

Medical Diagnostic Laboratories

2439 Kuser Road **Toll Free:** 877-269-0090 Hamilton, NJ 08690 **Fax:** 609-570-1050

www.mdlab.com

SPECIMEN

Type Source Collected Receiv

Received Reported

NasoSwab Nasopharyngeal 04/15/2024 04/16/2024 04/19/2024

2 PATIENT

MDL#

13364148

FINAL

DOE, JANE

555 MAIN ROAD

ANYTOWN, NJ 12345-6789

DOB: 11/30/1998 (Age 25)

Gender: Female
Ethnicity: Not provided
Patient ID: 82100
Home #: 123-456-7890

CLIENT

NPI: 0987654321

DOE FAMILY PRACTICE JOHN DOE, MD 1234 FIRST AVENUE

ANYTOWN, NJ 12345-6789

Tel: (555) 555-1234 **Fax:** 555-555-1235

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Pathogens Detected

1131 SARS-CoV-2 [COVID-19]

POSITIVE

*This test was developed and its performance characteristics determined by the laboratory. It has not been cleared or approved by the U.S. Food and Drug Administration. The FDA has determined that such clearance or approval is not necessary.

NasoSwab-1;1131:SARS-CoV-2 [COVID-19] by Reverse Transcription Real-Time PCR (CDC N1, N2, RP Assay)

This test was developed and its performance characteristics determined by Medical Diagnostic Laboratories L.L.C. This test has not been FDA cleared or approved. This test has been authorized by the FDA under an Emergency Use Authorization (EUA). This test 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 prior to Emergency Use Authorization for Coronavirus Disease-2019 during the Public Health Emergency" issued on February 29, 2020. FDA-independent review of this validation is pending. 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.

A positive result is provided for bacteria, virus, parasites, and/or fungal species when PCR amplification (real-time PCR), sequence information (Pyrosequencing), and/or sequencing analysis occurs above cut-off levels established by the laboratory. Pertinent reference intervals for the tests reported above are available from the laboratory upon request.

Medical Director, Jing-Jing Yang, M.D.

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05/20/2024