

CERTIFICATE OF ANALYSIS

IVD Directive and the Canadian Devices Regulations. the US Food and Drug Administration's Quality System Requirements, ISO 13485, European production and testing specifications required by Cepheid's Quality System, in compliance with This certificate is provided to confirm that the diagnostic kit specified below conforms to the

Product Name: Xpert® Xpress CoV-2/Flu/RSV plus

Cepheid Catalogue Part No.: XP3COV2/FLU/RSV-10

Kit Lot No.: 1001457628

Cartridge Lot No.: 63615

Kit Expiration Date: 2025 11 16

Legal Manufacturer

904 Caribbean Drive Cepheid

Sunnyvale, CA 94089

Cepheid Manufacturing Facility

)Solna)Newark

Sunnyvale
Lodi IVD (B2)

Lodi, CA 95240 121 N Guild Avenue

Functional Testing

Test Description	Acceptance Criteria	Test Result
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.

Signature of Quality Assurance,

Name: Camille Torres

HOW / LA/III

Title: Quality Systems Specialist