

# 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 IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.:

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**Valid from:**

2020-03-16

Valid until:

2024-05-26

Date, 2020-03-16

C.D.H

Christoph Dicks  
Head of Certification/Notified Body