

## **CERTIFICATE OF ANALYSIS**

This certificate is provided to confirm that the diagnostic kit specified below conforms to the production and testing specifications required by Cepheid's Quality System, in compliance with the US Food and Drug Administration's Quality System Requirements, ISO 13485, European IVD Directive and the Canadian Devices Regulations.

Directive and the Canad		on's Quality System Requirements, ISO 13485, Euro ices Regulations.	opean IVD
<b>Product Name:</b>	Хp	ert® Xpress CoV-2 plus	
Cepheid Catalogue Par	rt No.:	XP3SARS-COV2-10	
<b>Kit Lot No.:</b> 10014	154045		
Cartridge Lot No.:	16618		
<b>Kit Expiration Date:</b>	202	5-10-19	
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 U  Functional Testing	SA	Manufacturing Facility Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden	le ( Newark ( ) Lod
Test Description		Acceptance Criteria	Test Result
Negative		SARS-CoV-2 NEGATIVE	Passed
Positive		SARS-CoV-2 POSITIVE	Passed
Sara Mustafa Abdu Sara Mustafa Abdulla (Oct 28, 2024 09:29 GN	<u>lla</u>	s produced electronically and valid without a wet sig	nature.
Signature of Quality A	Assuran	ce, Date	

Name: Sara Mustafa Abdulla

Title: QA Analyst