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ISO 9001:2015 Certified

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Patient Name: CUNANAN, MARY ANN DARAY

Age : 27Y Gender: FEMALE Date of Birth: 11-13-1993
Address: 0448 PUROK 3, BRGY SAN ROQUE, SAN LUIS PAMPANGA

Physician : NO, PHYSICIAN

Referring Institution:

Nationality : FILIPINO

Transaction No. : 8622264

Laboratory No. | PID : 8021184695 | 2090459321

Date & Time Received: 09-15-2021 08:10 **Date & Time Received**: 09-15-2021 11:08 **Date & Time Reported**: 09-15-2021 14:58

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Passport No.: P5792235A

MOLECULAR BIOLOGY

Test Name: SARS-CoV-2 (causative agent of COVID-19) viral RNA RT-PCR

Qualitative Detection by Real-Time PCR

Result: SARS-CoV-2 NOT Detected

Interpretation: SARS-CoV-2 NEGATIVE

Specimen: Nasopharyngeal and Oropharyngeal Swab

Date & Time Collected: 09-15-2021 08:27

Methodology: This test utilizes the novel coronavirus (2019-nCoV) ORF 1ab Gene and the specific conserved

sequence of coding nucleocapsid protein N gene as the target regions which are designed for the conserved sequence of the double-target genes, to achieve detection of sample RNA through fluorescent signal changes. The PCR detection system uses the positive internal control, which monitors the presence of PCR inhibitors in test specimens by detecting whether the internal control

signal is normal, to avoid a false negative result.

Significance : This test is intended to be used to achieve qualitative detection of SARS-CoV-2, the causative agent

of COVID-19, extracted from nasopharyngeal swabs and oropharyngeal swabs specimen of patients.

Limitations : The detection of viral RNA is dependent on the viral load if the specimen was collected very early in

the infection. Pre-analytical variables (i.e. specimen quality, handling/ transport condition) may also

affect the results. Limit of detection for this is 200 copies/mL.

Test Platform: PCR Kit: CE IVD Marked Sansure Novel Coronavirus (2019-nCoV) Nucleic Acid Diagnostic Kit

RNA Extraction Method: Sansure Sample Release Reagent using One Tube Technology

Instrument: Sansure SLAN-96P PCR Machine

Note: The result obtained must be interpreted by the attending physician in correlation with patient's

pertinent clinical and ancillary findings.

Laboratory report must be clinically correlated by your attending physician.

LÝN M. RAMIREZ, RMT

Lic. No. 63568

Medical Technologist

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DR. PAULO GIOVANNI D. MENDOZA, FPSP Lic. No. 98131

Clinical Pathologist

JAE WARREN 6. ALVENDIA, RMT

OC Officer

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DR. DAPHNEC. ANG, MD, DPSP Lic. No. 102523

Molecular Pathologist