

Vijaya Diagnostic Centre

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LABORATORY TEST REPORT

Regn Date : 13/11/2021 14:32 Sample Collection : 13/11/2021 14:35

Name : MS. P AKHILA Print Date : 15/11/2021 11:40

Regn No : 352192483 Age / Sex : 20 Years / Female

Ref By : SELF Regn Centre : Kharkhana - 35 Sample Type : Swab Ref no. : S121406296387



COVID -19 TESTING - SARS -CoV-2 RNA

TEST NAME RESULT

SARS-CoV-2 (RdRp gene) : NEGATIVE

Method: Real Time RT-PCR

INTERPRETATION:

Result | Remarks

Positive | RNA specific to SARS-CoV-2 Detected.

Negative | RNA specific to SARS-CoV-2 NOT Detected.

Limit of Detection:

• Analytical lower unit of detection <150 viral genome equivalents/mL

Comments:

- Covid 19 Qualitative RT PCR test is an in vitro qualitative PCR assay for the qualitative detection of Novel Corona Virus 2019 in respiratory specimens
- Test is conducted on Nasopharyngeal swab/ Oropharyngeal swabs and other respiratory specimens collected in viral transport media.
- Detection of confirmatory(RdRp) genes indicates presence of SARS-CoV-2 RNA in the specimen tested.
- Negative results do not preclude SARS-CoV-2 infection and should not be used as the sole basis for
 patient management decisions. They must be correlated with clinical observations, patient history
 and epidemiological information.
- Mutations or polymorphisms in the primer and probe binding sites, presence of PCR inhibition due to host factors may also cause false negative results.
- Fresh sample for RT PCR can be considered after a gap of 2-4 days if there is a strong clinical suspicion/contact of Covid 19 patient
- Repeat sampling and testing of lower respiratory specimen is strongly recommended in a severe or progressive disease.
- This test is a qualitative assay and does not quantify viral load. Various host factors, variability in sample collection / site and techniques used by the laboratories can affect CT values. Therfore, CT values are not an absolute indication of viral load and should be interpreted with caution.

Note:

- ICMR-Registration Number : VIJAY001
- COVID-19 test is conducted with a kit approved by ICMR/CE-IVD/US-FDA.
- · Kindly consult Referring Physician/Authorized Government Hospital for appropriate follow up



MC-2657

DR. VITTAL M.D
CONSULTANT MICROBIOLOGIST

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