

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.

Directive and the Canad		s Regulations.	pean IVD
Product Name:	Xpert	® Xpress CoV-2 plus	
Cepheid Catalogue Par	rt No.:	KP3SARS-COV2-10	¥
Kit Lot No.: 10005	521437		
Cartridge Lot No.:	'01909		
Kit Expiration Date:	2023-0	6-25	
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 U	SA St	anufacturing Facility Solna Sunnyval epheid 4 Caribbean Drive annyvale, CA 94089 SA	e Newark OLodi
Test Description		Acceptance Criteria	Test Result
Negative		SARS-CoV-2 NEGATIVE	Passed
Positive		SARS-CoV-2 POSITIVE	Passed
If checked this document is produced electronically and valid without a wet signature. 12/22/2022			
Signature of Quality Assurance,		Date	

Name: ThuyTien Nguyen

Title: Quality Systems Specialist