



## **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

NINGBO TRUSTLAB INSTRUMENTS CO., LTD Facility(ies):

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.

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Valid from: 2019-07-15

Valid until: 2021-07-20

2019-07-15

1. Pumil

Stefan Preiß

Head of Certification/Notified Body

Date,