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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 094273 0007 Rev. 03

Manufacturer

**Shandong Weigao Group Medical
Polymer Co., Ltd.**

No.18 Xingshan Road
Torch Hi-tech Science Park
264210 Weihai, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile Prefillable Syringe Barrel (Without Needle) for single use, Sterile Precision Solution Filters for single use, Sterile Prefillable Syringe Stopper For Single Use, Sterile Pressure Connector Tube For Single Use, Sterile Dispensing Syringe for Single Use, Sterile Low Resistance Dispensing Syringe for Single Use, Sterile Irradiation-Resistance Dispensing Syringe with Needle for Single Use, Extension Tube for Single Use, Light-resistant Extension Tube for Single Use, Virus Sampling Kit for Single Use, Swab for Single Use.

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. 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:G2S 094273 0007 Rev. 03

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Christoph Dicks
Head of Certification/Notified Body