beurer	Title	**EC Declaration of Conformity**
	Date	2026-04-02

Manufacturer:	Beurer GmbH (see address in footer)
SRN:	DE-MF-000005422
Product category:	TENS devices
Product type:	EM 49 (Model: EM 1 R RELIEF+)
Intended use:	The device is intended to relieve pain using TENS (transcutaneous electrical nerve stimulation) technology.

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

(EU) 2017/745	Medical device regulation (MDR)
Basic-UDI-DI:	4211125EM49NY
Classification/applied rule(s):	Class IIa/rule 9
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.

Signed for and on behalf of:	Beurer GmbH
Place, date of issue:	Ulm, 2026-04-02
Name, function, signature, stamp:	Max Harder, Team Lead Regulatory Affairs Beurer GmbH Söflinger Straße 218 • 89077 Ulm