beurer health and well-being	Title	**EC Declaration of Conformity**
	File	GL Lancing device_EC DoC_Beurer_20200622

Manufacturer:	Beurer GmbH (see address in footer)
Product category:	Lancing device
Product type:	Beurer Lancing device

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:	16-380 Lancing Devices, Blood
Classification/applied rule(s):	Class I/rule 1
Conformity assessment procedure:	Annex VII
Certificate no. and validity:	D1311700042, valid to 2022-12-18

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, 22.06.2020
Name, function, signature, stamp:	Daniel Kämmerer, Team Leader RA Beurer GmbH Söflinger Straße 218 • 89077 Ulm