

## **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 Canadian Medical Device Regulations (CMDR).					
Product Name: Xpert® Factor II & Factor V					
Cepheid Catalogue Part No.: GXFIIFV-10					
Kit Lot No.: 1000	860788				
Cartridge Lot No.: 09003					
Kit Expiration Date: 2025-03-30					
Legal Manufacturer		anufacturing Facility	○ Solna	Sunnyvale	
Cepheid	Ce	epheid			
904 Caribbean Drive		4 Caribbean Drive			
Sunnyvale, CA 94089 U	SA Su	innyvale, CA 94089 USA			
Functional Testing					
Test Description	Acceptance Criteria		Test Result		
Normal (wild-type)	FII normal; FV normal		Passed		
Homozygous Mutant	FII homozygous; FV homozygous			Passed	

NEAD	antos	04/19/2023
Signatu	re of Quality Assurance	Date
Name:	Zenny Santos	
Title:	Quality Systems Specialist	t