

## EC Declaration of Conformity

Manufacturer: OMRON HEALTHCARE Co., Ltd.  
Single Registration Number: JP-MF-000007213  
Address: 53, Kunotsubo, Terado-cho, Muko, KYOTO, 617-0002 JAPAN  
European Authorised Representative: OMRON HEALTHCARE EUROPE B.V.  
Single Registration Number: NL-AR-000002683  
Address: Scorpium 33, 2132 LR Hoofddorp, The Netherlands  
Product Category: Clinical Electronic Thermometers  
Model (code): Flex Temp Smart (MC-343F-E4)  
Basic UDI-DI: 40156721136056  
MDR Classification: Class IIa (MDR Annex VIII Rule 10)

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 regulations: | Medical Device Regulation (EU) 2017/745  |
| Standards:                      | EN 1041:2008+A1:2013 EN ISO 10993-1:2020<br>EN 12470-3:2000+A1:2009 EN ISO 10993-5:2009<br>EN 60601-1:2006+A1:2013+A12:2014 EN ISO 10993-10:2013<br>EN 60601-1-2:2015 EN ISO 10993-11:2018<br>EN 60601-1-6:2010+A1:2015 EN ISO 10993-23:2021<br>EN 60601-1-11:2015 EN ISO 13485:2016<br>EN 62366-1:2015 EN ISO 14971:2019<br>EN ISO 15223-1:2016<br>EN ISO 17664:2017<br>EN ISO 80601-2-56:2017 +A1:2020 |
| 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 IX: HZ 2102042-1   |

|                                |   |
|--------------------------------|---|
| General applicable directives: | RoHS Directive 2011/65/EU, (EU)2015/863 and (EU)2017/2102 |
| Product Category for RoHS:     | Category 8 (Medical devices)                              |
| Standards:                     | EN IEC 63000:2018   |

Place / Date: Kyoto / February 28, 2023

Signature:

  
Name: Takefumi Nakanishi  
Position: General Manager  
Regulatory Affairs Department



Attachment to EC Declaration of Conformity No. OHQ(CS)-DoC(MDR)-3293608B

**Intended purpose of the model:**

The thermometer you purchased offers a safe, accurate and quick temperature reading. You can measure your temperature either in the anus (rectally), in the mouth (orally) or in the armpit (axillary).