



Patient Information	Specimen Information	Client Information
DOB: AGE: Gender: Phone: Patient ID: Health ID:	Specimen: Requisition: Lab Ref #: Collected: Received: Reported:	

Test Name	In Range	Out Of Range	Reference Range	Lab
MEASLES, MUMPS, AND RUBELLA (MMR) AB (IGG) PANEL, IMMUNE STATUS				
MEASLES AB (IGG), IMMUNE STATUS	>300.00		AU/mL	KS
AU/mL	Interpretation			
<13.50	Not consistent with immunity			
13.50-16.49	Equivocal			
>16.49	Consistent with immunity			
<p>The presence of measles IgG suggests immunization or past or current infection with measles virus.</p> <p>For additional information, please refer to http://education.QuestDiagnostics.com/faq/FAQ162 (This link is being provided for informational/educational purposes only.)</p>				
MUMPS VIRUS AB (IGG), IMMUNE STATUS	158.00		AU/mL	KS
AU/mL	Interpretation			
<9.00	Not consistent with immunity			
9.00-10.99	Equivocal			
>10.99	Consistent with immunity			
<p>The presence of mumps IgG antibody suggests immunization or past or current infection with mumps virus.</p>				
RUBELLA AB (IGG), IMMUNE STATUS	2.84		Index	KS
Index	Interpretation			
<0.90	Not consistent with immunity			
0.90-0.99	Equivocal			
> or = 1.00	Consistent with immunity			
<p>The presence of rubella IgG antibody suggests immunization or past or current infection with rubella virus.</p>				
HEPATITIS B SURFACE AB IMMUNITY, QN	826		> OR = 10 mIU/mL	KS
<p>Patient has immunity to hepatitis B virus.</p> <p>For additional information, please refer to http://education.questdiagnostics.com/faq/FAQ105 (This link is being provided for informational/educational purposes only).</p>				
CHLAMYDIA TRACHOMATIS RNA, TMA, UROGENITAL	NOT DETECTED		NOT DETECTED	KS
<p>See Endnote 1</p>				
NEISSERIA GONORRHOEAE				KS



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RNA, TMA, UROGENITAL <i>See Endnote 1</i>	NOT DETECTED		NOT DETECTED	
RPR (MONITOR) W/REFL TITER	NON-REACTIVE		NON-REACTIVE	KS
ABO GROUP AND RH TYPE ABO GROUP RH TYPE	O RH(D) POSITIVE			KS

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

Endnote 1 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.)



Patient Information	Specimen Information	Client Information
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Infectious Diseases

Test Name	Result	Reference Range	Lab
HIV 1/2 ANTIGEN/ANTIBODY, FOURTH GENERATION W/RFL			KS
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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