arrival)
- Standardizing definitions of turn-around-time (TAT)
- Assessing TAT with patient outcomes and length of stay

Changing Perspectives of TAT Tracking: Healthcare Implications

Study by Ervasti et al, Clin Chem Lab Med 2008

Proposed new concepts for TAT in the diagnostic process:

As a "Patient-oriented" view or the "whole process"

- Diagnostic TAT arrival to reporting of results (outcomes median 122 min)
- Clinical TAT arrival to order
- Laboratory TAT order to report/resulted

Changing Perspectives of TAT Tracking: Healthcare Implications

In Academic Emergency Medicine, 2010:17, Hwang et al noted:

- "Guidelines do not exist delineating times frames for when a troponin test should optimally be resulted in association with improved patient outcomes."
- " Prolonged laboratory TAT may delay recognition of conditions in the acutely ill, potentially affecting clinician decision-making and the initiation of timely treatment."
 - Outcomes median 107 minutes; "ordered to resulted"

SCPC

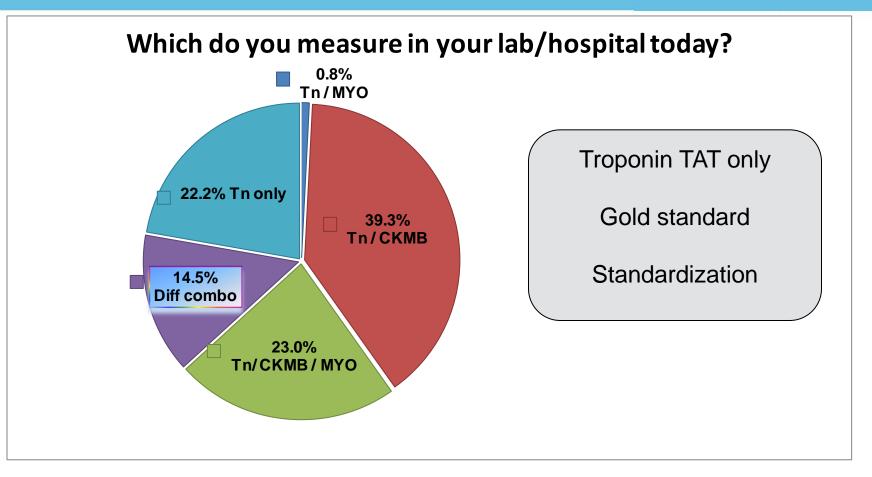
Cardiac Biomarkers and Troponin Turn-Around-Time Requirements

SCPC Cardiac Biomarker Requirements

Measuring TAT is a guideline driven recommendation

- No previous TAT requirement
 - SCPC requirement starting in 2012
 - Track and demonstrate improvements

Troponin Measurement



CLN – April 2009, vol 35, no 4



Yes	No	No Item					
				ponin "turnaround time" (TAT) is broken down into the at of ORDER to COLLECT.	4.4.1.		
		Baseline troponin TAT is broken down into the time segment of COLLECT to RECEIVE IN LAB. Baseline troponin TAT is broken down into the time segment of RECEIVE IN LAB to RESULT.					
		Baseline troponin TAT is broken down into the time segment of DOOR to RESULT.					
		Baseline troponin TAT is broken down into the time segment of DOOR to ORDER.Baseline troponin TAT is broken down into the time segment of ORDER to RESULT.					
		Baseline troponin TAT is broken down into the time segment of COLLECT to RESULT. 90% of baseline troponin TAT of ORDER TO RESULT or COLLECT to RESULT is within 60 minutes.					
		Yes	No	Item			
				90% of baseline troponin TAT of ORDER to RESULTS or COLLECT to RESULT is within 30 minutes.	4.4.9.		
Item							
evide	nced-b	ased seri	al tr	narker approach includes documentation of an oponin strategy that is consistent with the assay used ollowing green Tier III Items:	4.5.0.		
Yes	No	Item					
		The facility has a process in place to monitor the TAT of serial draws for troponin.					
	The cardiac biomarker protocol includes a serial troponin from arrival up to 6 hours. The protocol may last less than 6 hours if provocative cardiac testing or imaging takes place.						

