

**TEST CHANGE**

Leucine-Rich, Glioma-Inactivated Protein 1 Antibody, IgG CBA-IFA with Reflex to Titer, Serum

2009456, LGI1 IGG

**Specimen Requirements:**

**Patient Preparation:**

**Collect:** Serum separator tube.

**Specimen Preparation:** Separate serum from cells within 2 hours of collection. Transfer 1 mL serum to an ARUP [standard transport tube](#). [Standard Transport Tube](#). (Min: 0.2 mL)

**Transport Temperature:** Refrigerated.

**Unacceptable Conditions:** CSF or plasma. Contaminated, hemolyzed, or severely lipemic specimens.

**Remarks:**

**Stability:** After separation from cells: Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 year (avoid repeated freeze/thaw cycles)

**Methodology:** Semi-Quantitative Cell-Based Indirect Fluorescent Antibody

**Performed:** Wed

**Reported:** 1-8 days

**Note:** If LGI1 antibody IgG is positive, then LGI1 antibody IgG titer will be added. Additional charges apply.

**CPT Codes:** 86255; if reflexed, add 86256

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

**Interpretive Data:**

Leucine-rich, glioma-inactivated 1 protein (LGI1) IgG antibody may occur as part of the voltage-gated potassium channel (VGKC) complex antibodies.

The presence of LGI1 IgG antibody is mainly associated with limbic encephalitis, hyponatremia, and myoclonic movements. LGI1 IgG antibody is rarely associated with tumors but may occur infrequently in Morvan syndrome, neuromyotonia, and idiopathic epilepsy. The full-spectrum of clinical disorders associated with the LGI1 IgG antibody continues to be defined. Results should be interpreted in correlation with the patient's clinical history and other laboratory findings.

This indirect fluorescent antibody assay utilizes ~~leucine-rich, glioma-inactivated 1 protein (LGI1)~~

transfected cell lines for the detection and ~~semi-quantification~~ **semiquantification** of the LGI1 IgG antibody.

~~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.~~

Reference Interval:

Less than 1:10