



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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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 I in sterile conditions, sterilised systems or procedure packs)

No. G2S 095166 0017 Rev. 02

Manufacturer

**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):**

**-General non-active, non-implantable medical
devices
-Non-active devices for anaesthesia, emergency
and intensive care
-Bandages and wound dressings
(For detailed information please see attachment)**

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.: SH191044EXT01

Valid from: 2020-03-06

Valid until: 2024-05-26

Date, 2020-03-06

Christoph Dicks
Head of Certification/Notified Body

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

