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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 073684 0016 Rev. 02

Manufacturer:

Suzhou TexNet Co., Ltd.

RM1103-1104, 11/F, Block A
Xin Tian Xiang Bldg., 388 Suyu Road, SIP
215021 Suzhou
PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Suzhou TexNet Co., Ltd.
RM1103-1104, 11/F, Block A, Xin Tian Xiang Bldg., 388 Suyu
Road, SIP, 215021 Suzhou, PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Sterile Radiopaque Gauze Swabs, Sterile Radiopaque Lap
Sponges, Sterile Syringes for Single Use, Sterile Three-Way
Stopcocks for Single Use, Sterile Infusion Sets for Single
Use, Sterile Heparin Caps for Single Use, Scalp Vein Set for
Single Use, I.V. Cannula for Single Use, Hypodermic Needles
for Single Use, Disposable Urethral Catheters (Nelaton
Catheter), Disposable Nasal Oxygen Cannula, Oxygen Masks,
Disposable Suction Catheters, Endotracheal Tubes,
Disposable Feeding Tubes, Laryngeal Masks, Foley Catheters
(Latex), Surgical Gloves, Sterile Blood Lancets, Disposable
Surgical Blades, Alcohol Swabs/Pads, Silicone Foley
Catheters, Enteral Feeding Bag Set, Disposable Stomach
Tube

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.:

SH19620EXT01

Valid from:

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Valid until:

2024-05-26

Date,

2019-09-26

S. Preiß
Stefan Preiß
Head of Certification/Notified Body