



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

Production Quality Assurance System
 Directive 93/42/EEC on Medical Devices (MDD), Annex V
 (Devices in Class IIa, IIb or III)

No. G2 080946 0007 Rev. 03

Manufacturer: **Anji SPENQ Industrial Co., Ltd.**

F16, Building C
 Anji Chamber of Commerce Mansion
 No. 99 Tianhuangping South Road
 313300 Anji County, Zhejiang Province
 PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Anji SPENQ Industrial Co., Ltd.
 F16, Building C, Anji Chamber of Commerce Mansion, No. 99
 Tianhuangping South Road, 313300 Anji County, Zhejiang
 Province, PEOPLE'S REPUBLIC OF CHINA

**Product
 Category(ies):**

**Latex Foley Catheters, Oxygen Masks, Sterile Blood
 Lancets, Sterile Latex Surgical Gloves, Sterile Syringes
 for Single Use, Sterile Infusion Sets for Single Use,
 Sterile Intravenous Needles for Single Use, Sterile
 Hypodermic Needles for Single Use, Sterile Blood
 Transfusion Sets for Single Use, Nasal Oxygen
 Cannulae, Suction Catheters, Stomach Tubes, Feeding
 Tubes, Nelaton Catheter, Disposable Surgical Blades,
 Endotracheal Tubes, Laryngeal Mask, Reinforced
 Endotracheal Tube, Mucus Extractor, Tracheostomy
 Tube, Silicone Foley Catheter**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

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Valid from: 2020-01-29

Valid until: 2024-05-26

Date, 2020-01-29

Christoph Dicks
 Head of Certification/Notified Body