

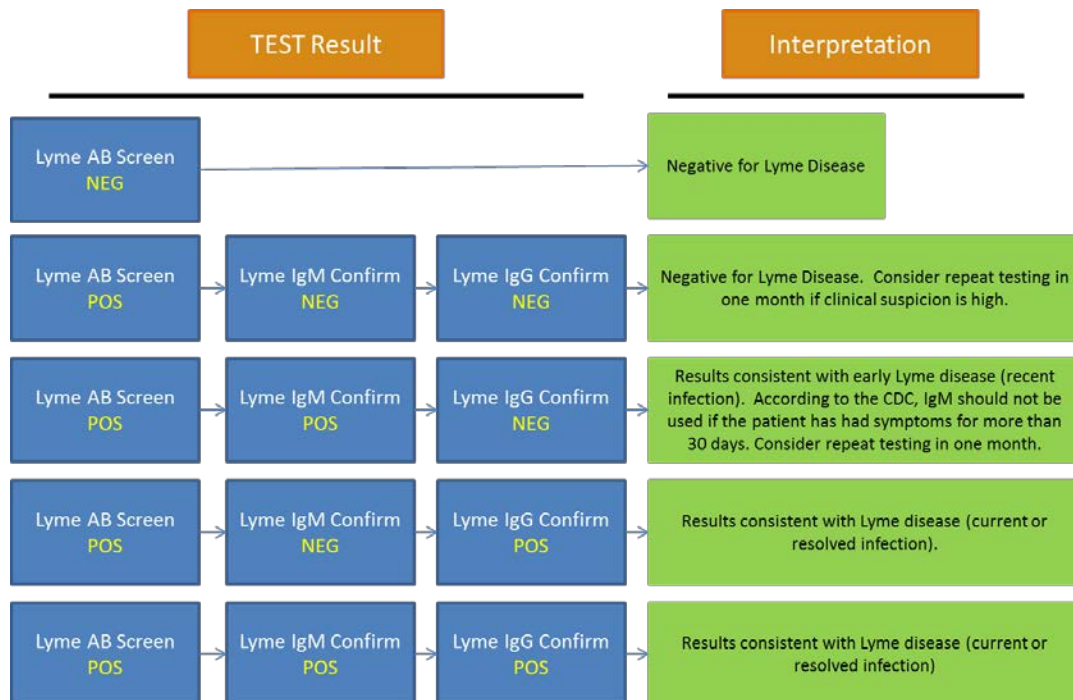
Lyme Disease Antibody Testing: Change to Assay

Date: March 20, 2019

Effective Date: April 17, 2019

The URM Clinical Microbiology Laboratory is instituting a **two tier testing algorithm for Lyme Disease**, replacing the single Lyme C6 antibody currently in use. At this time, the LYME C6 Peptide Antibody (IgG/IgM) will be used and resulted as a **Lyme antibody screening assay**. Specimens which test positive for antibodies to Lyme C6 peptide will then be reflexed to **confirmatory Lyme IgM and IgG ELFA assays**. This constitutes a modified two-tier assay (i.e. a screening EIA followed by a confirmatory EIA/ELFA).

The use of an EIA/ELFA as a second tier improves sensitivity compared to a traditional two tier testing algorithm which used a western blot as a confirmatory second step. The new testing algorithm will be reported as shown below.





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Testing Information:

Code: LYMAB

SPECIMEN: Serum Separator Tube with 1mL

SCHEDULE: Both tiers will be run Mon-Sat during the day. Expected turn-around-time is 24-72 hours.

Please contact Serology Supervisor, at 275-7801 with any questions.

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