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 ELASTIC THERAPEUTIC STOCKINGS STRONG COMPRESSION 180 DENIERS, coded and named:

- 835 collant 180 super cell
- 836 calza autoreggente 180 super cell
- 837 gambaletto 180 super cell

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 steril 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.
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