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|  | <b>Title</b> | <b>EC Declaration of Conformity</b> |
|                                                                                  | <b>File</b>  | GL 50_EC DoC_Beurer_20220420        |

  

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| Manufacturer:                                                                   | Beurer GmbH (see address in footer) |
| Product category:                                                               | Blood glucose monitor               |
| Product type:                                                                   | GL 50                               |
| The product specified above is in conformity with the following specifications. |                                     |

  

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| 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                                                                                                                          |
| <p>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:</p> |                                                                                                                                             |
| Certificate no. and validity:                                                                                                                                                                                                                    | D1311700053, valid to 2024-12-17<br><br>EN 61010-1:2010<br>EN 61010-2-101:2017<br>EN 61326-1:2013<br>EN 61326-2-6:2013<br>EN ISO 15197:2015 |

  

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| 2011/65/EU | Restrictions of the use of certain hazardous substances in electrical and electronic equipment (RoHS)<br><br>EN IEC 63000:2018 |
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| 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, 2022-04-20                                                                                                                                                               |
| Name, function, signature, stamp:                                                           | Daniel Kämmerer, Team Lead Regulatory Affairs<br><div style="text-align: center;">  </div> |