ALBERTA PRECISION LABORATORIES

Leaders in Laboratory Medicine

DATE:	2022 July 11
TO:	Edmonton Zone – Physicians, Nurses, Laboratory Directors, and Managers
FROM:	Clinical Biochemistry, North Sector, Alberta Precision Laboratories (APL)
	Implementation of high sensitivity troponin T (hs-TnT) testing at Misericordia Community Hospital and Grey Nuns Community Hospital

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Key Message

 The Roche high sensitivity troponin T (hs-TnT) assay will replace the Beckman hs-TnI assay at Misericordia Community Hospital on Tuesday July 19, 2022 at 10:00 AM, and Grey Nuns Community Hospital on Thursday July 21, 2022 at 10:00 AM (Table 1).

Site	Current assay	New assay	Go-Live date
*MCH	Beckman hs-Tnl	Roche hs-TnT	July 19, 2022
*GNH	Beckman hs-Tnl	Roche hs-TnT	July 21, 2022
RAH	Beckman hs-Tnl	Roche hs-TnT	End of September, date TBD

*This bulletin applies to the Misericrodia and Grey Nuns Community Hospital. MCH = Misericordia Community Hospital; GNH = Grey Nuns Community Hospital; RAH = Royal Alexandra Hospital; CCI = Cross Cancer Institute

TBD = to be determined

- Numerical values, 99th percentile upper reference limits (reference intervals), reporting limits, delta values, critical values, and interpretative comments will change (Table 2 and 3; click here for more details: <u>hsTnT Survival Guide; hsTnT Edmonton Zone Chest Pain Pathway</u>).
- Rule-in/rule-out cut points and a new chest pain algorithm involving 0 & 2 hour and 2 & 4 hour serial time points are also changing to provincially align reporting (Figure 1).
- For troponin collected in the outpatient/community setting only, critical troponin levels will be phoned to the ordering provider when the result is ≥ 53 ng/L (Table 2).

Why this is important

- Implementation of hs-TnT is part of a large-scale provincial project to implement the latest generation of Roche instruments at major urban hospitals. University of Alberta Hospital and Sturgeon Community Hospital have already implemented hs-TnT on June 19 and June 21, respectively. The Royal Alexandra Hospital will continue to use Beckman hs-TnI until the designated go-live date. See Table 1 for hs-TnT implementation schedule.
- Different assays are not comparable because they produce different values and use different cutoffs, therefore, Hs-TnI and hs-TnT and will not trend in Epic. See Appendix 1 for a complete list of all assays used across Edmonton Zone.
- The use of hs-TnT in conjunction with a 2-hour rapid protocol has been successfully used in other hospitals across Canada and has already been used in Calgary for over 8 years.



Background

- Evidence supports the Roche hs-TnT assay in rapid diagnostic decisions for patients presenting with chest pain and shows excellent clinical sensitivity (98.7 – 99.9%) at low levels to rule out acute myocardial infarction (AMI). Studies further support serial troponin measurements to calculate delta values at 0 & 2 hour and 2 & 4 hour intervals to rule in or exclude AMI (Figure 1).
- Additionally, elevated hs-TnT may not always represent AMI and could be due to other causes of myocardial injury. Clinical judgment is essential to ensure safe patient management.
- These changes are consistent with clinical practice guidelines and is supported by a joint Alberta Health Services (AHS)/APL working group representing Cardiology, Emergency Medicine, and Laboratory Medicine within Edmonton Zone*.

Action Required

- Be familiar with the transition from the Beckman hs-TnI assay to the Roche hs-TnT assay at Misericordia and Grey Nuns Community Hospitals.
- Be familiar with laboratory reporting changes and the new rapid chest pain pathway.
- Be aware of different troponin assays used within Edmonton Zone (Appendix 1). Do not interpret results interchangeably across sites with different assays.

Inquiries and feedback may be directed to

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This bulletin has been reviewed and approved by

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- Emergency Department: Drs. Brian Rowe, Shandra Doran, Dennis Lefebvre
- Cardiology: Drs. Sean van Diepen, Michelle Graham, and Robert Welsh



Table 2: Summary of new reporting scheme for the Roche hs-TnT assay

	Reporting details	Notes
Units	ng/L (whole numbers)	N/A
Reference interval	< 14 ng/L	Values above this limit will be flagged as high.
Critical Value	Only outpatient / community trop ≥ 53 ng/L Only outpatient / community trop critical value will be phoned ou ordering provider	
Reporting limits	3 ng/L to endpoint	N/A
Delta value	Reported for 0 - 2 hour delta and 2 - 4 hour delta	Reported if a previous hs-TnT value on the same patient and method is within 4 hours
Result comments	Pathway interpretive comments & method identification comments	N/A



Table 3: Interpretive comments reported with hs-TnT for in-hospital and community patients

A) <u>In-hospital</u> patients:

