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

Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 16 02 39452 025

Manufacturer:

Jiangsu Kangjin Medical Instrument Co., Ltd.
Zhenglu Town
213111 Changzhou
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

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

Product Category(ies):

Infusion Sets, Transfusion Sets,
Syringes, Scalp Vein Sets,
Injection Needles,
Blood Collection Needles for Single Use,
Light-resistant Infusion Sets for Single Use,
The Connection / Extension Tube for Single Use,
Biopsy Forceps for Single Use,
Infusion Sets with Precision Filters for Single Use,
Parenteral Nutrient Infusion Sets for Single Use,
Enteral Feeding Set for Single Use,
Grasping Forceps for Single Use,
Electrosurgical Snares for Single Use

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: SH16085EXT01

Valid from: 2016-03-17
Valid until: 2021-03-13

Date, 2016-03-17

Stefan Preiß

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

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Facility(ies):

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TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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