

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-10		
Kit Lot No.: 1001464973			
Cartridge Lot No.: 74404			
Kit Expiration Date: 2026-06-28			
Legal Manufacturer	Manufacturing Facility		
Cepheid AB	Cepheid AB	Solna	Sunnyvale
Röntgenvägen 5	Röntgenvägen 5	Some	O Builly vale
SE-17154 Solna	SE-171 54 Solna	_	
Sweden	Sweden	Lodi	

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

Signature of Quality Assurance,

Date

Name: Anton Andersson Engström

Title: QA Analyst