

Title:

Quality Systems Specialist

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		nents, 180 13483, European 1	IVD Direc	ctive and the	Cana-
Product Name: Xpe	ert® MRSA/SA Blood Cu	ulture			
Cepheid Catalogue P	art No.:GXMRSA/SABC	C-CE-10			
Kit Lot No.: 100142	1936				
Cartridge Lot No.: 2	21002				
Kit Expiration Date:	2026-01-18				
Legal Manufacture Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden		Manufacturing Facility Dephelo 04 Caribbean Drive Sunnyvale, CA 94089 ISA	<!--</th--><th>Solna Lodi</th><th>Sunnyvale</th>	Solna Lodi	Sunnyvale
Functional Testing	according to D36872, K	Rev. H			
Functional Testing of Test Description		Rev. H		Test Resu	lt
	Acce			Test Resu Passed	lt
Test Description	Acce MRSA Posi	eptance Criteria			elt
Test Description Positive Negative	Acce MRSA Posi MRSA Neg	eptance Criteria itive; SA Positive	without a	Passed Passed	

302-0555 Rev B CofA MRSASABC-CE

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