

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).

Product Name: Xpert [®] Factor II & Factor V		
Cepheid Catalogue Part No.: GXFIIFV-10		¥
Kit Lot No.: 1000169191		
Cartridge Lot No.: 07502		
Kit Expiration Date: 2021-02-14		E *
Legal Manufacturer	Manufacturing Facility	Os

Cepheid

904 Caribbean Drive Sunnyvale, CA 94089 USA Manufacturing Facility Solna Sunnyvale

Cepheid
904 Caribbean Drive
Symptotic CA 04080 US A

Sunnyvale, 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

Signature of Quality Assurance	Sept 3	0, 2019 Date
Name: Dilys Fung	7	
Title: Quality Assurance Supe	rvisor]