| beurer health and well-being | Title | EC Declaration of Conformity |
|---|---|---|
| | File | BM 49_EC DoC_Beurer_20200623 |

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|---|---|
| Manufacturer: | Beurer GmbH (see address in footer) |
| Product category: | Blood pressure monitor |
| Product type: | BM 49 |

The product specified above is in conformity with the provisions of the following Union legislation and harmonized standards.

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| 93/42/EEC | Medical device directive (MDD) |
| UMDNS code and name: | 16-174 Blood Pressure Monitor, electronic, manual |
| Classification/applied rule(s): | Class IIa/rule 10 |
| Conformity assessment procedure: | Annex II - excluding section 4 |

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:

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| Certificate no. and validity: | D1311700043, valid to 2024-05-26 |
| | IEC 60601-1:2012<br>EN 60601-1-2:2015<br>IEC 80601-2-30:2009 + A1:2013<br>ISO 81060-1:2007<br>EN 1060-3:1997 + A2:2009<br>ISO 81060-2:2013 |
| 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.

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| Signed for and on behalf of: | Beurer GmbH |
| Place, date of issue: | Ulm, 23.06.2020 |
| Name, function, signature, stamp: | Daniel Kämmerer, Team Leader RA<br>Beurer GmbH<br>Söflinger Straße 218 • 89077 Ulm |