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Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
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Product Service

# EC Certificate

Production Quality Assurance System  
Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in Class IIa, IIb or III)

**No. G2 093011 0006 Rev. 01**

**Manufacturer: Ningbo Foyomed Medical Instruments Co., Ltd.**

Room 805-806  
No. 299 of Jiangnan Yipin Garden  
Hi-Tech Zone  
315040 Ningbo  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Sterile Nonactive Medical Devices and Active Medical Devices**  
(for detailed information see following pages)

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

**Report No.:** SH19994EXT01

**Valid from:** 2019-11-26

**Valid until:** 2024-05-26

**Date,** 2019-11-26

Christoph Dicks  
Head of Certification/Notified Body

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