beurer health and well-being	Title	**EC Declaration of Conformity**
	File	EM 29_EC DoC_Beurer_20200217

Manufacturer:	Beurer GmbH (see address in footer)
Product category:	TENS device
Product type:	EM 29

The product specified above is in conformity with the provisions of the following Union legislation and harmonized standards.

93/42/EEC	Medical device directive (MDD)
UMDNS code and name:	13-782 Stimulator, Electroanalgesia, transcutaneous, electrical nerve stimulation
Classification/applied rule(s):	Class IIa/rule 9
Conformity assessment procedure:	Annex VII in conjuction with annex V

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:	D1311700038, valid to 2024-05-26
	EN 60601-1:2006 + A11:2011 + A1:2013 EN 60601-1-2:2015 IEC 60601-2-10:2012
2011/65/EU	Restrictions of the use of certain hazardous substances in electrical and electronic equipment (ROHS)

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, 17.02.2020
Name, function, signature, stamp:	Daniel Kämmerer, Team Leader RA Beurer GmbH Söflinger Straße 218 • 89077 Ulm