



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 036336 0054 Rev. 03

Manufacturer:

**Zhejiang Kindly Medical
Devices Co., Ltd.**

No.758, 5th Binhai Road
Binhai Industrial Park, Longwan District
325025 Wenzhou, Zhejiang Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

**Disposable Needles, Scalp Vein Sets, Blood-Collecting
Needles, Huber Needles, Fistula Needles, Anaesthesia
Needles, Dental Needles for Single Use, Sterile I.V. catheter
for single use, Disposable Insulin Pen Needle, Sterile Biopsy
Needles for single use, Sterile Percutaneous Vertebroplasty
Kit for single use, Sterile Irrigation Needles for Single Use,
Safety Needles, Safety Scalp Vein Sets, Safety Blood-
Collecting Needles, Safety I.V. Catheter for Single Use, Safety
Fistula Needles, Luer Adapter, Safety Blood Lancet, Syringes,
Infusion Sets, Transfusion Sets, Burette-Type Infusion Sets,
Sterile Intravascular Catheter Introducer for Single Use,
Sterile Syringes for Insulin for Single Use, Sterile Disinfecting
Cap 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. 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:G10363360054Rev.03

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Christoph Dicks
Head of Certification/Notified Body