

December 18, 2025

HEMATOLOGY

Lupus Anticoagulant**Date effective: December 22, 2025**

Background Information: Provincially standardized Lupus Anticoagulant testing, currently performed at the Health Sciences Centre Hemostasis Laboratory, utilizes two assays in the testing algorithm (Dilute Russell Viper Venom [DRVV] and Time and lupus-sensitive APTT [PTT-LA] with and without added phospholipids). The existing PTT-LA assay will transition to a Hexagonal Phase Phospholipid PTT-LA kit assay (Precision Biologic, HEXLA kit). This test utilizes the Werfen (previously known as Instrumentation Laboratory or IL) ACL TOP series coagulation analyzers.

Clinical Practice Change:

- PTT-LA results utilizing the HEXLA kit are reported as the Delta value (change in value) between the PTT-LA Start (Lupus sensitive APTT) and PTT-LA Correct (Lupus sensitive APTT with added hexagonal phospholipid).
 - 'PTT-LA Delta' value is reported in seconds(s).
 - The reference range is ≤ 10.9 (s).
 - PTT-LA Delta is reported on the patient report when it exceeds the reference range.
 - PTT-LA Test Ratios are not performed and will no longer appear on patient reports.
- Lupus Anticoagulant interpretation will continue to be reported as 'Demonstrated' or 'Not Demonstrated' on all requests.

References/Resources:

Please refer to the Laboratory Information Manual (LIM) [Lab Information Manual](#)

System Improvements:

- Streamlined laboratory workflow for Lupus Anticoagulant testing and consumables.
- Improved turnaround time for result reporting.
- Improved utilization of laboratory resources.

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