

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.

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/Flu/RSV plus				
Cepheid Catalogue Part No.: XP3COV2/FLU/RSV-10				
Kit Lot No.: 1000541872 Cartridge Lot No.: 21811				
Kit Expiration Date: 2023-10-08				
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA Functional Testing	Manufacturing Facility Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA		mnyvale di IVD (B2)	
Test Description	Acceptance Crite	eria	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.

1	nethornall	280CT WW
Signa	ture of Quality Assurance,	Date
Name	: Ma Stacybelle Peralta	
Title	Quality Systems Specialist	