Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 081681 0021 Rev. 00

Manufacturer:

Shanghai Kindly Medical Instruments Co., Ltd.
No.925 Jinyuan yi Road
201803 Shanghai
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Sterile Seldinger needle, Intervention accessories kit, Guidewire, Introducer tool kit, Introducer set, Bone cement syringe, Manifold kit, Stopcocks, Manifold, Infusion pumps, Pressure transducer for single use, Extension tube, Foley catheter, Safety syringe, Angiography Catheter, Y-connector Pack, Sterile Infusion connector and accessory for single use, Guiding Catheter, Micro Catheter.**

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.: BJ19703071

Valid from: 2019-10-29
Valid until: 2024-05-26

Date, 2019-10-29

Christoph Dicks
Head of Certification/Notified Body

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

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Facility(ies):

Shanghai Kindly Medical Instruments Co., Ltd.
No.925 Jinyuan yi Road, 201803 Shanghai, PEOPLE'S REPUBLIC OF CHINA

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