

CID : 2200146708

Name : MRS.MOHITA UPADHYAY

Age / Gender : 33 Years / Female

Consulting Dr.

Reg. Location : Maitri Park, Chembur East

Authenticity Check

Use a OR Code Scanner Application To Scan the Code

Collected

Reported

: 02-Jan-2022 / 08:29

: 02-Jan-2022 / 21:37

### Real time Qualitative RT-PCR detection of 2019-nCOV RNA / COVID-19 RNA **RESULT PARAMETER**

SARS-CoV-2 SARS-CoV-2: Not Detected (Negative)

Kit description-E gene as screening and ORF1ab/RdRp or N gene as target gene, Cutoff: <35

## Clinical Significance:

ICMR Registration No: Andheri-Mumbai-SUBUR001, Pune-SUDIIPLPMH

Specimen: Nasopharyngeal & Oropharyngeal swab in VTM Method: Real time RT-PCR

- Ct Value indicates the infectivity and not the severity of infection.
- \* ICMR Recommended kits are used for reporting. All the positive cases will be notified to ICMR for further surveillance.
- \* Clinical correlation with patient history, radiology findings and co-infection with other viruse infections is necessary to be determined especially in cases with Border line positive Ct Values Borderline Positive cases (Ct value>30) may be give variable results on repeat testing. The possible reasons could be the variations in kit and instruments used.

### Limitations:

- Optimum Specimen types and timing of peak viral levels during infections caused by 2019-nCOV have not been determined. Collection of multiple specimens (Types & Time Points) may be necessary in view of suspected clinical history. The repeat specimen may be considred aftee a gap of 2-4 days after the collection of first specimen for additional testing if required(other respiratory pathogens)
- \* Negative results do not preclude SARS-CoV-2 infection and should not be used as the sole basic for patient management decisions.
- \* This test is qualitative assay and does not qaulify viral load. Various host factors, Viral factors, Variability in the sample collection /site and techniques used by laboratory can effect the ct values.

Therefore, Ct values are not an absolute indication of viral load and should be interpreted with caution.

# Factors leading to false nagetive RT-PCR report:

- Inadequate specimen collection, poor qaulity of sample and non-representative sample.
- Technical reasons- PCR inhibitor, analytical sensitivity of kit used.
- \* Active recombination &/mutations in target gened used for detection of SARS-CoV-2 virus

- 1. Diagnostic detection of 2019-n-CoV by real-time RT-PCR, Berlin Jan 17th, 2020.
- 2.Labcorp COVID-19 RT-PCR test EUA Summary/ COVID 19 RT-PCR test (Laboratory corporation of America)
- \* Sample processed at Molecular Diagnostics Laboratory, CPL, Andheri West
  \*\*\* End Of Report \*\*\*







**Dr.HEENA SATAM** M.Sc. Microbiology, PhD. **Biochemistry Molecular Biologist** 

**Dr.SHASHIKANT DIGHADE** M.D. (PATH) **Pathologist** 

Page 1 of 1

ADDRESS: 2nd Floor, Aston, Sundervan Complex, Above Mercedes Showroom, Andheri West - 400053

HEALTHLINE - MUMBAI: 022-6170-0000 | OTHER CITIES: 1800-266-4343

For Feedback - customerservice@suburbandiagnostics.com | www.suburbandiagnostics.com