

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).

	Regulations (CMDR).	ective and the Cana-
Product Name: Xpe	rt® MTB/RIF Ultra	
Cepheid Catalogue F	art No.: GXMTB/RIF-ULTRA-50	
Kit Lot No.: 100115	55419	
Cartridge Lot No.:	45506	
Kit Expiration Date:	2025-02-09	
Legal Manufacture Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden	Manufacturing Facility Cepheid 121 N Guild Avenue Lodi, CA 95240 USA	Solna Sunnyvale Lodi
Functional Testing Test Description	according to D25862, Rev. AM 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 HEIGH; Rif Resistance NOT DETECTED or MTB DETECTED HIGH; Rif Resistance NOT DETECTED	Passed
Mutant Control	MTB DETECTED VERY LOW;RIF Resistance DETECTED MTB DETECTED LOW;RIF Resistance DETECTED of MTB DETECTED MEDIUM;RIF Resistance DETECTED or MTB DETECTED HIGH;RIF Resistance DETECTED	Passed
Negative	MTB NOT DETECTED	Passed
If checked, this do	pocument is produced electronically and therefore valid without a 29AU62023 y Assurance, Date	wet signature
Name: Chong Mou	a	
Title: Quality Systems Specialist I		