



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 102891 0002 Rev. 00

Manufacturer:

**Jiangsu Ruitian Medical
Technology Co.,Ltd.**

5 Floor East, Block B3
HuTang Tech Industrial Park, HuTang Town
Wujin
213000 Changzhou
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Disposable Biopsy Forceps,
Disposable Endoscopic Injection Needles,
Disposable Polypectomy Snare**

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.: SH19147201

Valid from: 2020-01-30

Valid until: 2024-05-26

Date, 2020-01-20

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認 證 證 書 ◆ CERTIFICADO ◆ CERTIFICAT

