ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 18 03 56464 018

Manufacturer: **Jiangsu Webest Medical Product Co., Ltd.**
Yingchun Road, Industrial park
211700 Xuyi, JiangSu
PEOPLE'S REPUBLIC OF CHINA

EC-Representative: **Shanghai International Holding Corp. GmbH (Europe)**
Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies):
Sterile Infusion Sets for Single Use,
Sterile Syringes for Single Use,
Sterile Three-way Stopcocks for Single Use,
Sterile Heparin Caps for Single Use,
Sterile Dental Needles for Single Use,
Scalp Vein Set for Single Use,
I.V. Cannula for Single Use,
Hypodermic Needle for Single Use,
Infusion Set with Burette, Transfusion Set,
Three Way Stopcock and Extension Tube,
Syringe for Insulin,
I.V. Flow Regulator for Single Use,
Solution Administration Set for Infusion Pump

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

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Date, 2018-04-13

Stefan Preiß

04052767931833

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

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Facility(ies):

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Yingchun Road, Industrial park, 211700 Xuyi,
JiangSu, PEOPLE'S REPUBLIC OF CHINA

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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