withings

EU DECLARATION OF CONFORMITY

We,

Withings SA 2, rue Maurice Hartmann, 92130 Issy-les-Moulineaux – France

Declare under our sole responsibility of the manufacturer, that the product:

Product name: BPM Connect Brand name: Withings Model: WPM05

Risk Classification: IIa

is in conformity with the relevant Union harmonization Legislation:

Directive

93/42/CEE as amended by 2007/47/EC – Annex I and the certified quality system per Annex II, excluding (4) (Medical device)

| Directive | 2014/53/EU (RED) |
|-----------|-------------------|
| Directive | 2011/65/EU (RoHS) |

The conformity with the essential requirements of the 93/42/CEE has been demonstrated against the following standards:

| Quality Management System | EN ISO 13485:2016 |
|------------------------------|---|
| | EN ISO 14971:2012 |
| | |
| Medical Electrical Equipment | EN 60601-1-11:2010, 2015 |
| | EN 6060-1 :1995 + A2 :2009 |
| | EN 1060-3 :1997 + A2 :2009 |
| | EN 81060-2-30 :2010,2013 |
| | EN 15223-1 :2016 |
| | |
| EMC | EN 60601-1-2-2015 (EN 55011 :2016) |
| | EN 301 489-1 V2.1.1 (2017-02) + EN 301 489-17 |
| | V3.1.1 (2017-02) |
| | |
| RF spectrum use | EN 300 328 v2.1.1 |
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| Biocompatibility | ISO 10993-1:2009 |

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The conformity assessment procedure referred to in Directive 93/42/CEE has been by Ente Certificazione Macchine, Via Ca Bella 243, 40053 Valsamoggia, Castello di Serravalle, Italy.

Thus, $\mathbf{C}\mathbf{\epsilon}$ is placed on the product

Signed on behalf of Withings SA, in Issy-les-Moulineaux, July 5th, 2019.

Name:Xavier DebreuilPosition:Product director

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