

Title EC Declaration of Conformity

Date 2023-10-19

Manufacturer: Beurer GmbH (see address in footer)

SRN: DE-MF-000005422

Product category: CPR devices

Product type: RH 112

The product specified above is in conformity with the following specifications.

(EU) 2017/745 Medical device regulation (MDR)

Basic-UDI-DI: 4211125RH112EY

Classification/applied

rule(s):

Class I/rule 13

Conformity assessment

procedure:

not applicable for class I devices

Certificate no. and validity: D1311700051, valid to 2023-11-29

2011/65/EU Restrictions of the use of certain hazardous substances in electrical and electronic

equipment (RoHS)

EN IEC 63000:2018

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, 2023-10-19

Name, function, signature,

stamp:

Werner Meternek, Director Quality Management & Regulatory Affairs

Beurer GmbH Söflinger Straße 218 · 89077 Ulm