



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 16 07 78455 013

Manufacturer: Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road
Mazhang District
524094 Zhanjiang
PEOPLE'S REPUBLIC OF CHINA



EC-Representative: Shanghai International Holding Corp. GmbH (Europe)

Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies): Foley Catheter (silicone), Nelaton Catheter (latex), Endotracheal Tubes, Anesthesia Kits, Suction Catheter for Single Use in the Respiratory Tract, Disposable Urethral Catheterization Set, Urethral Catheters (PVC), Stomach Tube (PVC), Reinforced Endotracheal 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.: SH16090EXT01

Valid from: 2021-09-17

Valid until: 2026-09-18

Date, 2021-09-17

Stefan Preiß



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

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Zhanjiang, PEOPLE'S REPUBLIC OF CHINA****Zhanjiang Star Enterprise Co., Ltd.
No. 49 Jinchuan Road, 524094 Zhanjiang,
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