

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): M7 Intelli IT (HEM-7322T-E)
Classification for MDD: Class IIa(MDD 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 under the premises of the manufacturer.

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

Directives

General applicable directives: Standards	Medical Device Directive 93/42/EEC EN 980:2008 EN 1041:2008 EN 1060-1:1995+A2:2009 EN 60601-1:2006 EN 60601-1-6:2010 EN 62304:2006 EN 62366:2008 EN ISO 10993-1:2009 EN ISO 10993-10:2010 EN ISO 14971:2012 ISO 81060-2:2013	EN 1060-3:1997+A2:2009 EN 60601-1-2:2007 EN ISO 10993-5:2009
Notified Body: Address: ID No: Certificate Registration No:	TÜV Rheinland LGA Products GmbH Tillystrasse 2, 90431 Nuremberg, Germany Notified under number 0197 to the EC Commission Annex II : HD 60100203 0001	

General applicable directives: Standards	Radio Equipment Directive 2014/53/EU EN 300 328 V2.1.1 EN 301 489-17 V3.1.1 EN 60950-1:2006+A11:2009+A1:2010+A12:2011+A2:2013	EN 301 489-1 V2.1.1 EN 62479:2010
General applicable directives: Standards	RoHS Directive 2011/65/EU EN50581:2012	

Place / Date: Kyoto / May 8, 2017

Signature:



Name: Takefumi Nakanishi

Position: General Manager

Customer Satisfaction Management Division