Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz 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 088861 0008 Rev. 03

Manufacturer:

Zibo Qiaosend Medical Articles Co., Ltd
No.2, Gaoyuan East Road
256300 Gaoqing county, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Disposable Infusion Sets, Sterile Syringes, Blood Transfusion Sets, Sterile Hypodermic Needle for Single Use, Intravenous Infusion Needle for Single Use, Extension line for single use, Intravenous Cannula 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. See also notes overleaf.

Report No.: BJ20090601

Valid from: 2020-02-24
Valid until: 2024-01-26

Date, 2020-02-24

Christoph Dicks
Head of Certification/Notified Body

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

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Facility(ies):

Zibo Qiaosend Medical Articles Co., Ltd
No.2, Gaoyuan East Road, 256300 Gaoqing county, Shandong Province, PEOPLE'S REPUBLIC OF CHINA

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