EC DECLARATION OF CONFORMTIY

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

Via Enzo Ferrari 2 – 30037 Scorzè (Ve), Italia

herewith declares under its own responsibility, that the product

LaMagneto Pro

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.

Scorzè, 04/08/2022 — *Place, date*

MASSIMO MARCON — *Legal Representative*

I.A.C.E.R. Srl
Via Enzo Ferrari 2 - 30037 Scorzè (VE) – Italia/Italy - Tel.: (+39) 041/5401356 - Email: iacer@iacer.it
PEC: iacer@pec.it - Web: www.itechmedicaldivision.com - Cod. Fisc./P.IVA/VAT N.: IT00185480274
R.E.A.: VE N. 120250 - M. VE001767 - Codice SDI/SDI Code: SUBM70N - Cap. Soc.: € 1.000.000,00 i.v.