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 052126 0043 Rev. 03

Manufacturer:

TaiDoc Technology Corporation
B1-7F, No. 127, Wugong 2nd Road, Wugu Dist.
24888 New Taipei City
TAIWAN

Product Category(ies): **Blood Glucose Plus Blood Pressure Monitoring System, Blood Glucose Plus Blood Pressure Meter, Thermometer, Blood Pressure Meter, Blood Pressure Monitoring System, Vital Signs Monitor, Pulse Oximeter, Nebulizer, Lancing Device With Sterile Blood Lancet, Sterile Blood Lancet, ECG Recorder, SpO2 Sensor, Temperature Monitor, Electronic Nasal Aspirator, Electric Breast Pump and Manual Breast Pump.**

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.: TW2001103

Valid from: 2020-03-23
Valid until: 2024-05-26

Date, 2020-03-23

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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