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Patient Name And Address: Mr. CHARAN SINGH

Name Mr. CHARAN SINGH Collection Date 10-Jan-2021 02:50 PM Age/Sex 34.3 Years/Male **Ordering Physician** Specimen Received 10-Jan-2021 05:20 PM 11-Jan-2021 09:55 AM Institution Walk-in Report Date Lab ID 112101100064 Specimen Collected by Self **Print Date** 11-Jan-2021 10:24 AM Vial ID 10830063 Status **Final Report**

Research	& 1	Development

Test Name Status Result Unit	Reference Interval

SARS-COV-2 Real-Time PCR, Qualitative *

Specimen Information Nasal and Throat Swab

Clinical Indication Asymptomatic (Required for travelling purpose)

Method: Real-time Polymerase Chain Reaction (RTPCR)

Results-COVID

E Gene Not Detected S Gene Not Detected SARS-CoV-2 Detection Negative

Comments

This nucleic acid based test is useful for detection of coronavirus disease 2019 (COVID-19) illness due to severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)

This test **should be requested only on** patients meeting current clinical and/or epidemiologic criteria defined by ICMR, state, or local public health directives:

https://icmr.nic.in/content/covid-19 (https://icmr.nic.in/sites/default/files/upload_documents/Strategy_COVID19_testing_India.pdf)

Clinical Information

Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) is a positive-sense, single-stranded RNA virus that causes coronavirus disease 2019 (COVID-19). Like other coronaviruses that infect humans, SARS-CoV-2 can cause both upper and lower respiratory tract infection. Symptoms can range from mild (ie, the common cold) to severe (ie, pneumonia) in both healthy and immunocompromised patients. SARS-CoV-2 transmission occurs primarily via respiratory droplets. During the early stages of COVID-19, symptoms maybe nonspecific and resemble other common respiratory tract infections, such as influenza. If testing for other respiratory tract pathogens is negative, specific testing for SARS-CoV-2 may be warranted.

Test Description

Kit is based on real-time PCR technology for qualitative detection and differentiation of lineage B-betacoronavirus (B- βCoV) and severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) specific RNA.

The kit targets $B-\beta CoV$ (E gene) and SARS-CoV-2 (S gene). The assay includes a heterologous amplification system (Internal Control) to identify possible RT-PCR inhibition and to confirm the integrity of the reagents of the kit.

Interpretation

Result	Interpretation
Detected (Both S and E gene)	Provided sample is positive for SARS-CoV-2
Not Detected (Both S and E gene)	Provided sample is negative for SARS-CoV-2
Indeterminate (Only E or only S gene detected)	Repeat sample is recommended
Inconclusive	Repeat sample is recommended

A **Detected** result indicates that severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) RNA is present and suggests the diagnosis of coronavirus disease 2019 (COVID-19). Test result should always be considered in the context of patient's clinical history, physical examination, and epidemiologic exposures when making the final diagnosis.

Dr Jatinder Kaur, PhD

Director MolQ Laboratory Head Genomics and Molecular Biology, Biochemistry Dr Gulshan Yadav, MD

Consultant Pathology

Dr Megha Kaushik, PhD Consultant Molecular Biology A **Not Detected** result indicates that SARS-CoV-2 is not present in the patient's specimen. However, this result may be influenced by the stage of the infection and the quality of the specimen collected for testing.

An **Indeterminate** result suggests that the patient may be infected with a variant SARS-CoV-2 or SARS-related coronavirus. Additional testing with an alternative molecular method is recommended on a newly collection specimen.

An **Inconclusive** result indicates that the presence or absence of SARS-CoV-2 RNA in the specimen could not be determined with certainty after repeat testing in the laboratory, possibly due to RT-PCR inhibition. Submission of a new specimen for testing is recommended.

Cautions

- The sensitivity of the assay is dependent on the quality of the specimen collected for testing.
- The test is specific for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), and positive test results do not exclude the
 possibility of concurrent infection with other respiratory viruses.
- Negative results do not preclude infection with SARS-CoV-2 and should not be used as the sole basis for decisions on treatment or other patient care management.

Note

- ICMR Registration Number for COVID-19 is MLG001.
- Test is conducted as per the guidelines recommended by WHO.
- Kindly consult referring Physician/ Authorized Government Hospital for appropriate follow-up.

End of Report

For related test information on this accession, please visit website www.molq.in or email at contact@molq.in

The tests marked with an * are not accredited by NABL

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