OMRON

No: OHQ(CS)-DOC-110002B

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:	Accessory for Nebulizers
Model (-code):	C30-E-AC
Classification for MDD:	Class I(MDD Annex IX Rule 1)
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 and Harmonized Standards

General applicable directives:	Relevant regulations and harmonized standards
93/42/EEC Medical Device Directive (MDD)	EN 980:2008 EN 1041:2008 EN 60601-1:1990+A1:1993+A2:1995 EN 60601-1-2:2007 EN ISO 14971:2012
2011/65/EU Restriction of Hazardous Substances (RoHS)	EN50581:2012

Place / Date: Kyoto / October 3, 2014

Signature:

M. Yasue

Name: Norikazu Yasue

Position: General Manager
Customer Satisfaction Management Division