



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 IIa, IIb or III)

No. G2 073283 0046 Rev. 01

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):**

**Non-active devices for anaesthesia, emergency
and intensive care
Non-active devices for injection, infusion,
transfusion and dialysis
Non-active instruments
Bandages and wound dressings
Respiratory devices, devices including hyperbaric
chambers for oxygen therapy, inhalation
anaesthesia
Monitoring devices of vital physiological
parameters
Medical Gloves
(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 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 IIIa and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: SH19299EXT01

Valid from: 2020-03-16
Valid until: 2024-05-26

Date, 2020-03-16

Christoph Dicks
Head of Certification/Notified Body

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

NINGBO GREETMED MEDICAL INSTRUMENTS CO., LTD.

