

	Title	EC Declaration of Conformity
	File	GL 40_EC DoC_Beurer_20220208

Manufacturer:	Beurer GmbH (see address in footer)
Product category:	Blood glucose monitor
Product type:	GL 40
The product specified above is in conformity with the provisions of the following Union legislation and harmonized standards.	
98/79/EC	In vitro medical device directive (IVDD)
EDMA code and name:	21-06-01 Blood glucose meter
Classification:	Annex II/list B
Conformity assessment procedure:	Annex IV section 3
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:	D1311700050, valid to 2024-05-26
	EN 61010-1:2010 EN 61010-2-101:2017 EN 61326-1:2013 EN 61326-2-6:2013 EN ISO 15197:2015
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, 08.02.2022
Name, function, signature, stamp:	Daniel Kämmerer, Team Leader RA
	