



America

CERTIFICATE

No. QS6 024497 0031 Rev. 03

Certificate Holder: **NONIN MEDICAL, INC.**
13700 1st Avenue North
Plymouth MN 55441-5443
USA

Certification Mark:



Scope of Certificate: **Design, Development, Manufacturing, Servicing and Distribution of Oximeters, Pulse Oximeters, Cerebral Oximeters, Breathing Monitors and Non-Sterile Sensors**

Standard(s): **ISO 13485:2016**

Regulatory Authority(ies): **Australia TGA, Brazil ANVISA, Health Canada, USA FDA, MHLW / PMDA. See attached for listing of specific regulatory requirements.**

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. Validity of this certificate can be obtained by visiting the website www.tuvsud.com/ps-cert
TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID: **F001301**

Effective Date: **2023-01-21**

Expiry Date: **2026-01-20**

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Date of Issue: 2023-02-17

(Renee Walker)
Director, US Certification Body, MHS



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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002
 - Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil

- RDC ANVISA n. 665/2022 - Good Manufacturing Practices
 - RDC ANVISA n. 551/2021
 - RDC ANVISA n. 67/2009 - Vigilance

Canada

- Medical Device Regulations – Part 1- SOR 98/282

Japan

- MHLW Ministerial Ordinance No. 169 (2004), as amended by MHLW Ministerial Ordinance No.60 (2021)
 - Japan PMD Act (as applicable)

United States

- 21 CFR Part 803
 - 21 CFR Part 806
 - 21 CFR Part 807 – Subparts A to D
 - 21 CFR Part 820

Facility(ies):

NONIN MEDICAL, INC.
 13700 1st Avenue North, Plymouth MN 55441-5443, USA

Facility Scopes:

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