

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).

Drug Administration's Quality Systems Medical Devices Regulations	tem Requirements, ISO 13485, Europea (CMDR).	n IVD Directive and the Cana-
Product Name: Xpert® SA Nasa	al Complete	
Cepheid Catalogue Part No.: GX	KSACOMP-10	
Kit Lot No.: 1001459095		
Cartridge Lot No.: 29103		
Kit Expiration Date: 2026-11-15		
Legal Manufacturer Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA Functional Testing according to	Manufacturing Facility Cepheid 904 Caribbean Drive Sunnyvale, CA 94089 USA D D 36872 Rev. H	Solna Sunnyval Lodi
Test Description	Acceptance Criteria	Test Result
Positive	MRSA Positive; SA Positive	Passed
Negative	MRSA Negative; SA Negative	Passed
Molly Doan	oduced electronically and therefore vali Dec 9, 2024	d without a wet signature
Molly Doan (Dec 9, 2024 12:27 PST) Signature of Quality Assurance.	·	_
Signature of Quartey rissurance,	Date	

Name: Molly Doan

Title: Quality Systems Specialist

301-6217 Rev B SACOMP

Final Audit Report 2024-12-09

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

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