

## 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.:** 1001463824

**Cartridge Lot No.:** 14803

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

**Legal Manufacturer**

Cepheid AB  
Röntgenvägen 5  
SE-17154 Solna  
Sweden

**Manufacturing Facility**

Solna  Sunnyvale

Cepheid AB  
Röntgenvägen 5  
SE-171 54 Solna  
Sweden

***Functional Testing according to D36985 Rev: C***

<i>Test Description</i>	<i>Acceptance Criteria</i>	<i>Test Result</i>
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.

*Lava Maroof* 2025 01 01

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**Signature of Quality Assurance,**

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**Date**

**Name:** Lava Maroof

**Title:** QA Analyst