

## **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.

	lian Devices Regulations.	
Product Name:	Xpert® Xpress CoV-2 plus	
Cepheid Catalogue Par	rt No.: XP3SARS-COV2-10	
<b>Kit Lot No.:</b> 10014	448342	
Cartridge Lot No.:	16614	
Kit Expiration Date:	2025-09-07	
<b>Legal Manufacturer</b> Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 U	Manufacturing Facility  Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden	)Sunnyvale ( )Newark ( )L
Functional Testing	Sweden	
Functional Testing  Test Description	Acceptance Criteria	Test Result
-	T	Test Result Passed
•	Acceptance Criteria	

Date

Name: Lava Maroof

Signature of Quality Assurance,

Title: QA Analyst