

# Declaration of Conformity

We, the manufacturer

**Gustav Baehr GmbH**

Max-Eyth-Straße 39  
D-71332 Waiblingen  
SRN DE-MF-000007339

confirm under our sole responsibility that the product

**Baehr NEO**

Dry Technology Foot Care Device

Basis UDI-DI 4064667NEOE9

complies with the requirements of Regulation  
**(EU) 2017/745 on medical devices**

Intended Purpose

Active therapeutic foot care device for driving rotating instruments (cutters, abrasive caps, etc.) in podiatry – to alleviate foot complaints, to treat diseases and changes in the foot area - as well as in foot care. Podiatrists and chiropodists are trained specialists.

Conformity Assessment Procedure

according to Annex IX, Chapter 1 of regulation (EU) 2017/745 on Medical Devices

Classification

according to Annex VIII of regulation (EU) 2017/745 on Medical Devices  
the device has been classified as Class IIa (Rule 9)

Name, Address and Identification Number of the Notified Body:

mdc medical device certification GmbH  
Kriegerstraße 6, 70191 Stuttgart  
Identification Number 0483

This declaration – valid until May 31, 2029 – certifies the conformity of the above-mentioned product with the essential requirements according to Annex I of the regulation (EU) 2017/745. Conformity was determined by means of the above-mentioned conformity assessment procedure (see certificate D1122200020); the relevant provisions of the regulation and the applied standards were complied with.

Waiblingen, October 20<sup>th</sup>, 2025



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Michael Ladwig (Managing Director)