

## **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		ices Regulations.	85, European IVD
Product Name:	Хp	ert® Xpress CoV-2 plus	
Cepheid Catalogue Par	rt No.:	XP3SARS-COV2-10	
<b>Kit Lot No.:</b> 10014	157788		
Cartridge Lot No.:	16709		
<b>Kit Expiration Date:</b>	202	5-11-02	
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 U	SA	Manufacturing Facility Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden	Sunnyvale Newark Lodi
Functional Testing  Test Description		Acceptance Criteria	Test Result
Negative		SARS-CoV-2 NEGATIVE	Passed
Positive		SARS-CoV-2 POSITIVE	Passed
If checked this doc  Samaneh Vahid  Samaneh Vahid Gharavi (Nov 18, 20)	Ghara	s produced electronically and valid without a	wet signature.
Signature of Quality A	Assuran	ice, Date	

Name: Samaneh Vahid

Title: Senior QA Analyst