



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.: 1001411079

Cartridge Lot No.: 09501

Kit Expiration Date: 2026-02-15

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,

3/1/24
Date

Name: Jessica Ratzlaff

Title: Quality System Specialist