

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).

Prod	luct	Name:

Xpert® HIV-1 Qual XC

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

Kit Lot No.:

1001454058

Cartridge Lot No.:

14202

Kit Expiration Date:

2026-04-19

Legal Manufacturer

Manufacturing Facility

Solna Sunnyvale

Cepheid AB

Sweden

Röntgenvägen 5

Cepheid AB

SE-17154 Solna

Röntgenvägen 5 SE-171 54 Solna

Sweden

Functional Testing according to D36985 Rev: B

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,

Name: Robert Fiedler

Title:

QA Analyst