



Product Service

EC-CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 09 08 36336 030

Manufacturer:



Zhejiang Kindly Medical Devices Co., Ltd.

No.252, Yongqiang Road
Yongzhong, Longwan
325024 Wenzhou, Zhejiang province
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies):

Syringes, Infusion Sets, Transfusion Sets, Burette-Type Infusion Sets, Sterile Intravascular Catheter Introducer 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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: BJ981204

Valid until: 2019-12-27

Hans-Heiner Junker

Date, 2014-12-27



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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