



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

Page 1 of 2



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Facility(ies):

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