Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 088855 0010 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

Facility(ies):

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

Product Category(ies):

Aerosoltherapy Nebulizers, Breast Pump, Blood Pressure Measuring Equipment, Bright Light Therapy Lamp

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: GZ19195EXT01

Valid from: 2019-08-14
Valid until: 2024-05-26

Date, 2019-08-14

Stefan Preiß
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany