Ministry of Health & Family Welfare
Government of India

# Certificate for COVID-19 Vaccination

Issued in India by Ministry of Health & Family Welfare, Govt. of India

**Certificate ID 62276829591**

## Beneficiary Details

| | |
|---|---|
| Beneficiary Name / பலனாளியின் பெயர் | **Revathi** |
| Age / வயது | **37** |
| Gender / பாலினம் | **Female** |
| ID Verified / அடையாளச் சான்று | **Aadhaar # XXXXXXXX6180** |
| Unique Health ID (UHID) | |
| Beneficiary Reference ID | **31781326162170** |
| Vaccination Status / தடுப்பூசி நிலை | **Fully Vaccinated (2 Doses)** |

## Vaccination Details

| | |
|---|---|
| Vaccinated By / தடுப்பூசியை வழங்கியவர் | **Poonkodi** |
| Vaccination At / தடுப்பூசி வழங்கப்பட்ட இடம் | **THAKKOLAM APHC COVAXIN, Ranipet, Tamil Nadu** |

| Dose Number<br>டோஸ் எண் | Date of Dose<br>டோஸ் வழங்கப்பட்ட தேதி | Vaccine Name<br>தடுப்பூசியின் பெயர் | Batch Number<br>பேட்ச் எண் | Vaccine Type<br>தடுப்பூசி வகை | Manufacturer<br>உற்பத்தியாளர் |
|---|---|---|---|---|---|
| 1/2 | 01 Mar 2022 | COVAXIN | 37C521040 | COVID-19 vaccine, Inactivated virus | Bharat Biotech, India |
| 2/2 | 31 Mar 2022 | COVAXIN | 37C521040 | COVID-19 vaccine, Inactivated virus | Bharat Biotech, India |

"மருந்து மற்றும் மனவுறுதியுடன்
Together, India will defeat COVID-19"

- பிரதம மந்திரி நரேந்திர மோதி

In case of any adverse events, kindly contact the nearest Public Health Center/ Healthcare Worker/District Immunization Officer/State **Helpline No. 1075**

ஏதேனும் எதிர்மறை விளைவுகள் ஏற்பட்டால், தயவு செய்து அருகாமையிலுள்ள பொது சுகாதார மையம் / ஆரோக்கியப் பராமரிப்புப் பணியாளர் / மாவட்ட தடுப்பூசி அலுவலர் / மாநில உதவி எண். 1075ஐ தொடர்பு கொள்ளவும்.

COWIN
Winning Over COVID

**DIAGNOSTIC REPORT**

Patient Ref. No. 323000001136898

MC-5320

**CLIENT CODE :** C000132666

**CLIENT'S NAME AND ADDRESS :**
URL ENTERPRISES
152/10, LOTUS COLONY, ANNA NAGAR EAST,
CHENNAI
CHENNAI 600102
TAMIL NADU INDIA
9940200487

SRL DIAGNOSTICS
Veda Ranghaa Nivas, No:22/97,Dr.Ambedkar Road, Ashok Nagar Avenue
CHENNAI, 600083
TAMIL NADU, INDIA

**PATIENT NAME : REVATHI THAKKALI** PATIENT ID : **REVAF270585323**

ACCESSION NO : **0323VK001985** AGE : 37 Years SEX : Female ABHA NO :

DRAWN : 08/11/2022 10:00:00 RECEIVED : 08/11/2022 11:59:24 REPORTED : 08/11/2022 13:53:03

**REFERRING DOCTOR :** SELF CLIENT PATIENT ID :

**CLINICAL INFORMATION :**

ICMR REG NO: SRLDGNANCH

| Test Report Status **Final** | Results | Biological Reference Interval | Units |
|---|---|---|---|

## MOLECULAR BIOLOGY

**SARS COV -2 REAL TIME PCR**

| | | | |
|---|---|---|---|
| SARS-COV-2 RNA | NEGATIVE | | |

METHOD : REAL TIME RT PCR (OPEN SYSTEM)

**Comments**

SPECIMEN TYPE: NASOPHARYNGEAL & OROPHARYNGEAL SWAB
ORF1ab GENE: NOT DETECTED
N GENE: NOT DETECTED
ICMR SRF ID: 3356811773096

