

EC DECLARATION OF CONFORMITY (Medical Device Directive 93/42/EEC Annex VII)

Cederroth AB

Hereby declare that the Medical Device products listed below conform to the relevant provisions of the Swedish Law regarding Medical Devices (1993:584) and the current version of LVFS 2003:11, which is the Swedish implementation of the Medical Device Directive 93/42/EEC including amendments to date.

(For devices class I sterile and class IIa as verified by our Notified Body, # 0413)

Brand	REF	Name of product	Medical Device Class
Cederroth	726000	Cederroth Eye & Wound Cleansing Spray	IIa

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Teija Ålander
Quality Assurance and Regulatory Affairs Manager
Wound Care Division, Cederroth AB