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Zentralstelle der Länder  
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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 IIa, IIb or III)

**No. G1 038814 0087 Rev. 00**

**Manufacturer:** **Well Lead Medical Co., Ltd.**  
C-4 Jinhu Industrial Estate, Hualong  
511434 Panyu, Guangzhou  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):** **Drainage System, Oxygen Catheter, Silicone Drainage System, Breathing Circuit, Anesthetic Breathing Circuit, Anesthetic Breathing Circuit Kit, Rectal Pressure Catheter, Extraction Bag (Operation Use), Ureteral Stent Set, Urodynamic Catheter, Ureteral Access Sheath, Ureteral Dilation Balloon Catheter, Ureteral Stent, Ureteral Dilator, Urethral Dilator, Urological Guide Wire, Suction-Evacuation Access Sheath, Dilation Set, Stone Retrieval Basket, Ureteral Catheter, Urinary Nephrostomy Catheter**

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.:** SH19080CN01

**Valid from:** 2020-03-31

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

**Date,** 2020-03-31

Christoph Dicks  
Head of Certification/Notified Body

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 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT