

**TEST CHANGE** 

Chlamydia Antibody Panel, IgG & IgM by IFA 0065100, V CHLM PAN

Specimen Requirements:

Patient Preparation:

Collect: Plain red or 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. Standard Transport Tube. (Min: 0.415 mL) Parallel testing is preferred and convalescent specimens must be received within 30 days from receipt of acute specimens. Mark

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specimens plainly as "acute" or "convalescent."

Transport Temperature: Refrigerated.

Unacceptable Conditions: Contaminated, hemolyzed, or hyperlipemic sera.

Remarks:

Stability: After separation from cells: Ambient: 48 hours; Refrigerated: 2

weeks; Frozen: 1 year (avoid repeated freeze/thaw cycles)

Methodology: Semi-Quantitative Indirect Fluorescent Antibody (IFA)

Performed: Sun-Fri

Reported: 1-4 days

Note:

CPT Codes: 86631 x3; 86632 x3

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

Interpretive Data:

Refer to individual components. Refer to individual components.

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:

- < 1:64 *C. pneumoniae* IgG.
- < 1:64 C. psittaci IqG.
- < 1:64 C. trachomatis IgG.
- < 1:20 C. pneumoniae IgM.



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< 1:20 *C. psittaci* lgM. < 1:20 *C. trachomatis* lgM.