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 095166 0013 Rev. 01

Manufacturer:

NINGBO TRUSTLAB INSTRUMENTS CO., LTD
18-7, No.1 Building, Wante Business Centre
Hi-tech zone
315042 Ningbo
PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

NINGBO TRUSTLAB INSTRUMENTS CO., LTD
18-7, No.1 Building, Wante Business Centre, Hi-tech zone,
315042 Ningbo, PEOPLE'S REPUBLIC OF CHINA

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

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.: SH19104404

Valid from: 2019-07-15
Valid until: 2021-07-20

Date, 2019-07-15

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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