



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 073283 0045 Rev. 00

Manufacturer: **Ningbo Greetmed Medical Instruments Co., Ltd.**
16F-1, Building 1
No. 98 Chuangyuan Road, Hi-Tech Zone
315042 Ningbo, Zhejiang Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **All Silicone Foley Catheter,
Blood Bags,
Electrosurgical Pencils,
Condom,
Tracheostomy Tubes,
Urethral Catheters,
Pulse Oximeter,
Non-absorbable Surgery Suture,
Sterile Hydrocolloid Dressings**

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.: SH1929915
Valid from: 2019-08-27
Valid until: 2021-04-17

Date, 2019-08-27

Stefan Preiß
Head of Certification/Notified Body

TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT



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Facility(ies):

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