



## 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, the Canadian Medical Devices Regulations (CMDR), and the China National Medical Products Administration's Quality System Requirements.

**Product Name:** Xpert® Carba-R

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

**Kit Lot No.:** 1001459074

**Cartridge Lot No.:** 24102

**Kit Expiration Date:** 2026 05 17

**Legal Manufacturer**

Cepheid  
904 Caribbean Drive  
Sunnyvale, CA 94089 USA

**Manufacturing Facility**

Cepheid  
121 N Guild Avenue  
Lodi, CA 95240  
USA



Solna



Sunnyvale

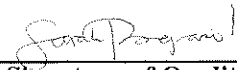


Lodi

*Functional Testing according to D18272, Rev. AF*

<i>Test Description</i>	<i>Acceptance Criteria</i>	<i>Test Result</i>
Low Positive	IMP DETECTED, VIM DETECTED, NDM DETECTED, KPC DETECTED, OXA48 DETECTED	Passed
High Positive	IMP DETECTED, VIM DETECTED, NDM DETECTED, KPC DETECTED, OXA48 DETECTED	Passed
Negative	IMP NOT DETECTED, VIM NOT DETECTED, NDM NOT DETECTED, KPC NOT DETECTED, OXA48 NOT DETECTED	Passed

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


01DEC 2024  


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**Signature of Quality Assurance,**
**Date**

**Name:** Sarah Bagasol

**Title:** Quality Assurance Specialist