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Zentralstelle der Länder  
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bei Arzneimitteln und  
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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 003960 0002 Rev. 00**

**Manufacturer:** **Better Life Medical Technology Co., Ltd.**  
1F(North), Bldg.19, No.8 Jinfeng Rd.  
Suzhou New District  
215163 Suzhou  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):** Better Life Medical Technology Co., Ltd.  
1F(North), Bldg.19, No.8 Jinfeng Rd., Suzhou New District,  
215163 Suzhou, PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Defibrillator Monitor**

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.

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Stefan Preiß