

Effective Date: February 20, 2024

TEST CHANGE

Autoimmune Vision Loss Panel, Serum 3016804. AIVLS

3016804, AIVLS	
Specimen Requirements:	
Patient Preparation:	
Collect:	Serum separator tube.
Specimen Preparation:	Separate serum from cells ASAP or within 2 hours of collection. Transfer 1 mL serum to an ARUP standard transport tube. (Min: 0.30 mL)
Transport Temperature:	Refrigerated.
Unacceptable Conditions:	Plasma. Contaminated, heat-inactivated, hemolyzed, or lipemic specimens.
Remarks:	
Stability:	After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month
Methodology:	Qualitative Immunoblot/Semi-Quantitative Cell-Based Indirect Fluorescent Antibody
Performed:	Varies
Reported:	1-8 days
Note:	If CV2.1 Antibody IgG Screen by IFA is positive, then CV2.1 Antibody IgG Titer by IFA will be added. Additional charges apply.
CPT Codes:	84182, 86255; if reflexed, add 86256
New York DOH Approval Status:	Specimens from New York clients will be sent out to a New York DOH approved laboratory, if possible.
Interpretive Data:	
Refer to report	
Reference Interval:	
Test Components Number	Reference Interval

Less than 1:10<u>0</u>

Negative

CV2-1 Ab IgG CBA-IFA Screen, Serum

Recoverin Ab, IgG by Immunoblot, Serum



Effective Date: February 20, 2024

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.