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EC CERTIFICATION

QUALITY MANAGEMENT SYSTEM CERTIFICATE

EU Regulation 2017/745 for Medical Devices, Annex IX Chapters I & III

We hereby declare that a conformity assessment based on a quality management system and technical documentation restricted to the aspects of manufacture concerned with the conformity of the devices with sterility requirements - has been carried out following the requirements of EU Regulation 2017/745 for Medical Devices.

We certify that the documentation conforms to the relevant provisions of the aforementioned regulation, and the result entitles the organization to use the CE 2862 marking on the products listed below.

Certificate Number:
28620139275

Initial Certification Date:
5 January 2023

Date of Certification Decision:
5 January 2023

Certificate Issue Date:
5 January 2023

Certificate Expiry Date:
28 November 2027

Medico Technology Co., Ltd

Room 201, 301 and 401 Building A No.10, Bao long 5th Road, Tongle Community, Baolong Street, Longgang District, Shenzhen, Guangdong, China

Manufacturer SRN: CN-MF-000013067

Authorised Representative Name
Luxus Lebenswelt GmbH
Kochstr.1, 47877, Willich, Germany

Scope:

Class I sterile devices

Brian Mather
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Intertek Medical Notified Body AB is a Notified Body in accordance with the requirements set out in EU Regulation 2017/745 on medical devices, with the identification number 2862.