

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 Adn IVD Directive and the Can	ninistration's Quality System Requadian Devices Regulations.	airements, ISO 1	3485, E	European	
Product Name: Xpert® Xpress CoV-2/Flu/RSV plus					
Cepheid Catalogue Part	No.: XP3COV2/FLU/RSV-	10			
Kit Lot No.: 1000571800					
Cartridge Lot No.: 21826					
Kit Expiration Date: 2023-11-05					
Legal Manufacturer	Manufacturing Facility	OSolna		nnyvale	
Cepheid	Cepheid	Newark	OLoo	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		3	Passed	
Positive	SARS-CoV-2 POSITIVE;Flu A POSITIVE;Flu B POSITIVE;RSV POSITIVE			Passed	

Positive	SARS-CoV-2 POSITIVE;Flu A POSITIVE;Flu B POSITIVE;RSV POSITIVE	
☐ If checked this documen	t is produced electronically and valid without a wet signature.	
Signature of Quality Assu		

Name: Molly Doan

Title: Quality Systems Specialist