

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/RIF	F-ULTRA-50		
Kit Lot No.: 1001460212			
Cartridge Lot No.: 73908			
Kit Expiration Date: 2026-05-24			
Legal Manufacturer Cepheid AB	Manufacturing Facility Cepheid AB		
Röntgenvägen 5	Röntgenvägen 5	Solna	Sunnyvale
SE-17154 Solna	SE-171 54 Solna Sweden	O 7 11	
Sweden	Owodon	Lodi	

Functional Testing according to D25862, Rev. AN

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 Samaneh Vahid Gharavi

Saman	eh Vahid Gharavi (Dec 10, 2024 13:40 GMT+1)	
Signature of Quality Assurance,		Date
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Name:	Samaneh Vahid	
Title:	Senior QA Analyst	
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