

## EC DECLARATION OF CONFORMTIY

## I.A.C.E.R. S.r.l

Via S.Pertini 24/A – 30030 Martellago (Ve), Italia

herewith declares under its own responsibility, that the products

MIO-CARE TENS
MIO-CARE FITNESS
MIO-CARE BEAUTY

UMDNS Code: **13762**Batch no.:

Series no.:

comply with the essential safety requirements of the European Medical Device Directive 93/42/EEC (transposed in Italy by the D.Lgs. 46/97), as modified by the Directive 2007/47/EC (D.Lgs.37/2010) and further modifications/integrations.

The products have been assigned to class IIa, according to Annex IX, rule 9 of the Directive 93/42/EEC (and further modifications/integrations) and bear the mark



Compliance of the concerned products with the Directive 93/42/EEC has been assessed and certified by the Notified Body

0068 – MTIC InterCert S.r.l.

Via G. Leopardi 14, Milano (MI) 20123, Italia
Certificate no.: 0068/QCO-DM/234-2020

following the certification procedure according to Annex II (excluding point 4) of the Directive 93/42/EEC. The devices comply with the following applicable standards:

EN 60601-1:2006 + A1:2013, EN 60601-1-2:2015, EN 60601-1-6:2010 + A1:2015, EN 60601-1-11:2015, • EN 60601-2-10:2015, EN ISO 14971: 2012, ISO 10993-1: 2009, ISO 10993-5: 2009, ISO 10993-10: 2013, IEC 62133:2012, EN 62304: 2006 +A1:2015, EN 62366-1:2015.

It is also claimed that:

- The devices do not incorporate, as an integral part, any substance or a human blood derivate of point 10 of Annex 1 (Directive 2007/47/CE);
- tissue of animal origin, provided for in Directive 2003/32/CE, haven't been used in production.

Martellago, 03/08/2020

Place, date

MASSIMO MARCON

Legal Representative