

SITE-SPECIFIC PROCEDURE CHANGES TO CONSIDER WHEN TRANSITIONING TO ABBOTT'S HIGH-SENSITIVITY POINT-OF-CARE TROPONIN ASSAY

LANGUAGE

- Update all procedure documents to be consistent with *i-STAT hs-TnI* cartridge.
- If any reference literature is indicated in the procedure, then update to align with hs-Tnl information (e.g. IFU, external guidelines)

CHECKS

- Revise all ranges within procedure and reporting systems (e.g. reportable, reference intervals)
- Add parameters according to facility procedure and clinical decisions
 - Sex-specific range
 - Overall range
 - Identify elevated, actionable, or critical ranges (if applicable)

NOTE: Update ranges in DE customization, if applicable

SPECIMEN COLLECTION

- Sample type and volume
- Collection device
- Sample stability

NOTE: Laboratory director is responsible for the approval of changes with signature and date of procedure document changes.

REAGENT / CONTROLS STORAGE CONDITIONS

- Expiration dates
- Room temperature
- · Refrigeration and freezer storage

NOTE: Refer to *i-STAT hs-Tnl* cartridge Instructions for Use (IFU) for comprehensive details on specimen preparation and reagent storage conditions

REPORTING

- Add parameters according to facility procedure and clinical decisions
 - Order and results code
 - Order and result name
 - Units of measure of new ranges
- Additional reportable items
 - Delta check
 - Result interpretation
 - LOINC and CPT (if applicable)

Visit the sites below for more information:

http://loinc.org/downloads/loinc-table/#users-guide https://www.codemap.com/abbott/userhome.cfm