

No. 8 Tongxing Road P.R. of China Tel.: +86 0513 85986718

## **Declaration of Conformity**

Manufacturer:	Little Doctor Electronic (Nantong) Co.,	Ltd.
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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, LD3, LD3a, LD23, LD23A, LD30, LD51, LD51A, LD51S

Classification (Medical Device Class IIa

Directive 93/42/EEC, Annex

IX):

**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, Ridlerstraße 65, 80339 Munich, Germany

Identification №: 0123

EC Certificate: G2 071993 0017 Rev.00

Expiration date of the Certificate: 2024-05-26

Data CE mark was affixed: October, 2005

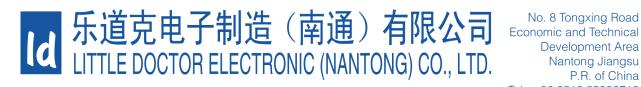
Place: People's Republic of China

Date: 2021-04-07

Signature:

Name:

Position: Director



P.R. of China Tel.: +86 0513 85986718

## Annex 2 of **Declaration of Conformity** for Digital Blood Pressure Monitor

Manufacturer:	Little Doctor Electronic (Nar	ntong) 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: Cuff

Model number: Cuff-LDA

Classification (Medical Device

Directive 93/42/EEC, Annex

IX):

Conformity:

Annex V\*

Class IIa

\*The classification & conformity assessment are applied as a complete Medical Device including its Cuff.

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.

Place: People's Republic of China

Date: 2021-04-07

Signature:

Name:

Director Position: