

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).

	Quality System Requirements Regulations (CMDR).	s, ISO 13485, European	i IVD Dire	ective and the Ca	ana-
Product Name: Xpo	ert® MRSA/SA Blood Culture				
Cepheid Catalogue P	art No.:GXMRSA/SABC-CE	E-10			
Kit Lot No.: 100146	51387				
Cartridge Lot No.: 2	22201				
Kit Expiration Date:	2026-10-25				
Legal Manufacture Cepheid AB Röntgenvägen 5 SE-17154 Solna Sweden	904 (	<u>Ifacturing Facility</u> eld Caribbean Drive yvale, CA 94089	0	Solna (	Sunnyvalo
	according to D36872, Rev.	Н			
	ū	H ace Criteria		Test Result	
Functional Testing	ū	ice Criteria		Test Result Passed	
Functional Testing  Test Description	Acceptan	sa Positive			
Test Description  Positive Negative	Acceptant MRSA Positive; MRSA Negative ocument is produced electronic	SA Positive ; SA Negative	l without a	Passed Passed	

## 302-0555 Rev B CofA MRSASABC-CE

Final Audit Report 2024-12-22

Created: 2024-12-22

By: Molly Doan (molly.doan@cepheid.com)

Status: Signed

Transaction ID: CBJCHBCAABAACMkR80VJotJn9rjzU4qxYfFG2i2OOdqk

## "302-0555 Rev B CofA MRSASABC-CE" History

Document created by Molly Doan (molly.doan@cepheid.com) 2024-12-22 - 8:12:07 PM GMT

Document emailed to Molly Doan (molly.doan@cepheid.com) for signature 2024-12-22 - 8:12:34 PM GMT

Document e-signed by Molly Doan (molly.doan@cepheid.com)
Signature Date: 2024-12-22 - 8:12:43 PM GMT - Time Source: server

Agreement completed. 2024-12-22 - 8:12:43 PM GMT

