

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 plus		Instructions for Use (IFU)	
		Part No. and Rev.: 302-8997	
Kit Lot No.: 1001445752			
Cartridge Lot No.: 09910			
Kit Expiration Date: 2025-0	8-24		
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA Functional Testing according	Manufacturing Facility Cepheid 121 N Guild Avenue Lodi, CA 95240 USA	Solna Sunnyval Lodi	e
Test Description	Acceptance Criter	ria Test Result	
Negative	SARS-CoV-2 NEGA	TIVE Passed	
Positive	SARS-CoV-2 POSIT	ΓΙVE Passed	
☐ If checked this document	is produced electronically and valid wit	thout a wet signature.	
Sorah Bogaco)	ol sept 2021		
Signature of Quality Assurance, Date		e	
Name: Sarah Bagasol			
Title: QualityAssurance Spe	ecialist		