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DATE:	2022 September 6	
TO:	All Central Zone Physicians and Healthcare Providers	
FROM:	Clinical Biochemistry, South Sector, Alberta Precision Laboratories (APL)	
RE:	Availability of High Sensitivity Troponin I (hsTnI) at Wetaskiwin Hospital and Care Centre	

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Key Message

- Effective September 13, 2022, high sensitivity troponin I (hs-TnI) on bioMérieux MINI VIDAS® will replace the current conventional (non-high sensitivity) Abbott iSTAT® TnI assay at Wetaskiwin Hospital and Care Centre.
- The use of hs-TnI will result in changes in reporting interpretation (new reference interval, critical value and units), and updated Urgent Care and Rural Hospital Vidas hs-cTnI Chest Pain Pathway (<u>https://insite.albertahealthservices.ca/Main/assets/tms/edc/tms-edc-vidas-hs-ctnI-pathway.pdf</u>) and reporting comments (Table 1)
 - New units: ng/L
 - New reference interval: 0 18 ng/L
 - New critical value: >= 100 ng/L
 - New Meditech test code for hs-Tnl ordering in Wetaskiwin: TROPIHST
 - New result test name in NetCare: Troponin I (Vidas) High Sensitivity

Why this is important

- Availability of hs-TnI in Wetaskiwin will better support clinicians in evidence-based interpretation of their troponin result and improve patient management.
- Recognize that hs-Tnl results are NOT interchangeable with conventional Tnl, or with hsTnT.

Background

- While hs-Tnl offers improved diagnostic performance relative to conventional troponin assays, hs-Tnl results
 alone cannot exclude all acute coronary syndrome presentations and high-risk clinical presentations remain
 high-risk, even if hs-Tnl concentrations are normal.
- Abnormal elevations in hs-Tnl do not necessarily represent acute myocardial injury or coronary ischemia; clinical judgment remains essential to ensure safe patient management.

Action Required

- All users of hs-TnI in Wetaskiwin should familiarize themselves with the Urgent Care and Rural Hospital Vidas hscTnI Chest Pain Pathway and accompanying comments.
- Be aware of differences between hs-TnI and hs-TnT for reference intervals and critical values, as result interpretation will change by test.

Effective

• September 13, 2022

Inquiries and feedback may be directed to

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This bulletin has been reviewed and approved by

- Heidi Paulin, MD, Regional Laboratory Medicine Site Chief for Red Deer Regional Hospital Centre
- Hossein Sadrzadeh, PhD, Section Chief, Clinical Biochemistry, South Sector
- Paul Klonowski, MD, Associate Medical Director, South Sector

Table 1: Updated bioMérieux MINI VIDAS® hsTnl Reporting

Current Reporting	New Reporting
(Abbott iSTAT® Tnl, non-high sensitivity assay)	(MINI VIDAS® hs-Tnl assay)
N/A	<6 ng/L:
	For patients with a non-ischemic ECG, a Troponin I, High Sensitivity of less than 6 on presentation AND at 2-hours is highly sensitive for excluding acute myocardial infarction (MI). Repeat troponin testing at 2-hours after the initial sample is recommended for all patients to reliably exclude MI. Please note that patients with ischemic ECG changes and /or high- risk clinical presentations should be considered for further evaluation irrespective of troponin results.
Tnl 0.02-0.04 μg/L:	hs-Tnl 6-18 ng/L:
Troponin I value not consistent with acute myocardial infarction, providing the sample was collected more than 6h from onset of symptoms. Repeat troponin testing after the initial sample is recommended for all patients to reliably exclude MI. Please note that patients with ischemic ECG changes and/or high-risk clinical presentations should be considered for further evaluation irrespective of troponin results.	Troponin I, High Sensitivity is below the upper reference limit (19 ng/L) and results are not consistent with myocardial infarction (MI) or injury. However, patients with acute symptoms (less than 6-hours since onset) or concerning clinical presentations should undergo repeat troponin testing at 2-hours after the initial sample. Repeat troponin testing at 2-hours after the initial sample is recommended for all patients to reliably exclude MI. A 2-hour change of 10 ng/L or more suggests an acute myocardial injury and may represent acute MI 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.
Tnl 0.05 –0.10 μg/L:	hs-Tnl 19-99 ng/L:
Troponin I value is inconclusive for acute myocardial infarction and may be due to myocardial injury. Repeat troponin testing after the initial sample is recommended for all patients to reliably exclude myocardial infarction. Please note that patients with ischemic ECG changes and/or high-risk clinical presentations should be considered for further evaluation irrespective of troponin results.	Troponin I, 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. Repeat troponin testing at 2-hours after the initial sample is recommended to assess for active myocardial injury. A 2-hour change of 10 ng/L or more suggests an acute myocardial injury and may represent acute MI 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.
Tnl >0.10 μg/L:	hs-Tnl >= 100 ng/L
Clear elevation of Troponin I consistent with acute myocardial injury or infarction in the appropriate clinical context.	Clear elevation of Troponin I, 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



Repeat troponin testing after the initial sample may be helpful to assess for ongoing myocardial iniury	helpful to assess for ongoing myocardial injury.
Tnl >0.10 µg/L may be observed in several nonthrombotic cardiac and systemic diseases (most commonly - acute PE, acute pericarditis, acute or severe HF, myocarditis, sepsis and/or shock).	