

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 I sterile as verified by Notified Body # 0413 according to Medical Device Directive 93/42/EEC, Annex II. EC-certificate No. 41315275-06.

REF	Name of product	Medical Device Class	EMDN
1910	Cederroth 4-in-1	I s	M03040199
1911	Cederroth 4-in-1 Mini	I s	M03040199

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