



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 088955 0007 Rev. 01

Manufacturer: **Shenzhen Caremed Medical**

Technology Co.,Ltd.

East side, 3/F, C building

Kelunte Low-carbon industries Park Gaofeng community, Dalang offices

Longhua district 518109 Shenzhen

PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): SpO2 sensors, Temperature probes

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.: GZ1919201

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

Date. 2020-03-23

Christoph Dicks Head of Certification/Notified Body