

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 provisions of the following Union legislation and harmonized standards.

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

Basic-UDI-DI: 4211125BMD0024XB

EMND code and name: C99 - CARDIOCIRCULATORY DEVICES – OTHER

Classification/applied rule(s): Class I/ rule 1

Conformity assessment procedure: not applicable for class I devices

Certificate no. and validity: D131170048, valid to 2023-11-29

EN 60601-1:2006 + A1:2013

EN 60601-1-2:2015

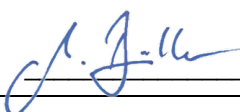
2011/65/EU Restrictions of the use of certain hazardous substances in electrical and electronic equipment (ROHS)

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, 01.12.2021

Name, function, signature, stamp: Marco Bühler, Managing Director


Beurer GmbH
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