



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 078097 0007 Rev. 00**

**Manufacturer:**

**Wuxi Jike Electronics Co., Ltd.**

2nd workshop floor & 3rd office floor of the 4th building  
No.29 Changjiang south road  
New District  
214028 Wuxi, Jiangsu Province  
PEOPLE'S REPUBLIC OF CHINA

**Facility(ies):**

Wuxi Jike Electronics Co., Ltd.  
2nd workshop floor & 3rd office floor of the 4th building, No.29  
Changjiang south road, New District, 214028 Wuxi, Jiangsu  
Province, PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Respiratory Humidifiers**

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.:**

SH19716EXT01

**Valid from:**

2019-10-29

**Valid until:**

2024-05-26

**Date,**

2019-10-29

Christoph Dicks

Head of Certification/Notified Body

ZERTIFIKAT ♦ CERTIFICATE ♦ CERTIFICADO ♦ CERTIFIKAT ♦ СЕРТИФИКАТ ♦ 認證證書

A4 / 07.17