

STATEMENT PROCEDURE PACK

REGULATION ON MEDICAL DEVICES (EU) 2017/745, ARTICLE 22

ORKLA CARE AB

SRN: Not assigned yet

This statement is issued in accordance with Article 22 of the Regulation on Medical Devices (EU) 2017/745.

The manufacturer has combined devices bearing a CE marking with other devices or products according to paragraph 1 of Article 22 of the Regulation on Medical Devices (EU) 2017/745, in a manner that is compatible with the intended purpose of the devices or other products and within the limits of use specified by their manufacturers, in order to place them on the market as a procedure pack.

We hereby declare that we have verified the mutual compatibility of the devices and, if applicable other products, in accordance with the manufacturers' instructions and have carried out their activities in accordance with those instructions.

We hereby declare that we have packaged the system or procedure pack and supplied relevant information to users incorporating the information to be supplied by the manufacturers of the devices or other products which have been put together.

We hereby declare that the activity of combining devices and, if applicable, other products as a procedure pack was subject to appropriate methods of internal monitoring, verification and validation.

Basic UDI-DI: 7310610502004D

EMDN (CND) code: V0501

GMDN code: 44047

Medical purpose: Assortments of products to be used in First Aid situations.
Single use pack.

List of products on next page.

REF	Name of Procedure Pack	
2596	Cederroth First Aid	Protection Kit
1893	Cederroth First Aid	Net Dressing
1880	Cederroth First Aid/ Salvequick	Wound Cleansers & Plasters
2030	Cederroth First Aid	First Aid Refill for large F/A cabinet
264000	Cederroth First Aid	Refill First Aid Panel
51011002	Cederroth First Aid	First Aid refill to station
51000026	Cederroth First Aid	Set of mixed DIN products

Solna 2021-04-29

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