Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

TÜV SÜD 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 Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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