ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 17 01 72857 008

Manufacturer:

Changzhou Jiafeng Medical Equipment Co., Ltd.
Zhongjiang Village, Sanhekou
Zhenglu Town, Wujin District
213115 Changzhou City, Jiangsu Province
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies):

Infusion Sets for Single Use (with Needle),
Sterile Hypodermic Syringes for Single Use (with Needle),
Disposable Sterile Needles

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.: SH1760308

Valid from: 2017-11-22
Valid until: 2020-05-25

Date, 2017-11-22

Stefan Preiß

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

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Facility(ies):

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Province, PEOPLE'S REPUBLIC OF CHINA

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