beurer health and well-being	**Title**	**EC Declaration of Conformity**
	Date	2023-10-27

Manufacturer:	Beurer GmbH (see address in footer)
SRN:	DE-MF-000005422
Product category:	Lancing Device
Product type:	LD 01 LD 02 LD 03 LD 04

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

(EU) 2017/745	Medical device regulation (MDR)
Basic-UDI-DI:	LD 01: 4211125LD01N8 LD 02: 4211125LD02NA LD 03: 4211125LD03NC LD 04: 4211125LD04NE
Classification/applied rule(s):	Class I/rule 1
Conformity assessment procedure:	not applicable for class I devices
Certificate no. and validity:	D1311700051, valid to 2023-11-29

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, 2023-10-27
Name, function, signature, stamp:	Werner Meternek, Director Quality Management & Regulatory Affairs ppa. Beurer GmbH Söflinger Straße 218 • 89077 Ulm

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