MAYO CLINIC
Mayo Medical Laboratories

Laboratory Service Report 1-800-533-1710

Patient Name	Patient ID	Age	Gender	Order #
SAMPLEREPORT,FBPTS AB	SA00053202	38	F	SA00053202
Ordering Phys CLIENT,CLIENT				**DOB** 05/16/1974
Client Order # SA00053202	**Account Information**			**Report Notes**
Collected 01/30/2013	C7028846-DLMP ROCHESTER 3050 SUPERIOR DRIVE ROCHESTER,MN 55901			
Printed 01/31/2013 14:39				

Test	Flag	Results	Unit	Reference Value	Perform Site*
B. pertussis Ab, IgA Immunoblot			REPORTED 01/31/2013 08:50		
B. pertussis Ab, IgA Immuno Interp		Negative		Negative	Y006
No IgA antibodies against Bordetella.					
B. pertussis, IgA Immunoblot PT		Negative			Y006
B. pertussis, IgA Immunoblot FHA		Negative			Y006
B. pertussis Ab, IgG Immunoblot			REPORTED 01/31/2013 08:50		
B. pertussis Ab, IgG Immuno Interp		Negative		Negative	Y006
No IgG antibodies against Bordetella FHA and PT detected.					
B. pertussis, IgG Immunoblot PT100		Negative			Y006
B. pertussis, IgG Immunoblot PT		Negative			Y006
B. pertussis, IgG Immunoblot FHA		Negative			Y006
B. pertussis Ab, IgM Immunoblot			REPORTED 01/31/2013 08:50		
B. pertussis Ab, IgM Immuno Interp		Negative		Negative	Y006
No IgM antibodies against Bordetella FHA and PT detected.					
This test was developed and its performance characteristics determined by ARUP Laboratories. The U.S. Food and Drug Administration has not approved or cleared this test; however, FDA clearance or approval is not currently required for clinical use. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions.					
B. pertussis, IgM Immunoblot PT		Negative			Y006
B. pertussis, IgM Immunoblot FHA		Negative			Y006
B. pertussis Abs IgG, M, A w/Reflex			REPORTED 01/31/2013 08:49		
B. pertussis Ab, IgG	H	6.5	U/mL	<=2.4	Y006
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Performing Site Legend on Last Page of Report

* Report times for Mayo performed tests are CST/CDT

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B. pertussis Ab, IgM	H	4.7	U/mL	<=1.1	Y006
BACKGROUND INFORMATION: B. pertussis Ab, IgM w/Reflex Recommend that treatment decisions be based on the result of the B. pertussis IgM immunoblot test instead of the ELISA test. B. pertussis IgM test by ELISA may produce false-positive results. This test was developed and its performance characteristics determined by ARUP Laboratories. The U.S. Food and Drug Administration has not approved or cleared this test; however, FDA clearance or approval is not currently required for clinical use. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions.					
B. pertussis Ab, IgA	H	3.9	U/mL	<=1.1	Y006
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* Performing Site:

Y006	ARUP Laboratories 500 Chipeta Way Salt Lake City, UT 84108	Lab Director: