

Applicant certifies that the following designated product

Product Name: Beauty Device

Model No.: TheraFace PRO

➤ **Is in fully conformity with the harmonized standard(s)**

-EN 55014-1:2017/A11:2020
-EN 55014-2:2015
-EN 61000-3-2:2014
-EN IEC 61000-3-2:2019
-EN 61000-3-3:2013/A1:2019
-EN 301 489-1 V2.2.3:2019
-EN 301 489-17 V3.2.4:2020
-EN 50663:2017
-EN 300 328 V2.2.2:2019
-EN 60335-1:2012/A2:2019
-EN 60335-2-32:2003/A2:2015
-EN 62233:2008
-EN 50564:2011
-EN 62479:2010
-EN 303 417 V1.1.1

➤ **Under the Council Directive**

2014/53/EU Radio Equipment Directive
2014/35/EU Low Voltage Directive
2014/30/EU Electromagnetic Compatibility
2011/65/EU (RoHS) Directive
2015/863/EU (RoHS amendment) Directive
EC regulation 278/2009:2009-04-06
EC regulation 2019/1782:2019-10-25
EC Regulation 1275/2008:2008-12-17
EC642/2009:2009-07-22
Eu617/2013:2013-06-26
EU 801/2013:2013-08-02

➤ **The declaration is the sole responsibility of the applicant**

Company Name : Therabody Inc.
Company address: 6100 Wilshire Blvd. Suite 200 Los Angeles, CA 90048-5107,USA

Signature: *CJ Frederick, AIA 13 JAN 2022*

Compliance Team (Therabody, Inc.)