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® BCR-ABL Ultra

Cepheid Catalogue Part No.: GXBCRABL-10

Kit Lot No.: 1001463944

Cartridge Lot No.: 43001

Kit Expiration Date: 2026-02-22

Legal Manufacturer
Cepheid
904 Caribbean Drive
Sunnyvale, CA 94089 USA

Manufacturing Facility
Cepheid
904 Caribbean Drive
Sunnyvale, CA 94089
USA

○ Solna ◉ Sunnyvale

○ Lodi

Functional Testing according to D16624, Rev. R.1

Test Description	*Acceptance Criteria*	*Test Result*
Negative	NEGATIVE [Sufficient ABL transcript]	Passed
~0.01% (IS)	POSITIVE	Passed
~0.1% (IS)	POSITIVE	Passed

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

Molly Doan
Molly Doan (Jan 9, 2025 09:33 PST)

Jan 9, 2025

Signature of Quality Assurance **Date**

Name: Molly Doan

Title: Quality Systems Specialist

301-9243 Rev B_ C of A Xpert BCR-ABL Ultra

Final Audit Report 2025-01-09

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"301-9243 Rev B_ C of A Xpert BCR-ABL Ultra" History

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