hs-TnT result (ng/L)	Comment	Flagging
< 5	For patients with a non-ischemic ECG, a Troponin T, High Sensitivity of 4 ng/L or less on presentation is highly sensitive for excluding acute myocardial infarction, provided the specimen was collected more than 3-hours from symptom onset. However, for patients with symptoms less than 3-hours duration or concerning clinical presentations, repeat troponin testing at 2-hours after the initial sample is recommended. Please note that patients with ischemic ECG changes and/or high-risk clinical presentations should be considered for further evaluation irrespective of troponin results.	Normal
5 -13	 Troponin T, High Sensitivity is below the upper reference limit (14 ng/L) and results are not consistent with myocardial infarction or injury. However, patients with acute symptoms (less than 6-hours) or concerning clinical presentations should undergo repeat troponin testing at 2-hours after the initial sample. A 2-hour change of 3 ng/L or less is highly sensitive for excluding acute myocardial infarction. A 2-hour change of 4-9 ng/L may indicate acute myocardial injury. Repeat clinical evaluation, ECG and troponin at 4-hours after the initial sample is recommended. A 2-hour change of 10ng/L suggests an acute myocardial injury and may represent acute myocardial infarction in the appropriate clinical scenario. Please note that patients with ischemic ECG changes and/or high-risk clinical presentations should be considered for further evaluation irrespective of troponin results. 	Normal
14 - 52	 Troponin T, High Sensitivity has a non-specific/non-diagnostic elevation. Interpretation is highly dependent on clinical presentation and patient history. New elevations are concerning; however, many patients have chronic elevations in troponin and measured concentrations near the patient's baseline are reassuring. Patients with acute symptoms (less than 6 hours) or concerning clinical presentations should undergo repeat troponin testing at 2-hours after the initial sample. A 2-hour change of 3 ng/L or less suggests acute myocardial infarction is unlikely. A 2-hour change of 4-9 ng/L may indicate acute myocardial injury. Repeat clinical evaluation, ECG and troponin at 4-hours after the initial sample is recommended. A 2-hour change of 10 ng/L suggests an acute myocardial injury and may represent acute myocardial infarction in the appropriate clinical scenario. Please note that patients with ischemic ECG changes and/or high-risk clinical presentations should be considered for further evaluation irrespective of troponin results. 	High
≥ 53	Clear elevation of Troponin T, High Sensitivity consistent with acute myocardial injury or infarction in the appropriate clinical context. Repeat troponin testing at 2-hours after the initial sample may be helpful to assess for ongoing myocardial injury.	Critical



B) Community patients

hs-TnT result (ng/L)	Comment	Flagging
< 14	Troponin T, High Sensitivity is below the upper reference limit (14 ng/L) and results are not consistent with myocardial infarction or injury, provided the specimen was collected more than 3 hours from the onset of symptoms. Patients with active symptoms, ischemic ECG changes and/or concerning clinical presentations should be considered for urgent evaluation irrespective of troponin results.	Normal
14 - 52	Troponin T, High Sensitivity has a non-specific/non-diagnostic elevation. Interpretation is highly dependent on clinical presentation and patient history. New elevations are concerning; however, many patients have chronic elevations in troponin and measured concentrations near the patient's baseline are reassuring. Patients with active symptoms. Ischemic ECG changes and/or concerning clinical presentations should be considered for urgent evaluation irrespective of troponin results.	High
≥ 53	Clear elevation of Troponin T, High Sensitivity consistent with myocardial injury or infarction. Interpretation is highly dependent on clinical presentation and patient history. Many patients have chronic elevations in troponin and measured concentrations near the patient's baseline are reassuring. New troponin elevations are concerning and urgent assessment in an emergency department may be indicated in the appropriate clinical context.	Critical

Figure 1: The new reporting scheme supports the following chest pain pathway which outlines the risk management of patients presenting with suspected AMI to acute care setting (see hsTnT Survival **Guide** for more details)

Part I



≤ 44 years ≥ 3 risk factors or

history of atherosclerotic

1 or 2 risk factors

No risk factors

disease

2

1

0 known

Diabetes

□ Obesity

Risk

Factors

Current smoker

HTN (diagnosed)
HL (diagnosed)
Family hx CAD

measured concentrations similar to the patient's baseline are reassuring. *** The indeterminate risk pathway arm relies on expert opinion, the

experience in Calgary, and our Edmonton Zone hs-Tnl chest pain pathway (initiated 2019). It aligns with current guideline recommendations.



Part II: Intermediate Risk



Note:

* Consider using a structured risk assessment tool such as the HEART score to aid decision making for all patients without indications for Cardiology consultation.

** For all patients with abnormal hs-TnT results, check the medical record for prior results. Many patients have stable abnormalities in hs-TnT and measured concentrations similar to the patient's baseline are reassuring.

***The indeterminate risk pathway arm relies on expert opinion, the experience in Calgary, and our Edmonton Zone hs-TnI chest pain pathway (started 2019). It aligns with current guideline recommendations.

otal (10 maximum)

≥ 3 risk factors or history of

disease 1 or 2 risk factors

atherosclerotic

No risk factors

known

1 0

2

1

0

2

0

HEART Score Interpretation

> 3x normal limit (43ng/L or greater)

1-3x normal limit (14-42ng/L) < normal limit (<14ng/L)

1

History

ECG

Age

Risk

Factors

hs-cTnT

(Peak)

Diabetes

Current smoker

HTN (diagnosed)

□ Family hx CAD □ Obesity

Low Risk	0-3
Moderate Risk	4-6
High Risk	7-10



Appendix 1: Summary of different troponin assays in Edmonton Zone after September 2022.

Different assays are not comparable because they produce different values and use different cutoffs.

Site	Troponin assay	Chest Pain protocol
University of Alberta Hospital		
Royal Alexandra Hospital*		
Grey Nuns Hospital	Roche hs-TnT	2-hour
Misericordia Hospital		
Sturgeon Hospital		
Strathcona Community Hospital	Beckman hs-Tnl	2 hour
Leduc Community Hospital		
Westview Health Center		3-hour
Northeast Health Centre		
Fort Saskatchewan Community Hospital		
East Edmonton Urgent Care Centre	Siemens Stratus	6-hour
Devon General Hospital	conventional TnI	
Redwater Health Center	7	
DynaLIFE Medical Labs (community)	Siemens Atellica hs-Tnl	Not Applicable

*Royal Alexandra Hospital will be implementing hs-TnT at end of September 2022. The Beckman hs-TnI assay and 3-hour ches