

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 Can	adian Devices Regulations.				
Product Name: Xpe	rt® Xpress CoV-2/Flu/RSV plus				
Cepheid Catalogue Part	No.: XP3COV2/FLU/RSV-	10			
Kit Lot No.: 1000416170					
Cartridge Lot No.: 04332	Assisting Assistance				
Kit Expiration Date: 202	3-05-21 ************************************				
Legal Manufacturer	Manufacturing Facility	OSolna		nnyvale	
Cepheid	Cepheid	Newark	di IVD (B2)		
904 Caribbean Drive	904 Caribbean Drive				
Sunnyvale, CA 94089	Sunnyvale, CA 94089				
USA	USA				
Functional Testing					
Test Description	Acceptance Criteria			Test Result	
Negative	SARS-CoV-2 NEGATIVE;Flu A NEGATIVE;Flu B NEGATIVE;RSV NEGATIVE			Passed	

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.

Signature of Quality Assurance, Date

Name: Emaela S. Quan-Tong

Title: Quality Assurance Supervisor