DECLARATION OF CONFORMITY CE

Calvenzano, December 20th 2010

LUROPAS del dottor Molinari & C. s.r.l., located in Calvenzano (BG), via Vivaldi 4, Italy,

DECLARES UNDER ITS OWN RESPONSIBILITY

that the following products that belong to the category of ANKLE BRACES coded and named:

- 537 cavigliera sporting
- 538 cavigliera steccata regolabile
- 540 cavigliera dynamic
- 560 cavigliera elastica a doppia forza
- 888 cavigliera air bi-valva

are classified according to the 93/42/EEC Directive
as **non active medical device class I.**

Besides, we declare that the above mentioned medical devices:
- are not instruments of measure
- are not used for clinics enquiries
- are commercialized in non sterile packaging
- satisfied the essential requirements of the Annex n. 1 and applicable instructions of the 93/42/EEC Directive.

LUROPAS, shall keep and have available to the Authority all the technical documentation specified in the Annex n. VII of the 93/42/EEC Directive for a period of five years from the last date of the making of the product.

President's signature

LUROPAS
del dott. Molinari & C. s.r.l.
Cap Soc. € 46.000,00 i.v.
Via Vivaldi, 4 - 24040 CALVENZANO (BG)
Tel 0363 [illegible] - Fax 0363 85281
Cod. Fisc. e Partita IVA 00383840169