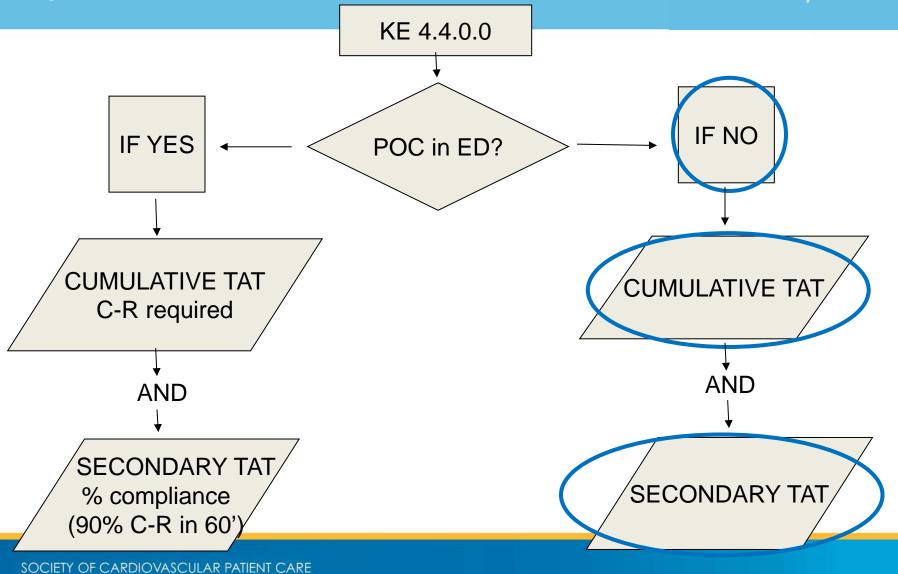
SCPC Accreditation Tool Example

SCPC ACCREDITATION & BIOMARKER TESTING: Key Element 4 - 4.4.0.0 overview

> Demonstrating a process for reviewing and assessing BASELINE Troponin TAT ED patients

Documentation requirements:

- Monthly or quarterly meeting notes ARE YOU INVOLVED?
 - Lab participates as an agenda item MUST BE ON CPC TEAM
 - Metrics, process and action plans discussed
- Minimum 6 months of data
- Goal times
- Required to provide TAT metrics: cumulative & secondary
 - Point-of-Care Testing (POCT) / Central Laboratory Analyzers



40

Where does Point-of-Care Testing (POCT) fit in with CP Accreditation?

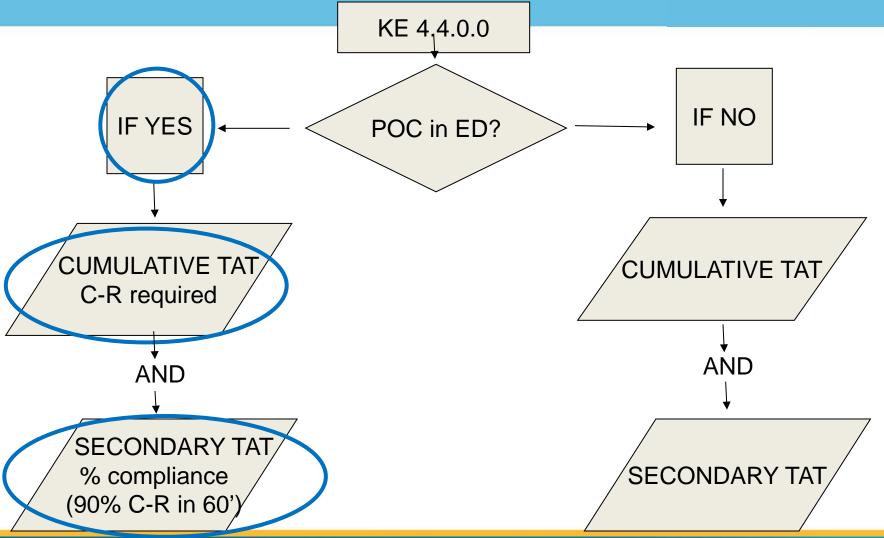
"To the extent that laboratory test TAT is <u>only one</u> <u>factor</u> impacting

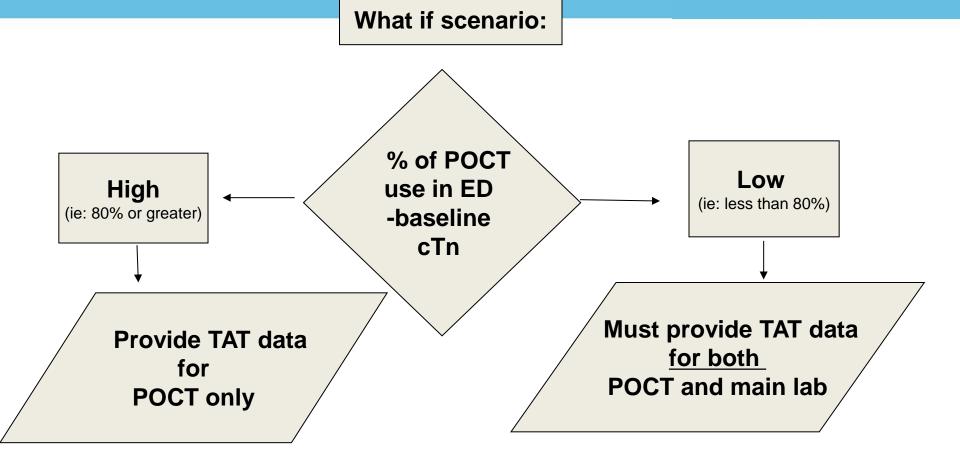
ED length of stay and patient outcomes,

it is unlikely that POCT alone, in the absence of an interdepartmental approach to ED operations,

will produce measurable improvements in outcomes."

