

TEST CHANGE

Coxiella burnetii (Q-Fever) Antibodies, IgG and IgM, Phase I and II with Reflex to Titer

2012634, Q-F GM		
Specimen Requirements:		
Patient Preparation:		
Collect:	Serum separator tube (SST).	
Specimen Preparation:	Separate serum from cells ASAP or within 2 hours of collection Transfer 1 mL serum to an ARUP <u>standard transport</u> <u>tube.Standard Transport Tube.</u> (Min: 0. <u>4</u> 3 mL) Parallel testing is preferred and convalescent specimens must be received within 30 days from receipt of the acute specimens. Mark specimens plainly as "acute" and "convalescent."	
Transport Temperature:	Refrigerated.	
Unacceptable Conditions:	Contaminated, hemolyzed, or severely lipemic specimens.	
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:	Mon, Wed, Fri	
Reported:	1-6 days	
Note:	For IgG or IgM testing, if any Phase I or Phase II screening result is Indeterminate or Positive, then titer(s) will be added. Additional charges apply.	
CPT Codes:	86638 x4; if reflexed add 86638 per titer	
New York DOH Approval Status:	This test is New York DOH approved.	
Interpretive Data:		
Refer to report.		
Reference Interval:		

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Test Number	Components	Reference Interval
	C. burnetii (Q- Fever) Ab, Phase I IgG	Negative
	C. burnetii (Q- Fever) Ab, Phase II IgG	Negative
	C. burnetii (Q- Fever) Ab, Phase I IgM	Negative
	C. burnetii (Q- Fever) Ab, Phase II IgM	Negative