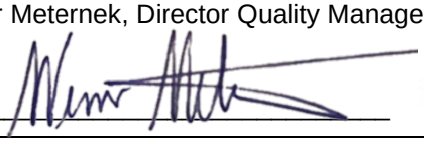
beurer health and well-being	**Title**	**EC Declaration of Conformity**
	Date	2025-02-03

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
Intended use:	The lancing device, in combination with a separate lancet needle, is intended for taking a blood sample for measuring glucose levels in human capillary blood. Use the lancing device only on the skin areas intended for taking the glucose measurement (fingertips).

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:	D1311700056, valid to 2026-11-29
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, 2025-02-03
Name, function, signature, stamp:	Werner Meternek, Director Quality Management & Regulatory Affairs ppa.  Beurer GmbH Söflinger Straße 218 • 89077 Ulm