

EC DECLARATION OF CONFORMITY

(Medical Device Directive 93/42/EEC Annex VII)

Orkla Care 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 IIa as verified by our Notified Body, # 0413 according to Medical Device Directive 93/42/EEC, Annex II, EC Certificate A2 41315275-06.

REF	Name of product	Medical Device Class	GMDN
726000	Cederroth Eye & Wound Cleansing Spray	IIa	63285

2021-04-30

Teija Ålander
Regulatory Affairs and Quality Manager
Wound Care, Orkla Care AB