

EC Declaration of Conformity

according to the

Medical Device Directive : 93/42/EEC. (amended by 2007/47/EC)

The undersigned, Pierre Chang, representing Chang Ming Trading Co., Ltd., 11F-10, No.188, Sec. 5, Nanking E. Rd., Taipei, Taiwan, R.O.C.; the supplier declares that the product described hereafter:

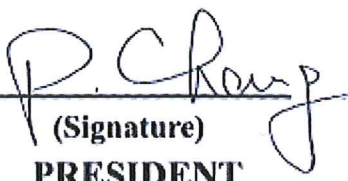
Product Name:
HI-CLEAN SPRAY

Model:
HI-CLEAN 550

Classification:
Class I

Provided that it is used and maintained in accordance with the generally accepted codes of good practice and the recommendations of the instruction manual, meets the essential safety and health requirements of the Medical Device Directive 93/42/EEC (amended by 2007/47/EC).

For the most specific risks of this equipment, safety and compliance with the essential requirements of the Directive has been based on elements of:
- The International Standard ISO 14971; 2000(E); Medical devices –
Application of risk management to medical devices.


(Signature)
PRESIDENT
(Position)



Signed in TAIWAN (place) on 27 May, 2013 (date)

Valid till 27 May, 2018 (date)