

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.

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Xpert® Xpress CoV-2/Flu/RSV plus

Cepheid Catalogue Part No.:

XP3COV2/FLU/RSV-10

Kit Lot No.: 1001449526

Cartridge Lot No.: 30121

Kit Expiration Date: 2025-09-21

Legal Manufacturer

Manufacturing Facility

Solna

Newark

Sunnyvale

Lodi IVD (B2)

Cepheid AB

Cepheid 904 Caribbean Drive

Röntgenvägen 5 SE-171 54 Solna

Sunnyvale, CA 94089

Sweden

USA

Functional Testing

Test Description	Acceptance Criteria	Test Result Passed
Negative	SARS-CoV-2 NEGATIVE;Flu A NEGATIVE;Flu B NEGATIVE;RSV NEGATIVE	
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.

Signature of Quality Assurance,

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Name: Mustafa Didehvar

Title: QA Analyst