

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.

<u> </u>	nadian Devices Regulations.	uirements, 180	13485, 1	European
Product Name: Xpe	ert® Xpress CoV-2/Flu/RSV plus			
Cepheid Catalogue Part	No.: XP3COV2/FLU/RSV-	10		
Kit Lot No.: 1001448997				
Cartridge Lot No.: 30107				
Kit Expiration Date: 202	5-09-14			
Legal Manufacturer	Manufacturing Facility	Solna		ınnyvale
Cepheid	Cepheid AB	Newark Lodi IVD (B		
904 Caribbean Drive	Röntgenvägen 5			
Sunnyvale, CA 94089	SE-171 54 Solna			
USA	Sweden			
Functional Testing				
Test Description	Acceptance Criteria			Test Result
	SARS-CoV-2 NEGATIVE:Flu A NEGATIVE:Flu B			

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	

If checked this document is produced electronically and valid without a wet signature.

2024-09-23

Signature of Quality Assurance, Date

Name: Alexander Avramidis

Title: QA Analyst