



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 039452 0033 Rev. 01**

**Manufacturer:**

**Jiangsu Kangjin Medical  
Instrument Co., Ltd.**

Zhenglu Town  
213111 Changzhou  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):** Infusion Sets, Transfusion Sets, Syringes, Scalp Vein Sets, Injection Needles, Light-resistant Infusion Sets 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 (Infusion Pump Type), Grasping Forceps for Single Use, Electrosurgical Snares for Single Use, Hot Biopsy Forceps for Single Use, Light-resistant Infusion Sets with Precision Filters for Single Use, Light-resistant Scalp Vein Sets for Single Use, Endoscopic Injection Needle for Single Use, Washing Pipe for Single Use, Endoscopic Hemoclip 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.:**

SH19085EXT01

**Valid from:**

2020-01-09

**Valid until:**

2024-05-26

**Date,**

2020-01-09

Christoph Dicks  
Head of Certification/Notified Body



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