

**NEW TEST**

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CV2 Antibody, IgG by CBA-IFA With Reflex to Titer, CSF  
3017001, CV2 CSF

**Specimen Requirements:**

**Patient Preparation:**

**Collect:** CSF.

**Specimen Preparation:** Transfer 0.5 mL CSF to an ARUP standard transport tube. (Min: 0.15 mL)

**Transport Temperature:** Refrigerated.

**Unacceptable Conditions:** Hemolyzed, contaminated, or severely lipemic specimens.

**Remarks:**

**Stability:** Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month

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

**Performed:** Thu

**Reported:** 1-8 days

**Note:** If CV2 Antibody IgG Screen by IFA, CSF is positive, then CV2 Antibody IgG Titer, CSF will be added. Additional charges apply.

**CPT Codes:** 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:**

CV2 antibodies aid in discriminating between chronic paraneoplastic neurological disorder (PND) and other inflammatory disorders of the nervous system. Anti-CV2 is associated with small-cell lung cancer and thymoma. A negative test result does not rule out a diagnosis of autoimmune neurologic disease. Results should be interpreted in correlation with the patient's clinical history and other laboratory findings.

This indirect fluorescent antibody assay utilizes CV2 transfected cell lines for the detection and semiquantification of the CV2 IgG antibody.

**Reference Interval:**

Test Number	Components	Reference Interval

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	CV2 Ab IgG CBA-IFA Screen, CSF	Less than 1:1
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**HOTLINE NOTE: Refer to the Hotline Test Mix for interface build information.**