

Vijaya Diagnostic Centre

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LABORATORY TEST REPORT

Regn Date : 26/08/2021 09:23 Sample Collection : 26/08/2021 09:28 Print Date Name : MR. KARKALA AMIT : 26/08/2021 20:40 : 30 Years/Male Regn No : 682028099 Age / Sex Ref By : SELF Regn Centre : Karmanghat - 68 : S474332024496347 Sample Type : Swab Ref no.

COVID -19 TESTING - SARS -CoV-2 RNA

TEST NAME

RESULT

SARS-CoV-2 : NEGATIVE

Method: Real Time RT-PCR

INTERPRETATION:

Result | Remarks

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

Toshive | Idivi specific to SARS-COV-2 Detected.

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

Limitations:

- 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 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.

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 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

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