CE Declaration of conformity

IACER S.r.l., headquartered in Italy, via S. Pertini 24/A 30030 Martellago (VE), declares on its own responsibility that following device:

Model:

- **GP-1**

Is manufactured in conformity with Council Directive 93/42/EEC (MDD) dated 14 June 1993 *(D. Lgs. 46/97 dated 24 February 1997 "Attuazione della Direttiva 93/42/CEE concernente i dispositivi medici)*, as amended by Council Directive 2007/47/EC dated 5 September 2007 *(D. Lgs. 37/2010 dated 25 January 2010)* and subsequent amendments / additions.

The above mentioned device is Class IIa equipment, with reference to Council Directive 93/42/EEC (MDD), annex IX rule 9 (and subsequent amendments / additions)

Notified Body: Kiwa Cermet Italia Spa, Via di Cadriano 23 – 40057 Cadriano di Granarolo (BO) ITALY

Certification path: Annex II (without point 4).

Martellago, 25/10/16

CEO

Caprara Mario