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

I-TECH LA8000

I-TECH LA10000

UMDNS Code**: 12299**

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 IIb, according to Annex IX, rule 9 of the Directive 93/42/EEC (and further modifications/integrations) and bear the mark

CE 0068

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/232-2020**

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 60601-2-22:2013, EN 60825-1: 2009, EN ISO 14971: 2012, EN 62304: 2006, EN 62366-1: 2008.

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, 19/06/2020

Place, date

MASSIMO MARCON

Legal Representative