



EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III
(Class IIa and Class IIb Devices)

No. G10 004665 0011 Rev. 01

Manufacturer:

Shenzhen Finicare Co., Ltd

201 and 301 of Building A22
Building A22 and A23
No.4 Industrial Park, Tantou Community, Songgang Street
Bao'an District
518105 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

SRN Manufacturer - CN-MF-000024364

Authorized Representative:

Riomavix S. L.
Calle de Almansa 55, 1D, 28039 Madrid, SPAIN

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s). The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.

The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.

The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples. All applicable requirements of the Testing, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with.

For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10 004665 0011 Rev. 01

Report No.: GZ25036501

Preceding Certificate No.: G10 004665 0011 Rev. 00

Valid from: 2025-12-29

Valid until: 2028-10-16

Date of Initial Issuance: 2023-10-17

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2025-12-29



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Classification: Class IIa
Device Group: Z1203020302 - NON-INVASIVE BLOOD PRESSURE
MONITORING INSTRUMENTS

Intended Purpose: -

Classification: Class IIa
Device Group: V030101 - THERMOMETERS
Intended Purpose: -

The validity of this certificate depends on conditions and/or is limited to the following: - none -

Revision History:

Rev.	Dated	Report	Description
00	2023-10-17	GZ2336501	Initial issuance
01	2025-12-29	GZ25036501	Amended: Change of certificate holder's data