1) **Manufacturer: HYPERICE, INC.**

Address: **525 Technology Dr. | Suite 100, Irvine, California 92618, USA**

2) Product(s):

Device name	Type/ model/ref number	First date of CE-compliance
Venom 2 Back	V20001	2018-03-29
Venom 2 Leg (knee)	V20002	2018-03-29
Venom 2 Shoulder Right	V20003	2018-03-29
Venom 2 Shoulder Left	V20004	2018-03-29

3) The product(s) described above is in conformity with:

Title	Document No.
RED Directive	2014/53/EU
LVD Directive	2014/35/EU
RoHS Directive	2011/65/EU
REACH	1907/2006

4) Additional information *(conformity procedure, Notified Body, CE certificate, Registration nr., etc.)*:

Conformity assessment procedure for CE marking:

2014/35/EU, Annex III

2014/53/EU, Annex II

Conformity to the essential requirements of the legislation(s) have been demonstrated by using the following standards:

Health and Safety (Art 3(1) (a)): EN 60335-1: 2012+AC:2014 + A11: 2014 + A13:2017 + A1:2019 + A14: 2019 + A2: 2019

EN 60335-2-32: 2003 + A1: 2008 + A2: 2015

EN 62479: 2010

EMC (Art.3(1) (b)): EN 301 489-1 v2.2.3

EN 301 489-17 v3.2.4

EN 55014-1: 2017+A11:2020

EN 55014-2: 2015

EN 61000-3-3: 2013+A1: 2019

EN IEC 61000-3-2: 2019

Spectrum (Art.3(2)): EN 300 328 v2.2.2

These Products are fully compliant and do not contain the restricted substances above levels noted in EU Directive 2011/65/EU.

Scott Lambert

Manager, Quality & Regulatory Systems

Hyperice, Inc.

2024-04-01