

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.

TVD Directive and the Can	adian Bevices Regulations.		
Product Name: Xpert® Xpress CoV-2/Flu/RSV plus			
Cepheid Catalogue Part No.: XP3COV2/FLU/RSV-10			
Kit Lot No.: 1001459033			
Cartridge Lot No.: 64401			
Kit Expiration Date: 2029	5 11 23		
Legal Manufacturer	Manufacturing Facility		innyvale
Cepheid	Cepheid	Newark OLo	odi IVD (B2)
904 Caribbean Drive	121 N Guild Avenue		
Sunnyvale, CA 94089	Lodi, CA 95240		
USA	USA	•	
Functional Testing			
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		Passed
☐ If checked this documen	t is produced electronically and va	lid without a wet signat	ure.
10-	12/06/2024		
Signature of Quality Assurance, Date			
Name: Jennifer Nguyen	v - 11 1 1 11 v 1 1 1 1 1 1 1 1 1 1		
Title: Quality Systems Spe	ecialist, 1945, 1944		į