beurer	Title	EU Declaration of Conformity
	Date	2026-11-17

Manufacturer: Beurer GmbH (see address in footer)

SRN: DE-MF-000005422

Product category: Nebulizers

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

Product name (Model/Type)		Basic-UDI-DI	intended use
Compressor IH	IH 15	4211125IH15NJ	Nebulizers (including compressor, ultrasonic, and mesh nebulizers) are medical devices for the nebulization of liquids and liquid medication (aerosols). This device produces aerosols by combining compressed air and liquid medication. The aerosol treatment is suitable for treating the upper and lower airways. By nebulizing 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.
	IH 16	4211125IH16NL	
	IH 18 (Type: IH 18/1)	4211125IH18/143	
	IH 24 kids (Type: IH 24)	4211125IH24NK	
	IH 26 kids (Type: IH 25/2)	4211125IH25/23V	
	IH 26 (Type: IH 25/2)		
	IH 28	4211125IH28NT	
	IH 60	4211125IH60NP	
	IH 61 (Type: IH 60)		
	Medel Air Kids (Type: IH 24)	4211125IH24NK	
	Medel Air Mini (Type: IH 16)	4211125IH16NL	
	Medel Air Plus (Type: IH25/2)	4211125IH25/23V	
Mesh IH	IH 47	4211125IH47NX	Nebulizers (including compressor, ultrasonic, and mesh nebulizers) are medical devices for the nebulization of liquids and liquid medication (aerosols). 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 nebulizing 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.
	IH 49 (Type: IH 51)	4211125IH51NN	
	IH 51		
	IH 51 kids (Type: IH 51)		
	IH 57	4211125IH57P2	
	Medel Air Silence (Type: IH 51)	4211125IH51NN	

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

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: D1311700065, valid to 2030-11-15

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.

beurer	Title	**EU Declaration of Conformity**
	Date	2026-11-17

Signed for and on behalf of:	Beurer GmbH
Place, date of issue:	Ulm, 2026-11-17
Name, function, signature, stamp:	Max Harder, Team Lead Regulatory Affairs Beurer GmbH Söflinger Straße 218 • 89077 Ulm

Beurer Template 410-0040 / rev26 / 2026-04-24