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

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 094490 0008 Rev. 00

Manufacturer:

VHMED (Nantong) Co., Ltd.
108 Qiantangjiang Road, Tongzhou Bay
226332 Nantong, Jiangsu
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

**Monopolar Electrode Surgical Instrument,
Bipolar Electrode Surgical Instrument,
Hemostatic Ligation Clips, Veress Needle,
Suction and Irrigation Tube Set,Trocar,
Specimen Retrieval Bag, Wound Protection Retractor,
Insufflation Tube, Port Closure Device,
Laparoscopic Instrument, Surgical Procedure Pack**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G1 094490 0008 Rev. 00

Report No.: SH201038EXT01

Valid from: 2021-03-18
Valid until: 2024-05-26

Date, 2021-03-18

Christoph Dicks
Head of Certification/Notified Body

Digitally signed by Iarovoi Petru
Date: 2022.11.23 11:37:28 EET
Reason: MoldSign Signature
Location: Moldova

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany