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I would to reschedule for a later date if possible.

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**MEDLAB2020, INC.**

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Lab Director: K. Cameron Campbell MD

PATIENT	SAMPLE	PHYSICIAN
Name: Theresa Styles	Specimen Type: Mid-turbinate	Name: Matthew Abinante
Date of Birth: 10/10/1971	Collection Date: 1/10/2023 06:35:49 PST	Address: 18800 Delaware St. Ste 800
Gender: F	Reported Date: 1/12/2023 09:34:26 PST	Huntington Beach, CA 92648
Accession #: 00-5486500		NPI: 1740685387
Passport Number:		
Passport Country:		

Testing Ordered COVID-19 (SARS-CoV-2) MOLECULAR ASSAY BY - RT-PCR

Tested Pathogens	Pathogen Type	Test Result
SARS-CoV-2	VIRUS	DETECTED

DETECTED (Positive): A positive result indicates that RNA from the SARS-CoV-2 virus was detected in the patient sample. Patients infected with this virus may be asymptomatic but are presumed to be contagious. Patient results will change over time. Positive results do not rule out bacterial co-infection with other viruses.

NOT DETECTED (Negative): A negative result means that the virus that causes COVID-19 was not found in your sample at the time the sample was taken. Negative results do not preclude SARS-CoV-2 virus infection and should not be used as the sole basis for diagnostic decisions. Negative results must be combined with clinical observations, patient history, and epidemiological information. For COVID-19, a negative test result for a sample collected while a person has symptoms usually means that COVID-19 did not cause the recent illness. However, it is possible for this test to give a negative result that is incorrect (false negative) in some people with COVID-19. This means that you could possibly still have COVID-19 even though the test is negative. If clinically indicated, repeat testing should be performed.

INDETERMINATE: An indeterminate result can occur when there is poor sample collection or the viral load is very low at the time of collection so to not elicit a positive or a negative result. If clinically indicated, repeat testing should be performed.

REFERENCE RANGE (Expected Result): Negative (Not Detected). The reference range is not the patient's test result.

This test has not been FDA cleared or approved. This test has been validated under an Emergency Use Authorization (EUA) guidance and it has been validated in accordance with the FDA's Guidance Document Policy for Diagnostics Testing in Laboratories Certified to Perform High Complexity Testing under CLIA. This test is only authorized for the duration of time the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of SARS-CoV-2 virus and/or diagnosis of COVID-19 infection under section 564(b)(1) of the Act, 21 U.S.C. 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.



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