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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 101975 0002 Rev. 00**

**Manufacturer:**

**Zhejiang Longmed Medical Technology Co., Ltd.**
Building 1, No. 926
Changhong east street
Fuxi street
Moganshan national high-tech zone
313200 Deqing, Huzhou, Zhejiang
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):**

**Wound Closure Device,**
**Single Use Fascial Closure System,**
**Single Use Abdominal Trocars and Suit**

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.:** SH18144001

**Valid from:** 2019-08-23
**Valid until:** 2024-05-26

**Date,** 2019-08-23

Stefan Preiß
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

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**Facility(ies):**

**Zhejiang Longmed Medical Technology Co., Ltd.
Building 1, No. 926, Changhong east street, Fuxi street, Moganshan national high-tech zone, 313200 Deqing, Huzhou, Zhejiang, PEOPLE'S REPUBLIC OF CHINA**

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT
TÜV SÜD


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