Final Report

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ACCESSION #: 1039785
REQUISITION #:
ORDERED: 06/19/2021 12:28 PM
COLLECTED: 06/19/2021 12:28 PM
RECEIVED: 06/19/2021 10:34 PM
REPORTED: 06/20/2021 02:50 PM
SPECIMEN TYPE:

Test Name	Result	Flag
COVID-19 Nasopharyngeal (NP)		
SARS-CoV-2*	NOT DETECTED	**NOT DETECTED**

You can view the Patient Fact Sheet here: http://exceltox.com/assets/img/EUA-Thermo-TaqPath-patient.pdf

*This test was performed by real time PCR methodology *This test has not been FDA cleared or approved; *This test has been authorized by FDA under an EUA for use by authorized laboratories; Exceltox Laboratory is authorized as a high complexity laboratory * This test has been authorized only for the detection of nucleic acid from SARS-CoV-2, influenza A and influenza B, not for any other viruses or pathogens; and This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner. * Negative results do not preclude acute SARS CoV-2 infection.

** This test has not been FDA cleared or approved; ** This test has been authorized by FDA under an EUA for use by authorized laboratories; ** This test has been authorized only for the presence of IgG and/or IgM antibodies against SARS-CoV-2, not for any other viruses or pathogens; and ** This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 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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