

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.

US Food and Drug Administ Directive and the Canadian	stration's Quality System Requirements, ISO 13485, Euro Devices Regulations.	ppean IVD
Product Name:	Xpert® Xpress CoV-2 plus	
Cepheid Catalogue Part N	No.: XP3SARS-COV2-10	
Kit Lot No.: 1001485	010	
Cartridge Lot No.: 17	204	
Kit Expiration Date:	2026-05-17	
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA Functional Testing	Manufacturing Facility Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden	le Newark Loc
Test Description	Acceptance Criteria	Test Result
Negative	SARS-CoV-2 NEGATIVE	Passed
Positive	SARS-CoV-2 POSITIVE	Passed
If checked this documed Mustafa Didehv Mustafa Didehvar (May 29, 2025 08:		nature.

© 2022 Cepheid. All rights reserved.

Title:

Name: Mustafa Didehvar

QA Analyst