

## EC Declaration of Conformity

Manufacturer: OMRON HEALTHCARE Co., Ltd.  
Single Registration Number: JP-MF-000007213  
Address: 53, Kunotsubo, Terado-cho, Muko, KYOTO, 617-0002 JAPAN  
European Authorised Representative: OMRON HEALTHCARE EUROPE B.V.  
Single Registration Number: NL-AR-000002683  
Address: Scorpius 33, 2132 LR Hoofddorp, The Netherlands  
Product Category: Electronic Sphygmomanometers/Blood Pressure Monitors  
Model (code): HBP-1320 (HBP-1320-E)  
Basic UDI-DI: 40156721112952  
MDR Classification: Class IIa (MDR Annex VIII Rule 10)

We herewith declare, under our sole responsibility, that the above mentioned product meets the provisions of the following European Union Regulations, Council Directives and Standards. All supporting documentation is retained at the premises of the manufacturer and the European Authorized Representative.

This Declaration of Conformity is valid in connection with all the shipping inspection reports for the respective batch of produced devices.

General applicable regulations:	Medical Device Regulation (EU) 2017/745
Standards:	EN 1041:2008+A1:2013      EN ISO 10993-1:2020 EN 1060-1:1995+A2:2009      EN ISO 10993-5:2009 EN 1060-3:1997+A2:2009      EN ISO 10993-10:2013 EN 60601-1:2006+A1:2013      EN ISO 13485:2016 EN 60601-1-2:2015      EN ISO 14971:2012 EN 60601-1-6:2010+A1:2015      EN ISO 15223-1:2016 EN 62304:2006+A1:2015      EN ISO 81060-2:2019+A1:2020 EN 62366-1:2015 EN 80601-2-30:2010+A1:2015

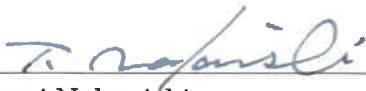
Notified Body:	TÜV Rheinland LGA Products GmbH
Address:	Tillystrasse 2, 90431 Nuremberg, Germany
ID No:	Notified under number 0197 to the EC Commission
Certificate Registration No:	Annex IX: HZ 2102042-1

General applicable directives:	Radio Equipment Directive 2014/53/EU
Standards:	

General applicable directives:	RoHS Directive 2011/65/EU, (EU)2015/863 and (EU)2017/2102
Product Category for RoHS:	Category 8 (Medical devices)
Standards:	EN IEC 63000:2018

Place / Date: Kyoto / May 9, 2023

Signature:

Name:   
Position: Takefumi Nakanishi  
General Manager  
Regulatory Affairs Department

Attachment to EC Declaration of Conformity No. OHQ(CS)-DoC(MDR)-9546059

**Intended purpose of the model:**

The device is a digital monitor intended for use in measuring blood pressure and pulse rate in adult and pediatric patient population with arm circumference ranging from 12 cm to 50 cm (from 5 inches to 20 inches).