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

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 076260 0010 Rev. 00

Manufacturer: **Shenyang Canta Medical Tech. Co., Ltd.**
No.76-39 Shenbei Road
Daoyi Economic Development Zone
Shenbei New District
110136 Shenyang
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Oxygen Concentrator for Medical Use, Sleep Apnoea Breathing Therapy Devices.**

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.: BJ19777071

Valid from: 2020-03-02
Valid until: 2024-05-26

Date, 2020-03-02

Christoph Dicks
Head of Certification/Notified Body

Digitally signed by Matei Vasile
Date: 2021.03.15 14:04:48 EET
Reason: MoldSign Signature
Location: Moldova

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

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