

Title

EC Declaration of Conformity

Date 2025-08-06

Manufacturer: Beurer GmbH (see address in footer)

SRN: DE-MF-000005422

Product category: Nebulizer

Product type: IH 57

Intended use: Nebulisers (including compressor, ultrasonic and mesh nebulisers) are medical

> devices for the nebulisation of liquids and liquid medication. This device produces aerosols by combining an oscillating mesh with holes and a liquid medication. The aerosol treatment is suitable for treating the upper and lower airways. By nebulising and inhaling the medication prescribed/recommended by your doctor, you can prevent diseases affecting the airways, or in the case that you contract such an

illness, you can alleviate symptoms and speed up your recovery.

The product specified above is in conformity with the following specifications.

(EU) 2017/745 Medical device regulation (MDR)

Basic-UDI-DI: 4211125IH57P2

Classification/applied

rule(s):

Class IIa/rule 12 and rule 20

Conformity assessment

procedure:

Annex IX, Chapter I

The notified body mdc medical device certification GmbH, located at Kriegerstr. 6, 70191 Stuttgart, Germany, identification number 0483, issued in the course of the mentioned conformity assessment procedure the following certificate:

Certificate no. and validity: D1311700063, valid to 2026-04-07

2011/65/EU Restrictions of the use of certain hazardous substances in electrical and electronic

equipment (RoHS)

EN IEC 63000:2018

This declaration of conformity is issued under the sole responsibility of the manufacturer.

Signed for and on behalf of: Beurer GmbH

Place, date of issue: Ulm, 2025-08-06

Name, function, signature,

stamp:

Werner Meternek, Director Quality Management & Regulatory Affairs

Beurer GmbH Söflinger Straße 218 + 89077 Ulm

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Beurer Template 410-0040 / rev21 / 2025-08-01