

Title

EC Declaration of Conformity

File

ME 90_EC DoC Beurer 20200622

Manufacturer:

Beurer GmbH (see address in footer)

Product category:

Electrocardiograph

Product type:

ME 90

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:

12-599 ECG-Monitor

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

EN 60601-1:2006 + A1:2013

EN 60601-1-2:2015 IEC 60601-2-25:2011 IEC 60601-2-47:2001

2014/53/EU

Radio equipment directive (RED)

EN 62479:2010

EN 301 489-1 V2.1.1 (2017-02) EN 301 489-17 V3.1.1 (2017-02) EN 300 328 V2.1.1 (2016-11)

2011/65/EU

Restrictions of the use of certain hazardous substances in electrical and electronic equipment (ROHS)

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, 22.06.2020

Name, function, signature,

stamp:

Daniel Kämmerer, Team Leader RA

/ Beurer GmbH /Sötlinger/Straße 218 · 89077 Ulm

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