

Chlamydia IgM, Immunofluorescence, Serum

Test ID: CHLM

Useful for:

Assessing IgM antibody levels to aid in the clinical diagnosis of *Chlamydia pneumoniae* or *Chlamydia psittaci* infections

Methods:

Micro-Immunofluorescent Antibody (MIF) Assay

Reference Values:

Chlamydia pneumoniae <1:10

Chlamydia psittaci <1:10

Specimen Requirements:

Collection container:

Preferred: Serum gel	
Acceptable: Red top	
Submission Container:	Plastic vial
Specimen Volume:	0.3 mL
Minimum Volume:	0.15 mL

Specimen Stability Information:

Specimen Type	Temperature	Time
Serum	Refrigerated (preferred)	30 days
	Frozen	30 days

Cautions:

• Specimens collected too early during primary infection may not contain detectable antibodies. If chlamydial infection is suspected, a second specimen should be collected 10 to 21 days later and tested in parallel with the original specimen.

- During a primary *Chlamydia* infection, the early antibody response may be cross-reactive with multiple *Chlamydia* species.
- This assay does not report antibodies detected against *Chlamydia trachomatis*. Sera from suspected cases
 of lymphogranuloma venereum (LGV) should be tested by a Lymphogranuloma Venereum Differentiation
 Antibody Panel. LGV testing is not performed by Mayo Clinic Laboratories; call 800-533-1710 for
 assistance with ordering.
- Due to the limited sensitivity and specificity of *Chlamydia* serologic tests, patients with suspected *C trachomatis* infection should be tested by a molecular method (eg, CTRNA / *Chlamydia trachomatis*, Nucleic Acid Amplification, Varies) when clinical manifestations are present.

CPT Code:

86632 x 2

Day(s) Performed: Monday through Friday

Report Available: 1 to 4 days

Questions

Contact Dunisha Messmer, Laboratory Technologist Resource Coordinator at 800-533-1710.