



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
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)

**No. G1 073767 0013 Rev. 00**

**Manufacturer: Shenzhen Aeon Technology Co., Ltd.**

RM6H02, Block 27-29  
Tianxia IC Industrial Park, Majialong  
No.133 of Yiyuan road, Nantou Street  
Nanshan District  
518052 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):**

Shenzhen Aeon Technology Co., Ltd.  
RM6H02, Block 27-29, Tianxia IC Industrial Park, Majialong,  
No.133 of Yiyuan road, Nantou Street, Nanshan District, 518052  
Shenzhen, PEOPLE'S REPUBLIC OF CHINA

Shenzhen Aeon Technology Co., Ltd. Bao'an Branch.  
3/F, Block B, Bldg 6, Industrial Zone of Yusheng, No. 467 of 107  
National Highway, Gushu intersection, Xixiang Street, Bao'an  
District, 518126 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

Dongguan Tianyuan Medical Devices Co., Ltd.  
5/F, Bldg A, No.68 of Junma Road, Xinmalian Village, Dalang  
Town, 523797 Dongguan, PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Fetal Dopplers, Pulse Oximeters,  
Nebulizers and Infrared Thermometers**

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 Preis