Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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 095178 0004 Rev. 01

Manufacturer: **Baisheng Medical Co.,Ltd.**
No.11 Fusheng Road, Xinhui District
529100 Jiangmen City, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Electrosurgical generator & Accessories, Electrosurgical pencil, Electrosurgical pads, Electrosurgical electrode, Electrosurgical forceps and Cable, Disposable skin staplers**

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. See also notes overleaf.

Report No.: GZ1925401

Valid from: 2020-02-11
Valid until: 2024-05-26

Date, 2020-02-11

Digitally signed by Ionașcu Andrei
Date: 2022.01.28 10:55:08 EET
Reason: MoldSign Signature
Location: Moldova

Christoph Dicks
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