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Product Service

# EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

**No. G2S 104589 0003 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):**

**Nasopharyngeal Airways,**
**Oropharyngeal Airways**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** GZ1942901

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

Digitally signed by Guțan Ghenadie
Date: 2021.01.10 19:03:38 MSK
Reason: MoldSign Signature
Location: Moldova

**Date,** 2019-11-26

Christoph Dicks
Head of Certification/Notified Body


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

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**Facility(ies):**

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**Parameter:**


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