Lewandrowski, E. et al. Cardiac Marker Testing As Part Of An Emergency Department Point-of-Care Satellite Laboratory In A Large Academic Medical Center. Practical Issues Concerning Implementation. Point of Care. The Journal of Near Patient testing & Technology. Vol. 1, No.3, pp. 145-154.





Example: Troponin TAT

CBM TAT EXAMPLE COLLECTION TOOL: 90% Goal w/in 60 / inutes								
Year:	Total (n=ED Tnl)	# Collect to Result w/in 60 min	Collect- Result <=60 min (=C/B)	4.4.7.0 Goal				
January	522	479	92%	90%				
February	554	453	82%	90%				
March	590	522	88%	90%				
April	520	477	92%	90%				
Мау	517	468	91%	90%				
June	507	471	93%	90%				
July	544	514	94%	90%				
August	473	440	93%	90%				
September	491	452	92%	90%				
October	534	484	91%	90%				
November	494	435	88%	h%				
December	<u>490</u>	<u>463</u>	<u>94%</u>	4.4.8.0				
Totals:	6236	5658	<u>91%</u>	<u>0%</u>				

DATA SUBMISSION OPTIONS

College of American Pathologist (CAP) QM1 monitor

The Society has partnered with the College of American Pathologist who have created a validation tool which collects data to meet the Society requirements for TAT tracking.

Additional benefits are:

- One source collecting data for research to track the "diagnostic TAT" or "door to result' data through sampling
- Great for facilities with large volumes
- Benchmarking
- Estimates trending of process improvement initiatives

4.5.1.0 – The facility has a process in place to monitor the TAT of serial draws for Troponin

Key concept: "Windows of Scheduled Time" • *Assessment and documentation of serial draw time points*

SCPC ACCREDITATION & BIOMARKER TESTING: Facility Information Booklet

For both CENTRAL LAB and POC Cardiac Markers

Test

Which ones being used?

- Manufacturer
- Analyzer
- Hours drawn from arrival time (i.e. 0-3-6 hours, 0=Initial)
- Cut-point used for negative biomarker results
- Decision point used for positive tests
- Intermediate or "gray-zone" range (if applicable) for Troponin only

- Cannot be > 8 hours from 0 time
- Biggest area of discrepancy between manufacturers recommendations and the decision points being used



SCPC GENERAL FINDING

(results for discussion purposes only)

- Cycle III FIB data estimates: n=700
 - approximately half using POCT for CBM
 - ▶ 65% using the 99th percentile

Polling results from webinars

(results for discussion purposes only)

Does your facility have a cath lab that can perform PCI?	Yes No	45% 27%
	Not sure	28%
Does your facility transfer chest pain or AMI patients? (n ~ 300)	Yes No Not sure	33% 53% 12%
Are you using the 99th %ile? (n~220)	Yes No Not sure	60% 9% 31%
Are you using a diagnostic protocol of 0-3-6-? (n ~ 220)	Yes No Not sure	56% 25% 19%
Do you provide education to your physicians?	Yes	51%

(n ~ 200)

Accreditation Summary

Expectations and documentation requirements include:

- TAT tracking for POC and/or central lab analyzers
- Laboratory participation -CPC meetings-quarterly
- Metrics by time points; defined starting/end points; goals
- Communicate with the key representatives for the CPC
 - CPC Coordinator may be dual role with Heart Failure/Stroke
 - CPC Medical Director
 - Director of the Emergency Department
 - Director of Cardiology
 - Administration Director of Quality

Overall goal, improving patient throughput and care.

Best demonstrated practices have requirements for defining and tracking cumulative turn-around-time metrics of the whole-process or the patientcentric view.

As such, timeliness of the reporting of Troponin equals timeliness of the diagnosis for the appropriate delivery of care in the Acute Coronary Syndrome patient population.

Validation Tool Resource

College of American Pathologists www.cap.org QM1 Monitor

SCPC Resources:

www.scpcp.org info@scpcp.org *Subject line:*

SCPC Laboratory Subject-Matter-Expert : Ruth Cantu



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