



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/Flu/RSV *plus*

Instructions for Use (IFU)

Catalogue Part No.: XPRS4PLEX-10

Part No. and Rev.:

302-8057 Rev A

Kit Lot No.: 1001451800

Cartridge Lot No.: 62024

Kit Expiration Date: 2025 10 05

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 D47377, Rev. L

| <i>Test Description</i> | <i>Acceptance Criteria</i> | <i>Test Result</i> |
|-------------------------|--|--------------------|
| Negative | SARS-CoV-2 NEGATIVE;Flu A NEGATIVE;Flu B NEGATIVE;RSV NEGATIVE | Passed |
| Positive | SARS-CoV-2 POSITIVE;Flu A POSITIVE;Flu B POSITIVE;RSV POSITIVE | Passed |

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

10/19/24

Signature of Quality Assurance,

Date

Name: Mark Magno Shelor

Title: Quality Assurance Specialist