

## EC Declaration of Conformity

Manufacturer: OMRON HEALTHCARE Co., Ltd.  
Address: 53, Kunotsubo, Terado-cho, Muko, Kyoto 617-0002 JAPAN  
European Representative: OMRON HEALTHCARE EUROPE B.V.  
Address: Scorpius 33, 2132 LR Hoofddorp, The Netherlands  
Product Category: Electronic Sphygmomanometers/Blood Pressure Monitors  
Model (code): RS1 (HEM-6160-E)  
Classification for MDD: Class IIa (MDD Article 9 Annex IX Rule 10)  
Product Category for RoHS: Category 8 (Medical devices)

We herewith declare that the above mentioned product meets the provisions of the following European Committee Council Directives and Standards. All supporting documentation are retained at the premises of the manufacturer and the notified body.  
This Declaration of Conformity is valid in connection with the shipping inspection reports for the respective batch of produced devices.

### Directives

General applicable directives:	Relevant regulations and harmonized standards
93/42/EEC Medical Device Directive (MDD)	EN ISO 15223-1:2016 EN 1041:2008 EN 1060-1:1995+A2:2009 EN 1060-3:1997+A2:2009 EN 60601-1:2006+A1:2013 EN 60601-1-2:2015 EN 60601-1-6:2010+A1:2015 EN 60601-1-11:2015 EN 80601-2-30:2010+A1:2015 EN 62304:2006+A1:2015 EN 62366:2008+A1:2015 EN ISO 10993-1:2009 EN ISO 10993-5:2009 EN ISO 10993-10:2013 EN ISO 14971:2012 EN ISO 81060-2:2014
2011/65/EU Restriction of Hazardous Substances (RoHS)	EN50581:2012

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 II : HD 60100203 0001

Place / Date: Kyoto / September 10, 2018

Signature:

Name:   
Position: Takefumi Nakanishi  
General Manager  
Regulatory Affairs Department