**Interpretation(s)**
SARS COV -2 REAL TIME PCR-SARS-CoV-2, formerly known as 2019-nCoV, is the causative agent of the coronavirus disease 2019 (COVID-19). Main symptoms of the disease include fever, cough and shortness of breath. The virus is spread via person-to-person contact through respiratory droplets produced when a person coughs or sneezes. The SARS-CoV-2 RNA is generally detectable in nasopharyngeal/oropharyngeal swabs during the acute phase of infection. Positive results are indicative of active infection. Real Time PCR assay targets specific genes and can be used for diagnosis of SARS-CoV-2 virus infection which contributes to severe upper respiratory distress and complications.
Positive result indicates that RNA from SARS-CoV-2 was detected in the specimen, and the patient is considered infected with the virus and presumed to be contagious.
Negative test result for this test means that SARS-CoV-2 RNA was not detected in the specimen above the limit of detection of the assay.
Limitations:
• Negative results do not preclude COVID-19 and should not be used as the sole basis for patient management decisions. Negative results must be combined with clinical observations, patient history, and epidemiological information.
• Positive results do not rule out bacterial infection or co-infection with other viruses.
• Optimum specimen types and timing for peak viral levels during infections caused by 2019-nCoV have not been determined. Collection of multiple specimens (types and time points) from the same patient may be necessary to detect the virus.
• Follow-up testing may particularly be important if patient has a clinical picture of viral pneumonia, a potential exposure history, and/or radiographic findings (chest CT o MRI scan) consistent with COVID -19 pneumonia. However repeat testing in the near-term after clearance (within 90 days) should be avoided as prolonged shedding of non-viable virus is not uncommon
• Ct values generated from different assay systems within the same laboratory, or from different laboratories, are not directly comparable and do not necessarily reflect t same viral load due to inter-assay and inter-laboratory variability.
• Variation in timing of sample collection, fluctuations in virus shedding, and difference between detection limit of different testing methods within same or different labs could lead to variation in results particularly during initial phase of infection.
• If the virus mutates in the rRT-PCR target region, 2019-nCoV may not be detected or may be detected less predictably. Inhibitors or other types of interference may produce a false negative result.
• The performance of this test has not been established for monitoring treatment of 2019-nCoV infection.
Note: Test is performed using ICMR approved Kit.
References:
1. Laboratory testing for coronavirus disease 2019 (COVID-19) in suspected human cases. Interim guidance. World Health Organization.
2. Druce et al. JCM. 2011
3. N. Engl. J. Med. 2020, 382, 929–936

****End Of Report****

**Please visit www.srlworld.com for related Test Information for this accession**



Scan to View Details

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**DIAGNOSTIC REPORT**

Patient Ref. No. 323000001136898

SRL Diagnostics

MC-5320

**CLIENT CODE :** C000132666

**CLIENT'S NAME AND ADDRESS :**
URL ENTERPRISES
152/10, LOTUS COLONY, ANNA NAGAR EAST,
CHENNAI
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S. Satheesh Kumar

**Dr. S.Satheesh Kumar,PH.D**
**Consultant Microbiologist**

Srinivas C.N.

**Dr.C.N.Srinivas**
**Vice President – Technical & Head of HLA & TI**

**CONDITIONS OF LABORATORY TESTING & REPORTING**

1. It is presumed that the test sample belongs to the patient named or identified in the test requisition form.
2. All tests are performed and reported as per the turnaround time stated in the SRL Directory of Services.
3. Result delays could occur due to unforeseen circumstances such as non-availability of kits / equipment breakdown / natural calamities / technical downtime or any other unforeseen event.
4. A requested test might not be performed if:
   i. Specimen received is insufficient or inappropriate
   ii. Specimen quality is unsatisfactory
   iii. Incorrect specimen type
   iv. Discrepancy between identification on specimen container label and test requisition form
5. SRL confirms that all tests have been performed or assayed with highest quality standards, clinical safety & technical integrity.
6. Laboratory results should not be interpreted in isolation; it must be correlated with clinical information and be interpreted by registered medical practitioners only to determine final diagnosis.
7. Test results may vary based on time of collection, physiological condition of the patient, current medication or nutritional and dietary changes. Please consult your doctor or call us for any clarification.
8. Test results cannot be used for Medico legal purposes.
9. In case of queries please call customer care (91115 91115) within 48 hours of the report.

**SRL Limited**
Fortis Hospital, Sector 62, Phase VIII,
Mohali 160062



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