



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 041808 0054 Rev. 05

Manufacturer

**Shanghai Kindly Enterprise
Development Group Co., Ltd.**

No 658 Gaochao Road
201803 Shanghai
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile 'Urinal Bag, Irrigation Syringe, Medical Tube (Rectal Tube), Irrigation Sets, Umbilical cord clamp, Protective cap, Transfer spikes, Oral syringes, Enteral syringes, Infusion connector, Syringes without needle, Infusion sets without needle, Transfusion sets without needle, Infusion Sets with Burette without needle, safe syringes without needle, Syringes with re-use prevention feature without needle, Drawing-up filter straw, Blunt Filter Needle, Zero residual small volume syringe without needle, Neuraxial fit tip syringe' for Single Use.

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G2S 041808 0054 Rev. 05](http://www.tuvsud.com/ps-cert?q=cert:G2S_041808_0054_Rev.05)

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Christoph Dicks
Head of Certification/Notified Body