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CLIA#: 33D0985206

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Client: LABWORQ

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Phys: LABWORQ

Patient: HEE, CAMERON CHARLES

215 WEST 95 STREET 3A

NY, NY 10025

**DOB.** 05/23/1998 **Age:** 23

**Phone:** ( ) - **ID#:** R0000051988

 Accession:2112290931
 Coll. Date: 01/10/22
 Recv. Date: 01/10/22
 Print Date: 01/12/22

 Chart#AL-140344
 Coll. Time: 13:42
 Recv. Time: 13:42
 Print Time: 08:54

3985

First reported on: 01/11/22 Final report date: 01/11/22

Test Name Within Range Out Of Range Ref.Range Units Previous Result (Date)

## **GENETIC AND MOLECULAR TESTING**

SARS-CoV-2 RNA PCR NEGATIVE

Tech:RICHA Date:01/11/22 19:22

The SARS-CoV-2 test is intended for the qualitative detection of nucleic acid from SARS-CoV-2(

who meet COVID-19 clinical and/or epidemiological criteria. For lower respiratory tract specimens, the assay is submitted for authorization by FDA under the Emergency Use Authorization (EUA).

## Disclaimer:

This test has not been Food and Drug Administration (FDA) cleared or approved and has been authorized by FDA under an Emergency Use Authorizatio (EUA). The 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 SARS-CoV-2 under Section 564(b)(1) of the Act, 21 U.S.C. section 360bbb-3(b)(1), unless the authorization is terminaed or revoked sooner. Alliance Laboratories is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA), 42 U.S.C section 263a, to perform high complexity tests.

Fact Sheet for Healthcare Providers: https://www.cdc.gov/coronavirus/2019-nCoV/hcp/index.html

Fact Sheet for Patients:

https://www.cdc.gov/coronavirus/2019-ncov/your-health/index.html

SARS-CoV-2 Collection NP SWAB

**Report Status: FINAL** 

END OF REPORT \_\_\_\_\_

Originally Reported On:01/10/22 18:54