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Zentralstelle der Länder
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bei Arzneimitteln und
Medizinprodukten
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ZLG-BS-244.10.08



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 101172 0003 Rev. 01

Manufacturer: **Changsha DeepMed
Medical Technology Co., Ltd.**
Room 709 - 717, Building 1
Haiping Park
No. 229, Guyuan Road, Hi-tech Zone
410205 Changsha
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): Infusion Pumps and Syringe Pumps

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:G11011720003Rev.01

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Date, 2020-09-14

Christoph Dicks
Head of Certification/Notified Body