

EU-Declaration of Conformity

We, the manufacturer

Gustav Baehr GmbH

Max-Eyth-Straße 39

71332 Waiblingen

Germany

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, July 06th, 2026



Michael Ladwig (Managing Director)