Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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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 094558 0005 Rev. 01

Digitally signed by Neer Ghennadii
Date: 2024.08.30 11:18:19 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Manufacturer: **Jiangyin Fanmei Medical Device Co., Ltd.**
No.19 Yishan Rd, Lingang Street
214400 Jiangyin
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Infusion Sets with Needles,Intravenous Needles, Safety Syringe, Auto-disable Syringe, Disposable Blood Transfusion Sets, Sterile Surgical Gloves, Feeding Tube, Stomach Tube, Suction Catheter, Nelaton Tube, Nasal Oxygen Cannula, Oxygen Mask, Disposable Blood Collection Needles, Syringe for Insulin, Sterile Syringe for Single Use (with Needle), Sterile Hypodermic Needle 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 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.

Report No.: SH191005EXT01

Valid from: 2020-01-07
Valid until: 2024-05-26

Date, 2020-01-07

Christoph Dicks
Head of Certification/Notified Body

REPUBLICA MOLDOVA IDNO 1003600060028 SRL «BECOR» PENTRU FACTURI INTREPRINDEREA MIXTA mun. CHISINAU

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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