







**Innovation, Responsibility**

**Shenzhen MedKe Technology Co., Ltd**

深圳市曼克科技有限公司

Item	P/N	Description	Ref.image
1	T1305	Philips Skin temperature probe,3m,1pcs/bag	
2	P9325A	Philips adult finger sensor,3m,1pcs/bag	
3	G5224S	Philips one-piece ECG cable,5 leads,Snap,IEC,1pcs/bag	
4	040-000545-00	Mindray BeneHeart defibrillator Cable D6/D3 MR6702 040-000545-00	



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
www.zlg.de  
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 085432 0004 Rev. 00**

**Manufacturer:**

**Shenzhen MedKe Technology Co., Ltd**

4/F, Bldg.A1, Anle Ind. Zone, Hangcheng RD., Baoan Dist.  
518126 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):**

Shenzhen MedKe Technology Co., Ltd  
4/F, Bldg.A1, Anle Ind. Zone, Hangcheng RD., Baoan Dist.,  
518126 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): SpO2 sensors**

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.:**

GZ18114EXT01

**Valid from:**

2019-02-28

**Valid until:**

2024-01-16

**Date,**

2019-02-28

Stefan Preiß

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A4 / 07.17



# Certificate

No. Q5 085432 0005 Rev. 01

**Holder of Certificate:** **Shenzhen MedKe Technology Co., Ltd**  
4/F, Bldg.A1, Anle Ind. Zone, Hangcheng RD., Baoan Dist.  
518126 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):** Shenzhen MedKe Technology Co., Ltd  
4/F, Bldg.A1, Anle Ind. Zone, Hangcheng RD., Baoan Dist.,  
518126 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

**Certification Mark:**



**Scope of Certificate:** **Design and Development, Production and Distribution of Medical Compressors and Nebulizers system, Medical accessories (including SpO2 sensors, Temperature probes, Patient cables and Lead wires, BP accessories and Infusors)**

**Applied Standard(s):** EN ISO 13485:2016  
Medical devices - Quality management systems -  
Requirements for regulatory purposes  
(ISO 13485:2016)  
DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

**Report No.:** GZ1911401  
**Valid from:** 2020-02-17  
**Valid until:** 2023-01-16

**Date,** 2020-02-17

Christoph Dicks  
Head of Certification/Notified Body

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