

partment of Pathology Effective Date: October 7, 2024

TEST CHANGE

Varicella-Zoster Virus Antibody, IgG, CSF 0054444, VZECSF

Specimen Requirements:

Patient Preparation:

Collect: CSF.

Specimen Preparation: Transfer 0.5 mL CSF to an ARUP <u>standard transport</u>

tube. Standard Transport Tube. (Min: 0.3 mL)

Transport Temperature: Refrigerated. Also acceptable: Frozen.

Unacceptable Conditions: Contaminated, heat-inactivated, hemolyzed, or xanthochromic

specimens.

Remarks:

Stability: Ambient: 8 hours; Refrigerated: 2 weeks; Frozen: 1 year

Methodology: Semi-Quantitative Chemiluminescent Immunoassay

Performed: Sun-Sat

Reported: Within 24 hours

Note:

CPT Codes: 86787

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:

The detection of antibodies to varicella-zoster in CSF may indicate central nervous system infection. However, consideration must be given to possible contamination by blood or transfer of serum antibodies across the blood-brain barrier.

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:

Effective October 7, 2024 February 18, 2020

0134.9 S/COIV or less

Negative - No significant level of IgG antibody to varicella-zoster virus detected.



135.0-164.9 IV

Equivocal—Repeat testing in 10-14 days may be helpful.

1165.0 S/COIV or greater

Positive - IgG antibody to varicella-zoster virus detected, which may indicate a current or past varicella-zoster infection.

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HOTLINE NOTE: There is a numeric map change associated with this test. Refer to the Hotline Test Mix for interface build information.

HOTLINE NOTE: There is a unit of measure change associated with this test. Refer to the Hotline Test Mix for interface build information.