

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	508131				
Cartridge Lot No.: 08	702				
Kit Expiration Date:	2024-09-01				
Legal Manufacturer			Manufacturing Facility	○ Solna	Sunnyvale
Cepheid 904 Caribbean Drive			Cepheid 904 Caribbean Drive		
Sunnyvale, CA 94089 USA		Sunnyvale, CA 94089 USA			
Functional Testing					
Test Description	Acceptance Criteria			Test Result	
Normal (wild-type)	FII normal; FV normal			Passed	
Homozygous Mutant	Mutant FII homozygous; FV homozygous				Passed
				3	

Signature of Quality Assurance

Name: Rhoziel Cusi

Title: Quality Systems Specialist