



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 18 04 03601 002

Manufacturer:

Zhejiang MDKingdom Technology Co., Ltd.

508, 5th Floor, 28 Cang Ling Road
Huzhen Town, Jinyun County
321404 Lishui City, Zhejiang Province
PEOPLE'S REPUBLIC OF CHINA



EC-Representative:

LAIPUDE GmbH

Schwarzenberger Str.46
47226 Duisburg
GERMANY

Product Category(ies):

**Enteral Feeding Pump,
Infusion Pump, Syringe Pump.**

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.:

BJ1722501

Valid from:

2018-07-12

Valid until:

2023-07-11



Date, 2018-07-12

Stefan Preiß

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

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