

EC Declaration of Conformity

Manufacturer: OMRON HEALTHCARE Co., Ltd.
Address: 24, Yamanouchi Yamanoshita-cho, Ukyo-ku, Kyoto
615-0084 JAPAN

European Representative: OMRON HEALTHCARE EUROPE B.V.
Address: Kruisweg 577, 2132 NA Hoofddorp, The Netherlands

Product Category: Accessory for Electroanalgesic Transcutaneous
Stimulators

Model Name (-code): E Pads (HV-OM3PAD-E)
Classification: Class I (MDD Annex IX Rule 1)

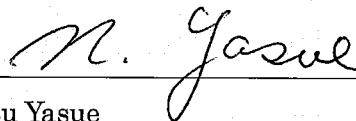
We herewith declare that the above mentioned product meets the provisions of the following European Committee Council Directives and Standards. All supporting documentation are retained under the premises of the manufacturer.

Directives

General applicable directives: Medical Device Directive (MDD) 93/42/EEC
Standards: EN ISO 10993-1:2009
EN ISO 10993-5:2009
EN ISO 10993-10:2009

Place / Date: Kyoto, Japan / June 15, 2011

Signature:



Name: Norikazu Yasue

Position: General Manager
Customer Satisfaction Management Division