



## 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).

**Product Name:** Xpert® Xpress Strep A

**Cepheid Catalogue Part No.:** XPRSTREPA-10

**Kit Lot No.:** 1001471366

**Cartridge Lot No.:** 41811

**Kit Expiration Date:** 2026-01-25

**Legal Manufacturer**

Cepheid  
904 Caribbean Drive  
Sunnyvale, CA 94089  
USA

**Manufacturing Facility**

Cepheid  
121 N Guild Avenue  
Lodi, CA 95240  
USA

Solna

Sunnyvale

Lodi

**Functional Testing according to D27089, Rev. R**

<i>Test Description</i>	<i>Acceptance Criteria</i>	<i>Test Result</i>
Negative	Strep A NOT DETECTED	Passed
LOW Positive	Strep A DETECTED	Passed
HIGH Positive	Strep A DETECTED	Passed

If checked, this document is produced electronically and therefore valid without a wet signature

2/16/25

\_\_\_\_\_  
Signature of Quality Assurance,

\_\_\_\_\_  
Date

**Name:** Mark Magno Shelor

**Title:** Quality Assurance Specialist