



## 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 Devices Regulations (CMDR).

**Product Name:** Xpert® Factor II & Factor V

**Cepheid Catalogue Part No.:** GXFIIFV-10

**Kit Lot No.:** 1001465277

**Cartridge Lot No.:** 20102

**Kit Expiration Date:** 2026 12 27

**Legal Manufacturer**

Cepheid  
904 Caribbean Drive  
Sunnyvale, CA 94089 USA

**Manufacturing Facility**

Cepheid  
121 N Guild Avenue  
Lodi, CA 95240  
USA

Solna

Sunnyvale

Lodi

***Functional Testing according to D16900, Rev. T***

<i>Test Description</i>	<i>Acceptance Criteria</i>	<i>Test Result</i>
Normal (wild-type)	FII normal; FV normal	Passed
Homozygous Mutant	FII homozygous; FV homozygous	Passed

If checked, this document is produced electronically and therefore valid without a wet signature

  
\_\_\_\_\_  
Signature of Quality Assurance,

  
\_\_\_\_\_  
Date

**Name:** Asia Taylor Torres

**Title:** Quality Assurance Specialist