

## Quest Diagnostics

## SPECIMEN INFORMATION

SPECIMEN: LV340301F  
REQUISITION: 00954919  
Lab ref no:

COLLECTED: 2019/02/14 12:54  
RECEIVED: 2019/02/14 12:55  
REPORTED: 2019/02/19 14:43

## PATIENT INFORMATION

**John, Smith**  
DOB: September 11, 1984  
AGE: 20  
GENDER: Male  
FASTING: Unknown

Clinical Info:

REPORT STATUS: FINAL

## ORDERING PHYSICIAN

**Jane, Doe**

## CLIENT INFORMATION

2019-02-19 14:43:00 -0800  
Lab Testing API  
280 Madison Avenue  
Room 912, 9th Floor  
New York, NY 10016

Test Name	Result	Flag	Reference Range	Lab
<b>ABO GROUP AND RH TYPE</b>				
ABO GROUP	A	NORMAL		01
RH TYPE	RH(D) POSITIVE	NORMAL		01

**CHLAMYDIA/N. GONORRHOEAE RNA, TMA, UROGENITAL**

CHLAMYDIA TRACHOMATIS RNA, TMA, UROGENITAL	NOT DETECTED	NORMAL	NOT DETECTED	02
NEISSERIA GONORRHOEAE RNA, TMA, UROGENITAL	NOT DETECTED	NORMAL	NOT DETECTED	02
COMMENT		NORMAL		02
This test was performed using the APTIMA COMBO2 Assay (Gen-Probe Inc.).				

The analytical performance characteristics of this assay, when used to test SurePath specimens have been determined by Quest Diagnostics.

**HEPATITIS B SURFACE AB IMMUNITY, QN**

HEPATITIS B SURFACE AB IMMUNITY, QN <5 LOW > OR = 10 mIU/mL 01

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).

**HIV 1/2 ANTIGEN/ANTIBODY, FOURTH GENERATION W/RFL**

HIV AG/AB, 4TH GEN	NON-REACTIVE	NORMAL	NON-REACTIVE	01
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.

**MMR (IGG) PANEL (MEASLES, MUMPS, RUBELLA)**

MEASLES ANTIBODY (IGG)	115.00	NORMAL	AU/mL	01
AU/mL Interpretation				
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<25.00 Negative				
25.00-29.99 Equivocal				
>29.99 Positive				

A positive result indicates that the patient has antibody to measles virus. It does not differentiate between an active or past infection. The clinical diagnosis must be interpreted in conjunction with clinical signs and symptoms of the patient.

MUMPS VIRUS ANTIBODY (IGG)	205.00	NORMAL	AU/mL	01
AU/mL Interpretation				
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<9.00 Negative				
9.00-10.99 Equivocal				
>10.99 Positive				

A positive result indicates that the patient has antibody to mumps virus. It does not differentiate between an active or past infection. The clinical diagnosis must be interpreted in conjunction with clinical signs and symptoms of the patient.

RUBELLA ANTIBODY (IGG)	1.63	NORMAL	index	01
Index Interpretation				
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<0.90 Not consistent with Immunity				
0.90-0.99 Equivocal				
> or = 1.00 Consistent with Immunity				

The presence of rubella IgG antibody suggests immunization or past or current infection with rubella virus.

**RPR (DX) W/REFL TITER AND CONFIRMATORY TESTING**

RPR (DX) W/REFL TITER AND CONFIRMATORY TESTING	NON-REACTIVE	NORMAL	NON-REACTIVE	01
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**Performing Laboratory Information:**

01: Quest Diagnostics-Wood Dale, 1355 Mittel Blvd, Wood Dale IL, phone: , Medical Director: MD Anthony V Thomas