

## 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  
SRN: CN-MF-000047483

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  
SRN: PL-AR-000042267

Product name: Compressor Nebulizers

Model number: LD-221C

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

Basic UDI-DI: 697368616NEBVV

UMDNS CODE: 12712

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.

Quality Management System (QMS) in accordance with Article 10(9) MDR is in place.

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 No: 0123

EC Certificate: G2 071993 0017 Rev.00  
Validity period: from: 2020-02-25 / until: 2024-05-26

Confirmation Letter: CL 071993 0019 Rev. 00  
Expiration date of the Certificate: 2028-12-31

Data CE mark was affixed: October, 2005

Place: People's Republic of China

Date: 2025-05-01

Signature:

Name: GAO JIAWEN  
Position: Director

