



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 IIa, IIb or III)

**No. G1 104553 0001 Rev. 00**

**Manufacturer:** **Shenzhen Yimi Life Technology Co., Ltd**  
305, Building A, Tengbo Industrial Park  
Changshangjiang Street, Longbei Village  
Pingshan District  
518118 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

### Product Category(ies): Pulse Oximeter

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.:** GZ1940901

**Valid from:** 2020-03-20

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

**Date,** 2020-03-20

Christoph Dicks  
Head of Certification/Notified Body