



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V

(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 087117 0010 Rev. 01

Manufacturer

Weigao Medical International Co., Ltd.

No.1 Weigao Road
High-tech Industrial Development Zone
264210 Weihai, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

MedNet GmbH

Borkstrasse 10, 48163 Muenster, GERMANY

Product Category(ies):

Sterile Prefilled Syringe composite (without needle) for single use, Sterile Precision Solution Filters for single use, Sterile Infusion Connector without Needles for single use, Sterile Heparin Cap for single use, Sterile Lachrymal Cannula Needle for single use, Urine Bags, Feeding Set, Oxygen Masks, Sterile pressure connector tube for Single Use, Inflation device.

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:

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2024-04-16

Date,

2019-01-18

Stefan Preiß



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CHINA