

## **TEST CHANGE**

Specimen Requirements:         Patient Preparation:         Collect:       Serum separator tube.         Specimen Preparation:       Separate serum from cells ASAP or within 2 hours of collection. Transfer 0.3 mL serum to an ARUP standard transport tube_Standard Transport Tube. (Min: 0.1 mL)         Transport Temperature:       Refrigerated.         Unacceptable Conditions:       Room temperature specimens. Plasma, CSF, or other body fluids. Contaminated, heat-inactivated, hemolyzed, icteric, or severely lipemic specimens.         Remarks:       Stability:         After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 year.         Methodology:       Semi-Quantitative Enzyme-Linked Immunosorbent Assay         Performed:       Tue, Thu, Sat Mon, Wed, Fri         Reported:       1-Z4 days         Note:       2         CPT Codes:       83516 x2         New York DOH Approval Status:       This test is New York DOH approved.	Ganglioside (GM1) Antibodies, IgG and IgM 0050591, GM1 PAN			
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Performed: Tue, Thu, Sat Mon, Wed, Fri   Reported: 1-74 days   Note: Sat	Stability:			
Mon, Wed, Fri       Reported:     1-74 days       Note:        CPT Codes:     83516 x2	Methodology:	Semi-Quantitative Enzyme-Linked Immunosorbent Assay		
Note: CPT Codes: 83516 x2	Performed:			
CPT Codes: 83516 x2	Reported:	1- <u>7</u> 4 days		
	Note:			
New York DOH Approval Status: This test is New York DOH approved.	CPT Codes:	83516 x2		
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Interpretive Data:

Ganglioside antibodies are associated with diverse peripheral neuropathies. Elevated antibody levels to ganglioside-monosialic acid (GM1) are associated with motor or sensorimotor neuropathies, particularly multifocal motor neuropathy. Anti-GM1 may occur as IgM (polyclonal or monoclonal) or IgG antibodies. These antibodies may also be found in patients with diverse connective tissue diseases as well as normal individuals. These tests by themselves are not diagnostic and should be used in conjunction with other clinical parameters to confirm disease.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.



Component	Interpretation
GM1 Antibody IgG	29 IV or less: Negative 30-50 IV: Equivocal 51- 100 IV: Positive 101 IV or greater: Strong Positive
GM1 Antibody, IgM	29 IV or less: Negative 30-50 IV: Equivocal 51- 100 IV: Positive 101 IV or greater: Strong Positive

## Reference Interval:

Test Number	Components	Reference Interval
	GM1 Antibody, IgG	50 IV or less
	GM1 Antibody, IgG	
		Component(s) Interpretation
		GM1 Antibody IgG 29 IV or less 30- 50 IV 51-100 IV 101 IV or greater Positive Strong Positive
	GM1 Antibody, IgM	50 IV or less