OMRON

No: OHQ(CS)-QS-091187F

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:	Electroanalgesic Transcutaneous Stimulators
Model (-code):	E4(HV-F128-E)
Classification:	Class IIa(MDD Annex IX Rule 9)

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 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:	93/42/EEC Medical Device Directive (MDD)
Standards:	EN 980:2008 EN 1041:2008 EN 60601-1:1990+A1:1993+A2:1995 EN 60601-1-2:2007 EN 60601-1-4:1996+A1:1999 EN 60601-2-10:2000+A1:2001 EN 62304:2006 EN 62366:2008 EN ISO 10993-1:2009 EN ISO 10993-5:2009 EN ISO 10993-10:2010 EN ISO 14971:2012
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 / May 29, 2015
Signature:	
Name:	Norikazu Yasue
Position:	General Manager Customer Satisfaction Management Division