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SOP for Dual Target Detection of COVID-19 (ORF1ab/N Gene)	Reviewed By	
	Approved By	

	Name and position	Signature/Stamp
<b>Author(s)</b>		
<b>Verifiers</b>		Stamp:
<b>Approver</b>		

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1.	PPHRL Lab Staff on duty
2.	Quality Management Record
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<b>Revise before:</b>	After one year of approver/ as per need

**Changes in this version compared to previous version:**

To be done after revision as per international accreditation guidelines
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## **Purpose**

The purpose of this SOP is to define procedure for dual target detection of Detection of COVID-19 (ORF1ab/N Gene) guidelines to use Rotor Gene Q for the

## **Scope**

The detection of COVID-19 genes (ORF1 ab and N gene) will be done on the Rotor-Gene Q instrument, which is designed to perform real-time and end-point thermal cycling using the polymerase chain reaction (PCR), and high-resolution melting analysis (HRM) in molecular biology applications.

## **Responsibility**

PACP Advanced Diagnostic Lab (Authorized) Staff on duty.

## **Aim**

The aim is to detect the COVID-19 target genes (ORF1ab and N gene) qualitatively against the Standards provided in the Fluorescence qPCR kit.

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## Material and equipment required

- Rotor Gene Q Instrument
- Rotor-Gene Q chamber
- Rotor Gene Q holder
- Laptop
- UPS
- PPE (Annex I)
- Disinfectant (Sodium hypochlorite and Ethanol)
- Tips
- Pipettes
- Reaction plates
- PCR strips
- PCR Amplification kits
- Adsorbent paper
- PPE (BSL2/BSL3 level working)
- Discarder (waste box)

**Assay Setup Procedure:** Assay set up for COVID-19 Amplification must be carried out in Biosafety Cabinet (BSC II) in the designated room (labelled as Amplification Unit COVID-19). Please see the BSCII SOPs for further information.

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**Reference:**

<https://www.asu.edu/ehs/documents/asu-bsc-sop.pdf>.

[www.geneodx.com](http://www.geneodx.com)

**ANNEXURE**

**Annexure I**

**PPE**

- Eye protection (Safety glasses)
- Disposable nitrile gloves
- Lab coat/tyvek (Disposable)
- Closed shoes with shoe cover
- Respirators may be required for some procedures. The nature of the Personal Protective Equipment (PPE) worn may vary according to the findings obtained from the biological risk assessment.