

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.

IVD Directive and th	e Callaulali	Devices Regula	mons.			
Product Name:	Xpert® X	press CoV-2/F	lu/RSV plus	S		
Cepheid Catalogue	Part No.:	XP3COV2	2/FLU/RSV-	-10		
Kit Lot No.: 10014	44791					
Cartridge Lot No.:	19902					
Kit Expiration Date	2025-08-	31				
Legal Manufacture Cepheid 904 Caribbean Driv Sunnyvale, CA 940 USA	e e	Manufacturin Cepheid AB Röntgenväge SE-171 54 S Sweden	en 5	Solna Newark	_	nnyvale di IVD (B2)
Functional Testing						
Test Description		A	Acceptance Crit	teria		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 doc Lava Maroof Lava Maroof (Oct 1, 2024 16:03 GMT+2)		oduced electron	nically and va	alid without a we	et signatu	ire.

Lava Maroof (Oct 1, 2024 16:03 GMT+2)	
Signature of Quality Assurance,	Date

Name: Lava Maroof

Title: QA Analyst