

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 and ISO 13485.

Product Name: Xpert [®] Xpress CoV-2/Flu/RSV <i>plus</i>		Instructions for Use (IFU) Part No. and Rev.:		
Catalogue Part No.: XPRS4PLEX-10		302-8057 ▼ Rev A		
Kit Lot No.: 1001462582		302-8055 Rev	02-8055 Rev C	
Cartridge Lot No.: 64007				
Kit Expiration Date: 2025 1	2 07			
<u>Legal Manufacturer</u> Cepheid	<u>Manufacturing Facility</u> Cepheid	Solna	Sunnyvale	
904 Caribbean Drive	121 N Guild Avenue	Lodi		
Sunnyvale, CA	Lodi, CA 95240			
94089 USA	USA			
Functional Testing according Test Description	g to D47377, Rev. L Acceptance Criteria	1	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 document	is produced electronically and valid with	out a wet signature.		
Me	12(18)	124		
Signature of Quality Assurance, Date				
Name: Mark Magno Shelor	· .			
Title: Quality Assurance Sp	pecialist			