

TEST CHANGE

Borrelia burgdorferi Antibody, IgM by Immunoblot 0050253, LYME M WB

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S	necimen.	Red	uirements:

Patient Preparation:

Collect: Serum separator tube.

Separate serum from cells ASAP or within 2 hours of collection. **Specimen Preparation:**

Transfer 1 mL serum to an ARUP Standard Transport Tube.

Effective Date: May 15, 2023

(Min: 0.15 mL)

Refrigerated. **Transport Temperature:**

Unacceptable Conditions: CSF or plasma. Contaminated, heat-inactivated, severely

hemolyzed, severely lipemic, and severely icteric specimens.

Remarks:

Stability: After separation from cells: Ambient: 48 hours; Refrigerated: 2

weeks; Frozen: 1 year (avoid repeated freeze/thaw cycles)

Qualitative Immunoblot Methodology:

Performed: Sun-Sat

Reported: 1-42 days

Note: Current CDC recommendations for the serologic diagnosis of

Lyme disease are to screen with a polyvalent EIA test and confirm equivocal and positive with immunoblot. Both IgM and IgG immunoblots should be performed on specimens less than 4 weeks after appearance of erythema migrans. Only IgG immunoblot should be performed on specimens greater than 4 weeks after disease onset. IgM immunoblot in the chronic stage is not recommended and does not aid in the diagnosis of neuroborreliosis or chronic Lyme disease. Please submit

requests for appropriate immunoblot testing within 10 days.

CPT Codes: 86617

New York DOH Approval Status: This test is New York DOH approved.

Interpretive Data:

For this assay, a positive result is reported when any 2 or more of the following bands are present: 23, 39, or 41 kDa. All other banding patterns are reported as negative.

Reference Interval:

Effective Date: May 15, 2023



Effective August 15, 2011

Negative