RTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 08 88955 001

Manufacturer: **Shenzhen Caremed Medical Technology Co.,Ltd.**
Zone B, 3/F, 11 Building, Hebei
Industrial Area, Longhua office
Longhua New District
518109 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

EC-Representative: **C. K. Medical Int`l**
Laan van Cattenbroeck 10
3703 BM Zeist
THE NETHERLANDS

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.: 748485364

Valid from: 2015-02-10
Valid until: 2020-02-09

Date, 2015-02-13

Hans-Heiner Junker

616784

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

4.11