

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 Medical Device Regulations (CMDR).

Directive and the Co	madian Medical Devi	ice regulations (CIVIDIC).			
Product Name: Xpe	ert® Factor II & Factor	V			
Cepheid Catalogue Pa	art No.: GXFIIFV-10	0			
Kit Lot No.: 1001	127293				
Cartridge Lot No.: 09	303				
Kit Expiration Date:	2025-07-27				
Legal Manufacturer		Manufacturing Facility	Solna) Sunnyvale	
Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA		Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 US.	Cepheid		
Functional Testing					
Test Description		Acceptance Criteria		Test Result	
Normal (wild-type)	FII normal; FV normal			Passed	
Homozygous Mutant	FII homozygous; FV homozygous			Passed	
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Tanja Reed Tanja Reed (Aug 16, 2023 10:01 PDT)	Aug 16, 2023	
Signature of Quality Assurance	Date	
Name: Tanja Reed		
Title: Quality Systems Specialist		