

Patient Information	Specimen Information	Client Information
RILLION, ELIZA DOB: 06/18/1996 AGE: 24 Gender: F Phone: 678.603.7274 Patient ID: 4727	Specimen: AL575500E Requisition: 8707957 Lab Ref #: 00683244 Collected: 06/04/2021 / 13:30 CDT Received: 06/04/2021 / 23:17 CDT Reported: 06/07/2021 / 14:39 CDT	Client #: 49100638 QATL0000 BARTON, LINDSEY AUM NURSING CARE CENTER 7461 EAST DRIVE #102 MONTGOMERY, AL 36117

PROVIDER: HAVARD JYSSICA

SARS CoV 2 (COVID-19) Tests

Test Name	Result	Reference Range	Lab
SARS COV 2 AB (IGG) SPIKE, SEMI QN	2.06 H	<1.00 index	AT
Reference Range			
INDEX	INTERPRETATION		
<1.00	Negative		
> or = 1.00	Positive		
This test is intended to help identify individuals with antibodies to SARS-CoV-2 (COVID-19). The results of this semi-quantitative test should not be interpreted as an indication or degree of immunity or protection from reinfection. A test result that is 1.00 or more (Positive) means antibodies to SARS-CoV-2 were detected in the blood sample by the test. This could mean that the individual may have an immune response to a recent or prior infection with SARS-CoV-2. Positive results may occur after COVID-19 vaccination, but the clinical significance of a positive antibody result for individuals that have received a COVID-19 vaccine is unknown, and the performance of the test has not been established in COVID-19 vaccinees. False positive results for the test may occur due to cross-reactivity from pre-existing antibodies or other possible causes. A test result that is less than 1.00 (Negative) means that antibodies were not detected in the blood sample by the test. This could mean that the individual has not been previously infected with SARS-CoV-2. The clinical significance of a negative antibody result for individuals that have received a COVID-19 vaccine is unknown. The performance of the test has not been established in COVID-19 vaccinees. False negative results for the test may occur if the individual's antibodies have not reached a sufficient level for the test to be able to detect them. Antibodies can take up to two to three weeks (sometimes longer) to develop after someone is infected. How long antibodies to SARS-CoV-2 last after infection is not known. This test should not be used to diagnose an active SARS-CoV-2 infection. If an active infection is suspected, direct molecular or antigen testing for SARS-CoV-2 is recommended. Please review the "Fact Sheets" available for healthcare providers and patients using the following websites: https://www.QuestDiagnostics.com/home/Covid-19/HCP/antibody/fact-sheet8 https://www.QuestDiagnostics.com/home/Covid-19/Patients/antibody/fact-sheet8 Healthcare Providers: For additional information please refer to: http://education.questdiagnostics.com/faq/FAQ219 (This link is being provided for informational/educational purposes only.) This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by authorized laboratories. The FDA authorized labeling is available on the Quest Diagnostics website: www.QuestDiagnostics.com/Covid19.			
Physician Comments:			

PERFORMING SITE:

AT QUEST DIAGNOSTICS-ATLANTA, 1777 MONTREAL CIRCLE, TUCKER, GA 30084-6802 Laboratory Director: ANDREW N YOUNG, MD, PHD. CLIA: 11D0255931

LIST OF RESULTS PRINTED IN THE OUT OF RANGE COLUMN:

SARS COV 2 AB (IGG)			
SPIKE, SEMI QN	2.06 H	<1.00 index	AT
Reference Range			

Patient Information	Specimen Information	Client Information
RILLION, ELIZA DOB: 06/18/1996 AGE: 24 Gender: F Patient ID: 4727	Specimen: AL575500E Collected: 06/04/2021 / 13:30 CDT Received: 06/04/2021 / 23:17 CDT Reported: 06/07/2021 / 14:39 CDT	Client #: 49100638 BARTON, LINDSEY

LIST OF RESULTS PRINTED IN THE OUT OF RANGE COLUMN:

INDEX	INTERPRETATION
<1.00	Negative
> or = 1.00	Positive

This test is intended to help identify individuals with antibodies to SARS-CoV-2 (COVID-19). The results of this semi-quantitative test should not be interpreted as an indication or degree of immunity or protection from reinfection.

A test result that is 1.00 or more (Positive) means antibodies to SARS-CoV-2 were detected in the blood sample by the test. This could mean that the individual may have an immune response to a recent or prior infection with SARS-CoV-2. Positive results may occur after COVID-19 vaccination, but the clinical significance of a positive antibody result for individuals that have received a COVID-19 vaccine is unknown, and the performance of the test has not been established in COVID-19 vaccinees. False positive results for the test may occur due to cross-reactivity from pre-existing antibodies or other possible causes.

A test result that is less than 1.00 (Negative) means that antibodies were not detected in the blood sample by the test. This could mean that the individual has not been previously infected with SARS-CoV-2. The clinical significance of a negative antibody result for individuals that have received a COVID-19 vaccine is unknown. The performance of the test has not been established in COVID-19 vaccinees. False negative results for the test may occur if the individual's antibodies have not reached a sufficient level for the test to be able to detect them. Antibodies can take up to two to three weeks (sometimes longer) to develop after someone is infected. How long antibodies to SARS-CoV-2 last after infection is not known.

This test should not be used to diagnose an active SARS-CoV-2 infection. If an active infection is suspected, direct molecular or antigen testing for SARS-CoV-2 is recommended.

Please review the "Fact Sheets" available for healthcare providers and patients using the following websites:
<https://www.QuestDiagnostics.com/home/Covid-19/HCP/antibody/fact-sheet8>

<https://www.QuestDiagnostics.com/home/Covid-19/Patients/antibody/fact-sheet8>

Healthcare Providers: For additional information please refer to:
<http://education.questdiagnostics.com/faq/FAQ219>
(This link is being provided for informational/educational purposes only.)

Patient Information	Specimen Information	Client Information
RILLION, ELIZA DOB: 06/18/1996 AGE: 24 Gender: F Patient ID: 4727	Specimen: AL575500E Collected: 06/04/2021 / 13:30 CDT Received: 06/04/2021 / 23:17 CDT Reported: 06/07/2021 / 14:39 CDT	Client #: 49100638 BARTON, LINDSEY

LIST OF RESULTS PRINTED IN THE OUT OF RANGE COLUMN:

This test has been authorized by the FDA under an
Emergency Use Authorization (EUA) for use by authorized
laboratories. The FDA authorized labeling is available
on the Quest Diagnostics website:
www.QuestDiagnostics.com/Covid19.