LabCorp

SPECIMEN INFORMATION
ACCOUNT NO: 04138630
SPECIMEN: 06335802820
REQUISITION: 108017855
Lab ref no:

COLLECTED: 03/04/2022 02:15PM PST
RECEIVED: 03/05/2022
REPORTED: 03/05/2022 07:06PM ET

PATIENT INFORMATION
Kashiwao, Kotaro
DOB: December 22, 2015
AGE: 6
GENDER: Male
FASTING: Unknown
PATIENT ID: 108017855

Clinical Info:

REPORT STATUS: FINAL

ORDERING PHYSICIAN
Mota, Christina
NPI: 1114014263
CLIENT INFORMATION
Walgreens COVID-19

Test Name	Result	Flag	Reference Range	Lab
SARS-CoV-2, NAA				
SARS-CoV-2, NAA	Not Detected	NORMAL	Not Detected	01

This nucleic acid amplification test was developed and its performance characteristics determined by LabCorp Laboratories. Nucleic acid amplification tests include RT-PCR and TMA. This test has not been FDA cleared or approved. This test has been authorized by FDA under an Emergency Use Authorization (EUA). 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.
When diagnostic testing is negative, the possibility of a false negative result should be considered in the context of a patient's recent exposures and the presence of clinical signs and symptoms consistent with COVID-19. An individual without symptoms of COVID-19 and who is not shedding SARS-CoV-2 virus would expect to have a negative (not detected) result in this assay.

Performing Laboratory Information:

01: Integrated Genetics Sequenom, 3595 John Hopkins Court, San Diego CA, 921211121, phone: 877-821-7266, Director: MDPhD Phillip M Cacheris