ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-099

Product Service

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 088855 0016 Rev. 00

Manufacturer:

Globalcare Medical Technology Co., Ltd.
7th Building
39 Middle Industrial Main Road
European Industrial Zone, Xiaolan Town
528415 Zhongshan City, Guangdong Province
PEOPLE'S REPUBLIC OF CHINA

SRN Manufacturer: Not available at issuance date of this certificate

Authorized Representative:

Donawa Lifescience Consulting Srl
Piazza Albania, 10, 00153 Rome, ITALY

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 and certification regulation of TÜV SÜD Group have to be complied with.
For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10 088855 0016 Rev. 00

Report No.: GZ2019503

Valid from: 2021-10-29
Valid until: 2026-10-28

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2021-10-29

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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No. G10 088855 0016 Rev. 00

Classification:	IIa
Device Group:	Z12159002 - AEROSOL GENERATORS
Intended Purpose:	/
Classification:	IIa
Device Group:	Z12030205 - NON INVASIVE BLOOD PRESSURE MONITORS
Intended Purpose:	/
Classification:	IIa
Device Group:	Z12080303 - BREAST PUMPS
Intended Purpose:	/
Classification:	IIa
Device Group:	Z12040299 - GENERAL MEDICINE INSTRUMENTS FOR THERAPEUTIC TREATMENTS - OTHERS
Intended Purpose:	/
Classification:	IIa
Device Group:	V03010102 - ELECTRONIC THERMOMETERS AND END CAPPS
Intended Purpose:	/
Classification:	IIa
Device Group:	Z12050403 - ECG HOLTER RECORDERS
Intended Purpose:	/
Classification:	IIa
Device Group:	Z12030202 - MULTI-PARAMETER PATIENT MONITORS
Intended Purpose:	/
The validity of this certificate depends on conditions and/or is limited to the following:	-none-

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