



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 16 04 95178 002

Manufacturer:

Baisheng Medical Co.,Ltd.

No.11 Fusheng Road, Xinhui District
529100 Jiangmen City, Guangdong
PEOPLE'S REPUBLIC OF CHINA



EC-Representative:

GUIDIER CO.,LTD

KEMP HOUSE 152-160 City Road
London
EC1V 2NX
UNITED KINGDOM

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.:

7484032609

Valid from:

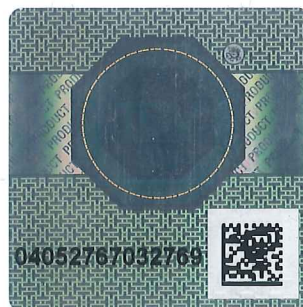
2016-11-08

Valid until:

2021-11-07

Date, 2016-11-08

Stefan Preiß



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

Page 1 of 2



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 16 04 95178 002**Facility(ies):**

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