

## **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® HIV-1 Qual XC

Cepheid Catalogue Part No.:GXHIV-QA-XC-CE-10

**Kit Lot No.:** 1001463825

Cartridge Lot No.: 14804

**Kit Expiration Date:** 2026-06-21

<u>Legal Manufacturer</u> <u>Manufacturing Facility</u> • Solna Sunnyvale

Cepheid ABCepheid ABRöntgenvägen 5Röntgenvägen 5SE-17154 SolnaSE-171 54 Solna

Sweden Sweden

## Functional Testing according to D36985 Rev: C

Test Description	Acceptance Criteria	Test Result
HIV-1 Qual XC PC Low	HIV-1 DETECTED	Passed
Negative	HIV-1 NOT DETECTED	Passed

✓ If checked this document is produced electronically and valid without a wet signature.

Signature of Quality Assurance,	Date
Mustafa Didehvar (Jan 6, 2025 09:28 GMT+1)	
Mustafa Didehvar	

Name: Mustafa Didehvar

Title: QA Analyst