

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 Cana	adian Devices Regulations.		
Product Name: Xper	rt® Xpress CoV-2/Flu/RSV plus		
Cepheid Catalogue Part	No.: XP3COV2/FLU/RSV-	10	
Kit Lot No.: 1000459319	4. 网络克勒克		
Cartridge Lot No.: 08718	TOTAL NEEDS		
Kit Expiration Date: 2023	3 07 09		
Legal Manufacturer	Manufacturing Facility	OSolna •	Sunnyvale
Cepheid	Cepheid	ONewark C)Lodi IVD (B2)
904 Caribbean Drive	904 Caribbean Drive		
Sunnyvale, CA 94089	Sunnyvale, CA 94089		
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 sig	gnature.
- ASZ	27 m 22		
Signature of Quality Assu	rance, Date		
Name: Donna A.	Samson		
Title: Quality System	ns Specialist		