

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.

Product Name:	Xpert® Xpress CoV-2 plus
Cepheid Catalogue Par	rt No.: XP3SARS-COV2-10
Kit Lot No.: 10014	458933
Cartridge Lot No.:	16715
Kit Expiration Date:	2025-11-16
Legal Manufacturer	Manufacturing Facility Solna Sunnyvale Newark Lodi

<u>Legal Manufacturer</u>	Manufacturing Facility	Solna	Sunnyvale Newark Lodi
Cepheid	Cepheid AB	Ŭ	0 0 0
904 Caribbean Drive	Röntgenvägen 5		
Sunnyvale, CA 94089 USA	SE-171 54 Solna		
•	Sweden		

Functional Testing

Test Description	Acceptance Criteria	Test Result
Negative	SARS-CoV-2 NEGATIVE	Passed
Positive	SARS-CoV-2 POSITIVE	Passed

If checked this document is produced electronically and valid without a wet signature.

Anton Engstrom

Signature of Quality Assurance,

Date

Name: Anton Andersson Engström

Title: QA Analyst