



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. 01**

**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

**EC-Representative:**

SUNGO Europe B.V.  
Olympisch Stadion 24, 1076DE Amsterdam, THE  
NETHERLANDS

**Product  
Category(ies):**

**Latex Foley Catheters, Oxygen Masks, Sterile Blood  
Lancets, Sterile Latex Surgical Gloves, Digital  
Thermometers, Blood Pressure Monitors, 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  
Cannulaes, 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.

**Report No.:** SH1860109

**Valid from:** 2018-12-03

**Valid until:** 2020-11-23

**Date,** 2018-12-03

Stefan Preiß

TÜV SÜD  
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認 證 證 書 ◆ CERTIFICADO ◆ CERTIFICAT



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