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Client: LABWORQ 3985
455 GRAHAM AVE
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Phys: LABWORQ

Patient: HEE, CAMERON CHARLES
215 WEST 95 STREET 3A
NY, NY 10025
DOB: 05/23/1998 **Age:** 23 **Sex:** M
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: 18:54
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)
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GENETIC AND MOLECULAR TESTING

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

SARS-CoV-2 RNA PCR **NEGATIVE**

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