No: OHQ(CS)-RAF-231051

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:	Electronic Sphygmomanometers/Blood Pressure Monitors
Model (code):	M2+ (HEM-7146-E)
Basic UDI-DI:	40156721139857T2
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 60601-1:2006+A1:2013+A12:2014	EN ISO 10993-5:2009
	EN 60601-1-2:2015 +A1:2021	EN ISO 10993-10:2013
	EN 60601-1-6:2010+A1:2015	EN ISO 13485:2016
	EN 60601-1-11:2015	EN ISO 14971:2019
	EN 62304:2006+A1:2015	EN ISO 15223-1:2021
	EN 62366-1:2015	EN ISO 81060-2:2019+A1:2020
	EN IEC 80601-2-30:2019	
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:	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 / December 13, 2023

Signature:

オムロンヘルスケア株式会社
〒617-0002 京都府向日市寺戸町九ノ坪53番地

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Position: General Manager
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