Hospidex

Declaration of Conformity

We hereby certify that the CE marked product

ALCO-PREP ®

- Has been classified as class I Medical Device following rule 1 of annex IX of the Medical Device Directive 93/42/EEC as amended by Directive 2007/47/EEC.
- Is in conformity with the essential requirements and meet the provisions of directive 93/42/EEC as amended by Directive 2007/47/EEC.

This declaration is made on the basis of

- The technical documentation in accordance with annex VII of the Medical Device Directive 93/42/EEC as amended by Directive 2007/47/EEC。

This certificate is valid for the above mentioned device, bearing the CE mark and originated & manufactured by Hospidex France SAS, Athélia 4, 297 avenue du Mistral, 13705 La Ciotat cedex, France in collaboration with Hospidex nv, Grijpenlaan 23, B-3300 Tienen, Belgium, distributor.

Wim Steegmans
Operations & Regulatory Affairs Manager

Tienen, Belgium,
April 1st, 2017

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