

## **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                            |
| <b>Kit Lot No.:</b> 10014 | 451793   |
| Cartridge Lot No.:        | 16703  |
| Kit Expiration Date:      | 2025-10-12   |
| Legal Manufacturer        | Manufacturing Facility Solna Sunnyvale Newark Lodi |

| <u>Legal Manufacturer</u> | Manufacturing Facility () | Solna 🔘 Sunnyvale ( | Newark OLodi |
|---------------------------|---------------------------|---------------------|--------------|
| Cepheid                   | Cepheid AB                | <b>U</b>            | 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.

Sara Mustafa Abdulla

Signature of Quality Assurance,

Date

Name: Sara Mustafa Abdull

Title: QA Analyst