



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 16 08 74618 006

**Manufacturer:**

**Jiangyin Jinfeng Medical  
Equipment Co., Ltd.**

No. 74 Zhoujia Village, Jinfeng Village  
Xuxiake Town  
214407 Jiangyin  
PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:**

**ZOUSTECH S.L.**

Pso. Castellana, 141 – Planta 19  
28046 Madrid  
SPAIN

**Product  
Category(ies):**

**Sterile Hypodermic Syringes For Single Use(with Needles),  
Infusion Sets For Single Use(with Needles),  
Sterile Hypodermic Needle For Single Use,  
Sterile Insulin Syringes For Single Use**

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.:**

SH1662201

**Valid from:**

2017-02-07

**Valid until:**

2022-02-06

**Date,** 2017-02-07

Stefan Preiß

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

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**Facility(ies):**

Jiangyin Jinfeng Medical Equipment Co., Ltd.  
No. 74 Zhoujia Village, Jinfeng Village, Xuxiake  
Town, 214407 Jiangyin, PEOPLE'S REPUBLIC OF  
CHINA