

Patient Information	Specimen Information	Client Information	
DOB: AGE: Gender: Phone: Patient ID: Health ID:	Specimen: Requisition: Lab Ref #: Collected: Received: Reported:	REQUEST A TEST - PW 7027 MILL RD STE 201 BRECKSVILLE, OH 44	
Test Name	In Range Out Of Ra		Lab
MEASLES AB (IGG), IMMUNE STATUS AU/mL Int	LA (MMR) AB (IGG) PANEL, IMMUNE 138.00 cerpretation	AU/mL	AT
13.50-16.49 Equ	consistent with immunity privocal insistent with immunity		
	sles IgG suggests immunization o ection with measles virus.	r	
http://education.Que	rmation, please refer to estDiagnostics.com/faq/FAQ162 provided for informational/ s only.)		
	101.00 cerpretation	AU/mL	AT
<9.00 No 9.00-10.99 Ec	ot consistent with immunity quivocal onsistent with immunity		
	os IgG antibody suggests immuniz infection with mumps virus.	ation	
RUBELLA AB (IGG), IMMUNE STATUS Index	3.79 Interpretation	Index	AT
<0.90 $0.90-0.99$ $> or = 1.00$	Not consistent with immunity Equivocal Consistent with immunity		
	ella IgG antibody suggests c or current infection with		AMD
CASCADING REFLEX T. PALLIDUM AB	Negative	Negative	71110
syphilis) were detection; does not eximple the infection; testing of weeks after this specific than the second testing of the second testing the second testing the second testing testing the second testing	pallidum (the agent causing cted in the specimen. This result collected very recent T. pallidum of a second specimen, collected ecimen, is recommended if the intent infection is high.	2-4	
HEPATITIS B SURFACE AB IMMUNITY, QN	<5 L	> OR = 10 mIU/mL	AT



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Test Name Out Of Range Reference Range Lab In Range Patient does not have immunity to hepatitis B virus. For additional information, please refer to http://education.questdiagnostics.com/faq/FAQ105 (This link is being provided for informational/ educational purposes only). CHLAMYDIA TRACHOMATIS AT RNA, TMA, UROGENITAL NOT DETECTED NOT DETECTED See Endnote 1 NEISSERIA GONORRHOEAE AT RNA, TMA, UROGENITAL NOT DETECTED NOT DETECTED See Endnote 1 ABO GROUP AND RH TYPE AT ABO GROUP

RH(D) POSITIVE

For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ111 (This link is being provided for informational/educational purposes only.)

Endnote 1

RH TYPE

The analytical performance characteristics of this assay, when used to test SurePath(TM) specimens have been determined by Quest Diagnostics. The modifications have not been cleared or approved by the FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

For additional information, please refer to https://education.questdiagnostics.com/faq/FAQ154 (This link is being provided for information/educational purposes only.)





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Infectious Diseases

Test Name	Result	Reference Range	Lab
HIV 1/2 ANTIGEN/ANTIBODY, FOURTH GENERATION W/RFL			AT
HIV AG/AB, 4TH GEN	NON-REACTIVE	NON-REACTIVE	

HIV-1 antigen and HIV-1/HIV-2 antibodies were not detected. There is no laboratory evidence of HIV infection.

PLEASE NOTE: This information has been disclosed to you from records whose confidentiality may be protected by state law. If your state requires such protection, then the state law prohibits you from making any further disclosure of the information without the specific written consent of the person to whom it pertains, or as otherwise permitted by law. A general authorization for the release of medical or other information is NOT sufficient for this purpose.

For additional information please refer to http://education.questdiagnostics.com/faq/FAQ106 (This link is being provided for informational/educational purposes only.)

The performance of this assay has not been clinically validated in patients less than 2 years old.

Physician Comments:

PERFORMING SITE:

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