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 069211 0023 Rev. 02

Manufacturer:

Hangzhou Bever Medical Devices Co., Ltd.
Building 2, No. 1-1, Houmuqiao
Yongle Village, Cangqian Street
Yuhang District
311121 Hangzhou
PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Hangzhou Bever Medical Devices Co., Ltd.
Building 2, No. 1-1, Houmuqiao, Yongle Village, Cangqian Street, Yuhang District, 311121 Hangzhou, PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Nelaton Catheters, Tracheal Tubes, Suction Catheters, Hydrophilic Urinary Catheter Kits, Reinforced Endotracheal Tubes, Hydrophilic Urinary Catheters, Tiemann Catheters, Feeding Tubes, Heat & Moisture Exchangers and Filters, Lubricating Jellies, Catheterization Kits, Hydrophilic Meatal Dilators

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.: SH1954411

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