Number: 6078824CE01

EU Quality Assurance Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex XI Part A

Manufacturer:

NINGBO RANOR MEDICAL SCIENCE & TECHNOLOGY CO., LTD.

NO.127 Fenghui Road, Wangchun Industrial Park, Haishu District
315100 Ningbo, Zhejiang
China
SRN ID.: CN-MF-000011211

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EU- Regulation which apply to them:

0344

Supplement to certificate: 6078827CN

Authorized Representative:

MedPath GmbH
Mies-van-der-Rohe-Strasse 8
80807 Munich, Germany

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant requirements of EU Regulation 2017/745, including all subsequent amendments for the above mentioned conformity assessment. The manufacturer/ authorized representative is subject to periodic surveillance as required for the applicable conformity assessment in accordance to Regulation 2017/745.

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Principal Certification Manager

First Issued: **14 December 2021** Date: **14 December 2021** Expiry date**: 1 December 2026**

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
T +31 88 96 83000 www.dekra-product-safety.com Company registration 09085396

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EU Quality Assurance Certificate

Conformity Assessment Regulation 2017/745 on Medical devices, Annex XI Part A

This certificate covers the following device(s) / groups of device(s):

Active non-implantable devices for monitoring of vital physiological parameters (MDA0203, class IIa)
- Infrared thermometers - Blood pressure monitors

Conditions for or limitations to the validity of this certificate:

- N/A

Certificate History

Identification of the Common Specifications and Harmonized Standards complied with are documented within the technical documentation and audit assessments carried out. These are traceable through the DEKRA Certification B.V. Certification Notice. The Certification Notice also identifies the necessary information related to the quality management system of the manufacturer, including facilities.

Date of Issue certificate	Certification Notice Reference	Action
14-12-2021	6078827CN02	first issue

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