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 104589 0002 Rev. 00**

**Manufacturer:** **Guangdong Ecan Medical Co., Ltd.**
Building 1, No. 222, Xindu Road, Chengjiao Street
Conghua District
510920 Guangzhou City, Guangdong Province
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):** **Urethral catheters, Laryngeal Mask Devices, Endotracheal Tubes, Tracheostomy Tubes, Endobronchial Tubes, Feeding Tubes, Suction Catheters, Suction tubing, PVC Stomach Tubes, Oxygen Masks, Tubing Set for Nebulizers**

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.:** GZ1942901

Digitally signed by Guțan Ghenadie
Date: 2022.11.12 20:33:10 EET
Reason: MoldSign Signature
Location: Moldova

**Valid from:** 2019-11-26
**Valid until:** 2024-05-26

**Date,** 2019-11-26

Christoph Dicks
Head of Certification/Notified Body

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT


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**Facility(ies):**

Guangdong Ecan Medical Co., Ltd.
Building 1, No. 222, Xindu Road, Chengjiao Street, Conghua District, 510920 Guangzhou City, Guangdong Province, PEOPLE'S REPUBLIC OF CHINA

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