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

TÜV SÜD Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 067303 0017 Rev. 00

Manufacturer:

Shenzhen Med-link Electronics Tech Co., LTD.
4th and 5th Floor, Building Two
Hualian Industrial Zone
Xinshi Community, Dalang Street, Longhua District
518109 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Pulse Oximeter Probes, Temperature Probes, Non-sterilized Disposable Neutral Electrodes , Temp-Pulse Oximeter, Digital Blood Pressure Monitors, Digital Infrared Thermometers, ECG Recorders, ETCO2 Mainstream and Sidestream Sensors, Micro Capnometer.**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: GZ1913601

Valid from: 2020-04-01
Valid until: 2023-10-23

Date, 2020-04-01

Christoph Dicks
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

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