



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 and ISO 13485.

Product Name: Xpert[®] Xpress CoV-2 *plus*

Instructions for Use (IFU)

Catalogue Part No.: XPRS-COV2-10

Part No. and Rev.:

302-8997 Rev. B

Kit Lot No.: 1001455097

Cartridge Lot No.: 21102

Kit Expiration Date: 2025 11 02

Legal Manufacturer

Cepheid
904 Caribbean Drive
Sunnyvale, CA
94089 USA

Manufacturing Facility

Cepheid
121 N Guild Avenue
Lodi, CA 95240
USA

Solna
 Lodi

Sunnyvale

Functional Testing according to D48538, Rev. J

<i>Test Description</i>	<i>Acceptance Criteria</i>	<i>Test Result</i>
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.

10 NOV 2024

Signature of Quality Assurance,

Date

Name: Nancy Inthavisak

Title: Quality Systems Specialist