

EC Declaration of Conformity Title

File

BM 54_EC DoC Beurer 20200807

Manufacturer:

Beurer GmbH (see address in footer)

Product category:

Blood pressure monitor

Product type:

BM 54

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:

16-174 Blood Pressure Monitor, electronic, manual

Classification/applied rule(s):

Class Ila/rule 10

Conformity assessment

Annex II - excluding section 4

procedure:

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:

D1311700043, valid to 2024-05-26

IEC 60601-1:2012 EN 60601-1-2:2015

IEC 80601-2-30:2009 + A1:2013

ISO 81060-1:2007

EN 1060-3:1997 + A2:2009

ISO 81060-2:2013

2014/53/EU

Radio equipment directive (RED)

EN 62479:2010 EN 50663:2017

EN 301 489-1 V2.2.3 (2019-11) Draft EN 301 489-17 V3.2.2 (2019-12)

EN 300 328 V2.2.2(2019-07)

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, 07.08.2020

Name, function, signature,

stamp:

Beurer GmbH Soninger Straße 218 · 89077 Ulm

Daniel Kämmeren Team Leader RA

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