

TEST CHANGE

Acetylcholine Receptor Binding Antibody with reflex to Muscle-Specific Kinase (MuSK) Ab, IgG

Effective Date: May 15, 2023

3001868, ACHR BIN R

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Specimen Requirements:		
Patient Preparation:		
Collect:	Serum Separator Tube (SST).	
Specimen Preparation:	Separate from cells ASAP or within 2 hours of collection. Transfer 0.5 mL serum to an ARUP Standard Transport Tube. (Min: 0.3 mL)	
Transport Temperature:	Refrigerated.	
Unacceptable Conditions:	Plasma. Contaminated, hemolyzed, or severely lipemic specimens.	
Remarks:		
Stability:	After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month (avoid repeated freeze/thaw cycles)	
Methodology:	Quantitative Radioimmunoassay (RIA)	
Performed:	Sun-Sat	
Reported:	2-8 days	
Note:	If Acetylcholine Receptor Binding Antibody result is less than or equal to 0.4 nmol/L then Muscle-Specific Kinase (MuSK) Ab, IgG (ARUP test code 300 <u>6</u> 1 <u>98</u> 576) will be added. Additional charges apply.	
CPT Codes:	83519; if reflexed, add <u>86255; if reflexed, add 86256</u> 83519	

Interpretive Data:

New York DOH Approval Status:

Approximately 85-90 percent of patients with myasthenia gravis (MG) express antibodies to the acetylcholine receptor (AChR), which can be divided into binding, blocking, and modulating antibodies. Binding antibody can activate complement and lead to loss of AChR. Blocking antibody may impair binding of acetylcholine to the receptor, leading to poor muscle contraction. Modulating antibody causes receptor endocytosis resulting in loss of AChR expression, which correlates most closely with clinical severity of disease. Approximately 10-15 percent of individuals with confirmed myasthenia gravis have no measurable binding, blocking, or modulating antibodies.

This test is New York DOH approved.



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.

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Component Interpretive Data
Acetylcholine
Binding Antibody
Negative 0.5
nmol/L or greater:
Positive

Reference Interval:

Test Number	•	Reference Interval
	Acetylcholine Binding Antibody	0.4 nmol/L or less

HOTLINE NOTE: There is a reflexive pattern change associated with this test. One or more orderable or component has been added or removed to the reflexive pattern. Refer to the Hotline Test Mix for interface build information.