

| Patient Information | Specimen Information | Client Information |
|---------------------|----------------------|--------------------|
|                     |                      |                    |

**COMMENTS:**

| Test Name                | In Range              | Out Of Range | Reference Range | Lab |
|--------------------------|-----------------------|--------------|-----------------|-----|
| CHLAMYDOPHILA PNEUMONIAE |                       |              |                 |     |
| AB (IGG, IGA, IGM)       |                       |              |                 |     |
| C. PNEUMONIAE AB (IGG)   | <1:64                 |              | titer           |     |
| C. PNEUMONIAE AB (IGA)   | <1:16                 |              | titer           |     |
| C. PNEUMONIAE AB (IGM)   | <1:10                 |              | titer           |     |
| INTERPRETATION           | ANTIBODY NOT DETECTED |              |                 |     |

REFERENCE RANGE:

IgG <1:64  
 IgA <1:16  
 IgM <1:10

The immunofluorescent detection of specific antibodies to Chlamydomphila pneumoniae may be complicated by crossreactive antibodies, non-specific antibody stimulation, or past exposure to similar organisms such as C. psittaci and Chlamydia trachomatis. IgM titers of 1:10 or greater usually indicate recent infection, and any IgG titer may indicate past exposure. IgA is typically present at low titers during primary infection, but may be elevated in recurrent exposures or in chronic infection.

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

**PERFORMING SITE:**

**SPECIMEN:**