

## Declaration of Conformity

We hereby certify that the CE marked product

### ***BENZA-CLEAN*** ®

- 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