

Product Name: Xpert® MTB/RIF Ultra

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 Device Regulations (CMDR).

Cepheid Catalogue Part No.: GXMTB/RIF-ULTRA-10				
Kit Lot No.: 1000148478				
Cartridge Lot No.: 13405				
Kit Expiration Date: 2020-02-16				
Legal Manufacturer		Manufacturing Facility		Sunnyvale
Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden		Cepheid AB Röntgenvägen 5 SE-171 54 Solna Sweden		
Functional Testing				
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 NO	T DETECTED		Passed
Signature of Quality Assurance Date Name: Kambiz Ghaderi				
Title: QA Analyst P/N 301-7659, Rev. B				