





## **EU Quality Assurance Certificate (MDR)**

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex XI Part A (Class I Devices in sterile condition, with measuring function or reusable surgical instruments)

No. G21 115419 0002 Rev. 00

Manufacturer: **WUXI EXANOVO MEDICAL INSTRUMENT** 

CO.,LTD.

C2-Liandong U Gu, Xibei Town

Xishan District

214194 Wuxi City, Jiangsu

PEOPLE'S REPUBLIC OF CHINA

SRN Manufacturer - CN-MF-000021396

Lotus NL B.V. **Authorized** 

Koningin Julianaplein 10, 1e Verd, 2595AA The Hague, THE Representative:

**NETHERLANDS** 

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 XI Part A of this regulation with a positive result.

As applicable the involvement of the notified body is limited to the aspects relating to:

- establishing, securing and maintaining sterile conditions,
- conformity of the devices with the metrological requirements,
- reuse of the device, in particular cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use.

The certified quality assurance system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/pscert?q=cert:G21 115419 0002 Rev. 00

Report No.: SH22190701

Valid from: 2023-11-02 Valid until: 2028-11-01

Christoph Dicks

Issue date: 2023-11-02 Head of Certification/Notified Body



**Product Service** 

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No. G21 115419 0002 Rev. 00

Classification: Class I

V03010104 - LIQUID EXPANSION THERMOMETERS **Device Group:** 

**Device Properties:** MDS 1010 - Devices with a measuring function

The validity of this certificate depends on conditions and/or is limited to the following:

-none-

**Revision History:** 

Rev. Dated Description Report 00 2023-11-02 SH22190701 Initial issuance