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 family product

LaMagneto

which includes the following models

☐ LaMagneto Basic ☐ LaMagneto ☐ LaMagneto Pro ☐ LaMagneto X

UMDNS Code**: 12415**

Batch no.:

Series no.:

has been designed and manufactured according to 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

CE 1936

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

1936 – TÜV Rheinland Italia
Via Enrico Mattei 3, 20010 Polignano Milanese (MI), Italia
Certificate no.: HD60134521

following the certification procedure according to Annex II (excluding point 4) of the Directive 93/42/EEC.

Martellago, 09/08/2021 — MASSIMO MARCON

Place, date — *Legal Representative*

I.A.C.E.R. Srl
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Vat Number: IT00185480274 - R.E.A.: VE N. 120250 - M. VE001767 - Share Capital: € 110.000,00 i.v.