



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

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 071993 0017 Rev. 00

Manufacturer:

**Little Doctor Electronic
(Nantong) Co., Ltd.**

No.8, Tongxing Road
Economic & Technical Development Area
226010 Nantong, Jiangsu
PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Little Doctor Electronic (Nantong) Co., Ltd.
No.8, Tongxing Road, Economic & Technical Development Area,
226010 Nantong, Jiangsu, PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

**Blood Pressure Monitor,
Digital Thermometer, Ultrasonic Nebulizer,
Compressor Nebulizer , Dental Oral Irrigator**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: SH1940401

Valid from: 2020-02-25
Valid until: 2024-05-26

Date, 2020-02-25

Christoph Dicks
Head of Certification/Notified Body



Declaration of Conformity

Manufacturer: Little Doctor Electronic (Nantong) Co., Ltd.
No. 8, Tongxing Road, Economic and Technical Development
Area, 226010 Nantong, Jiangsu, People's Republic of China

Facility: Little Doctor Electronic (Nantong) Co., Ltd.
No. 8, Tongxing Road, Economic and Technical Development
Area, 226010 Nantong, Jiangsu, People's Republic of China

EU Representative: Little Doctor Europe Sp. z o. o.
Zawila Str. 57G, 30-390, Krakow, Poland

Product name: Digital Blood Pressure Monitor

Model number: LD2
Cuff-LDA

Classification (Medical Device Directive 93/42/EEC, Annex IX): Class IIa

UMDNS CODE: 16157

Conformity: Annex V

We herewith declare that the above mentioned product meet the provisions of the following EC Council Directive and Standards. All supporting documentation are retained under the premise of manufacturer and the notify body.

General Applicable Directives: The DIRECTIVE 93/42/EEC on Medical Devices (MDD 93/42/EEC)

Notified Body: TÜV SÜD Product Service GmbH,
Certification Body, Rüdlerstraße 65, 80339 Munich, Germany

Identification No: 0123

EC Certificate: G2 071993 0017 Rev.00

Expiration date of the Certificate: 2024-05-26

Data CE mark was affixed: October, 2005

Place: People's Republic of China

Date: 2020-10-15

Signature: 

Name: GAO JIAWEI
Position: Director

