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	GMDN
1910	Cederroth	4-in-1	I s	59022
1911	Cederroth	4-in-1 Mini	I s	59022

2021-04-30

Teija Ålander
2021.04.30
12:03:31 +02'00'

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

Postadress/Postal address
Orkla Care AB
Box 1336
SE-171 26 Solna
www.cederroth.com

Besöksadress/Visiting address
Svetsarvägen 15
Solna
Sweden
www.orkla.com

Telefon/Telephone
010 142 64 00
International
+46 (0)10 142 64 00

Orgnr
556052-5478
VAT No:
SE556052547801