

## EU 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: Accessory for Electronic Sphygmomanometers/Blood Pressure Monitors  
Product Description: Blood Pressure Monitor Cuff  
Model (code): GS CUFF2 M (HXA-GCFM-PBE)  
Basic UDI-DI: 40156721115655  
MDR Classification: Class I (MDR Annex VIII Rule 1)

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 13485:2016 EN 60601-1-6:2010+A1:2015      EN ISO 14971:2019 EN 62366-1:2015      EN ISO 15223-1:2016 EN IEC 80601-2-30:2019      EN ISO 81060-2:2019+A1:2020 EN ISO 10993-1:2020 EN ISO 10993-5:2009 EN ISO 10993-10:2013

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 / July 24, 2023

Signature:

Name:   
Position: Takefumi Nakanishi  
General Manager  
Regulatory Affairs Department



Attachment to EU Declaration of Conformity No. OHQ(CS)-DoC(MDR)-9546008B

Intended purpose of the model:

This product is an upper arm cuff for OMRON non-invasive blood pressure monitors.