

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).

dian Medical Devices	Regulations (CMDR).		
Product Name: Xper	rt® MTB/RIF Ultra		
Cepheid Catalogue P	art No.: GXMTB/RIF-ULTRA-50		
Kit Lot No.: 100073	1191		
Cartridge Lot No.: 4	14522		
Kit Expiration Date:	2024 07 28		
Legal Manufacture Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden Functional Testing	Manufacturing Facility Cepheid 121 N Guild Avenue Lodi, CA 95240 USA according to D25862, Rev.) Solna	unnyval
Test Description	Acceptance Criteria	Test Result	
Wild Type Control	MTB DETECTED VERY LOW; Rif Resistance NOT DETECTED of MTB DETECTED LOW; Rif Resistance NOT DETECTED or MTB DETECTED MEDIUM; Rif Resistance NOT DETECTED or MTB DETECTED HEIGH; 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 Name: Trinh Nguyo	y Assurance, 2/15/23 Date	a wet signature	