



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 14 09 82107 006

Manufacturer: **Zhejiang Runqiang Medical Instruments Co., Ltd**
No. 618 Dade Road, Jiaxing Xiuzhou District
314031 Jiaxing City
PEOPLE'S REPUBLIC OF CHINA

EC-Representative: **Shanghai International Holding Corp. GmbH (Europe)**
Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies): **Disposable infusion pump, Disposable anaesthesia air filter and Medical filters, Disposable breathing Tubes Intended for Use with Anaesthesia apparatus and Ventilator, Disposable Urethral Catheterization Kit, Disposable Sterile Urethral Catheter.**

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.: BJ1479404

Valid from: 2014-12-15

Valid until: 2018-05-06



Hans-Heiner Junker

Date, 2014-12-16

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

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Facility(ies):

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