EU DECLARATION OF CONFORMITY

REGULATION ON MEDICAL DEVICES (EU) 2017/745, ANNEX IV

ORKLA WOUND CARE AB

SRN: SE-MF-000021041

This EU Declaration of Conformity is issued under the sole responsibility of Orkla Wound Care AB.

We hereby declare that the Medical Device products listed below conform to the Regulation on Medical Devices (EU) 2017/745.

Conformity assessment route:
The products are class I medical devices and have been CE marked in accordance with MDR 2017/745 Article 52 (7); Conformity assessment procedure of class I medical devices.

Basic UDI-DI:	7340210051003X		
EMDN (CND) code:	M04010102	**GMDN code:**	44990
Intended purpose:	The products are intended to be used as a mechanical barrier and for absorption of exudates from minor and larger superficial wounds.		

REF	Name of product		Risk class
6796	Salvequick	Detectable Finger bandage Refill	I
51030126	Salvequick	Detectable Fingertip/Regular Plasters Refill	I
51030127	Salvequick	Detectable Plasters Refill	I

Common Specifications applied: No applicable CS available

Solna 2024-08-06

Johanna Brinck

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