Benannt durch/Designated by
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 IIa, IIb or III)

No. G2 088705 0011 Rev. 00

Manufacturer:

FarmaSino Pharmaceuticals (Jiangsu) Co., Ltd.
No.100 JianYe Road
210004 Nanjing
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

**Sterile Hypodermic Syringes for Single Use,
Sterile Hypodermic Needles for Single Use,
Intravenous Needles for Single Use,
Disposable Infusion Sets,
Disposable Sterilized Latex Surgical Gloves,
Disposable Surgical Blades (with and without handle),
Blood Pressure Monitor,
Nasal Oxygen Cannula, Endotracheal Tubes,
Reinforced Endotracheal Tube,
Foley Catheter(latex), Transfusion Sets,
Digital Thermometer**

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.: SH19885EXT01

Valid from: 2019-06-11
Valid until: 2024-05-26

Date, 2019-06-11

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):

FarmaSino Pharmaceuticals (Jiangsu) Co., Ltd.
No.100 JianYe Road, 210004 Nanjing, PEOPLE'S REPUBLIC OF CHINA

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