

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, the Canadian Medical Devices Regulations (CMDR), and the China National Medical Products Administration's Quality System Requirements.

Product Name: Xpert® MTB/RIF Ultra					
Cepheid Catalogue Part No.: GXMTB/RII	F-ULTRA-50				
Kit Lot No.: 1001470366					
Cartridge Lot No.: 74809					
Kit Expiration Date: 2026-07-26					
Legal Manufacturer Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden	Manufacturing Facility Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden	Solna Lodi	Sunnyval		
Functional Testing according to D25862 Rev AP					

Test Description	Acceptance Criteria	Test Result
Wild Type Control	MTB DETECTED VERY LOW; Rif Resistance NOT DETECTED or MTB DETECTED LOW; Rif Resistance NOT DETECTED or MTB DETECTED MEDIUM; Rif Resistance NOT DETECTED or MTB DETECTED HIGH; Rif Resistance NOT DETECTED	Passed
Mutant Control	MTB DETECTED VERY LOW;RIF Resistance DETECTED or MTB DETECTED LOW;RIF Resistance DETECTED or MTB DETECTED MEDIUM;RIF Resistance DETECTED or MTB DETECTED HIGH;RIF Resistance DETECTED	Passed
Negative	MTB NOT DETECTED	Passed

If checked, this document is produced electronically and therefore valid without a wet signature Sara Mustafa Abdulla

ra Mustafa Abd	ulla (Feb 6, 2025 14:08 GMT+1)	
Signature of Quality Assurance,		Date
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Name:	Sara Mustafa Abdulla	
Title	QA Analyst	
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