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Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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ZLG-BS-244.10.08

Product Service

# EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

**No. G2S 081681 0017 Rev. 02**

**Manufacturer**

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

**Product Category(ies):**

**Sterile 'Inflation device, Pressure bandage, Angiography syringe' for single use, Infusion sets without needle, syringes without needle, Burette infusion sets without needle, Vaginal Dilator.**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. 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

Digitally signed by Perminov Andrei
Date: 2021.12.21 09:30:59 EET
Reason: MoldSign Signature
Location: Moldova


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

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**Facility(ies):**

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