



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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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 I in sterile conditions, sterilized systems or procedure packs)

No. G1S 102716 0002 Rev. 00

Manufacturer:

**JingFang Precision Medical
Device(Shenzhen) Co., Ltd.**

3/F, Bldg C, Jinwei Yuan
Crossing at Qingsong West Road and Qinglan 2nd Road
Julongshan, Grand Industrial Zone
Longtian Street, Pingshan District
518118 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

**Product
Category(ies):**

**Disposable Needle Guides for endocavity
or general purpose biopsy**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex II. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: GZ1838201

Valid from: 2019-08-16

Valid until: 2024-05-26

Date, 2019-08-16

Stefan Preiß
Head of Certification/Notified Body

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 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFICAT

A4 / 07.17

