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

TÜV SÜD 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 003509 0003 Rev. 01

Manufacturer	**Zibo Qray International Co., Ltd.** No.77 Zhanghuan Road 255000 Zibo City, Shandong PEOPLE'S REPUBLIC OF CHINA
EC-Representative:	ZOUSTECH S.L. Pso. Castellana, 141 – Planta 19, 28046 Madrid, SPAIN
Product Category(ies):	**Infusion Connector and Accessory for Single Use, Urine Bags, Sterile Vaginal Dilators for Single Use, Basic Dressing Pack, First Aid Kits, Disposable Surgical Pack**

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.

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Valid from:	2019-08-02
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Digitally signed by Bolea Petru
Date: 2022.09.14 07:14:36 EEST
Reason: MoldSign Signature
Location: Moldova

Date, 2019-08-02

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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Facility(ies):

Zibo Qray International Co., Ltd.
No.77 Zhanghuan Road, 255000 Zibo City, Shandong, PEOPLE'S REPUBLIC OF CHINA

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