



MOLECULAR LABORATORY - CHEST HOSPITAL

REPORT FORM FOR COVID-19

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| | | | |
|--------------|---------------|-----------|-----------------------|
| Name | : SARANYA S | Sample ID | : CH220204003 |
| UHID | : 2022/002364 | Collected | : 02/03/2022 12:45 pm |
| Gender | : Female | Received | : 02/03/2022 3:00 pm |
| Age | : 25 | Reported | : 03/03/2022 6:33 pm |
| Ref. Doctor | : DR GIRISH G | | |
| Reference ID | : | | |

DEPARTMENT OF MOLECULAR BIOLOGY

| Test Name | Sample | Result | Method |
|----------------------------|--|----------|---------------------------------|
| SARS CoV-2 qualitative PCR | Nasopharyngeal/Oropharyngeal swabs in VTM*/MTM** | Positive | RT-PCR-CRISPR Cas-9-based assay |

SRF ID : 3433/KZD/20220312

ICMR Registration number of the lab: **CHMLCK**

Result Interpretation

Negative: Absence of SARS Cov-2 in the given specimen.

Positive: Presence of SARS Cov-2 in the given specimen.

*VTM: Viral Transport Medium, **MTM: Molecular Transport Medium



COMMENTS:

TATA MD CHECK CRISPR SARS CoV-2 Kit 1.0, Powered by FELUDA is CRISPR - Case 9 based in-vitro diagnostic test kit, capable of detecting nucleic acid sequence of novel corona virus, Severe Acute Respiratory Syndrome Corona virus - 2(SARS CoV-2) in respiratory specimens (Nasopharyngeal/oropharyngeal swabs) of individuals suspected to have contracted corona virus disease COVID-19.

The test is designed to detect target sequence, in the S gene (Spike protein gene). In the RNA of SARS CoV-2, the assay uses the RNA extracted from nasopharyngeal/oropharyngeal swab specimens from patients suspected to have COVID 19. The RNA is first converted into cDNA (Complementary DNA) by reverse transcription and amplified through Polymerase Chain Reaction (PCR), specifically generated for the S target sequences. The complex is detected by a method FELUDA (FnCas9 Editor Linked Uniform Detection Assay) which used lateral flow strips to show the presence or absence of the target sequences as a visible band.

NOTE:

Negative result does not rule out the possibility of COVID-19 infection. Presence of inhibitors, mutations and insufficient RNA specific to SARS CoV-2 can influence the test results. Please correlate the results with clinical findings. A negative result with single upper respiratory tract sample does not rule out SARS CoV-2 infection. Hence in such cases a repeat sample should be sent. Lower respiratory tract samples like sputum, BAL, ET aspirate are appropriate sample especially in sever and progressive lung disease. COVID-19 test is conducted as per the kit approved by ICMR Kindly consult referring physician / authorized hospital for appropriate follow up.



Authorized signatory

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Status: Final Report, Printed On: 03/03/2022 6:33 pm

All steps in the testing procedures are performed in an NABL accredited; ICMR and Government of Kerala approved Laboratory

... End of the report ...

MC-4774