

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): M6 Comfort (HEM-7360-E)
Basic UDI-DI: 40156721117457
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 14971:2019 EN 60601-1-6:2010+A1:2015 EN ISO 15223-1:2016 EN 60601-1-11:2015 EN ISO 81060-2:2019+A1:2020 EN 62304:2006+A1:2015 EN 62366-1:2015 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 / February 28, 2023

Signature:

Name: 
Position: General Manager
Regulatory Affairs Department

Intended purpose of the model:

The device is a digital monitor intended for use in measuring blood pressure and pulse rate in adult patient population. The device detects the appearance of irregular heartbeats during measurement and indicates this via a symbol with the measurement results. It is mainly designed for general household use. The device can detect an irregular pulse suggestive of Atrial Fibrillation (Afib). Please note that the device is not intended to diagnose Afib. A diagnosis of Afib can only be confirmed by an Electrocardiogram (ECG). If the Afib symbol appears, consult with your physician.