

## EC Declaration of Conformity

Manufacturer: OMRON HEALTHCARE Co., Ltd.  
Address: 53, Kunotsubo, Terado-cho, Muko, KYOTO, 617-0002  
JAPAN  
European Authorised Representative: OMRON HEALTHCARE EUROPE B.V.  
Address: Scorpius 33, 2132 LR Hoofddorp, The Netherlands  
Product Category: Electroanalgesic Transcutaneous Stimulators  
Model (code): Avail (HV-F601T-E)

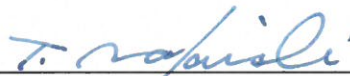
We herewith declare, under our sole responsibility, that the above mentioned product meets the provisions of the following European Union Regulations, Council Directives and Standards. All supporting documentation is retained at the premises of the manufacturer and the European Authorized Representative.

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

General applicable directive:	Radio Equipment Directive 2014/53/EU	
Standards:	EN 300 328 V2.2.2	EN 301 489-1 V2.2.3
	EN 301 489-17 V3.2.4	EN 62479:2010
	EN IEC 62368-1:2020+A11:2020	

Place / Date: Kyoto / July 15, 2022

Signature:



Name: Takefumi Nakanishi  
Position: General Manager  
Regulatory Affairs Department