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Zentralstelle der Länder  
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bei Arzneimitteln und  
Medizinprodukten  
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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)

**G1 049076 0016 Rev. 02**

**Manufacturer:** **Shenzhen Creative Industry Co., Ltd.**  
Floor 5, BLD 9  
BaiWangxin High-Tech Industrial Park  
Songbai Road, Xili Street  
Nanshan District  
518110 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Patient Monitor, Vital Signs Monitor, Fingertip Oximeter, Handheld Pulse Oximeter, Wrist Oximeter, Easy ECG Monitor, Spot-Check Monitor, SpO2 Probe, Sleep Screener, Multi Parameter Monitors for Capnography and Pulse Oximetry**

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.:** GZ1915302

**Valid from:** 2020-01-07  
**Valid until:** 2024-05-26

**Date,** 2020-01-07

Christoph Dicks  
Head of Certification/Notified Body

