

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: 7340210004003B

EMDN (CND) code: M04010199 **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
6444	Salvequick Textile Plasters Refill	I
644401	Salvequick Textile Plasters 1-p Refill	I
6454	Salvequick Textile Fingertip Plasters Refill	I
6470	Salvequick Textile Plasters XL Refill	I
6496	Salvequick Textile Finger Bandage Refill	I

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

Besöksadress/Visiting address
Svetsarvägen 15
Solna, Sweden
www.orklahealth.com/wound-care/
www.cederroth.com

Telephone
+46 (0)10 142 64 00
norgesplaster.no

Orgnr
556765-1756
VAT No:
SE556765175601

Common Specifications applied: No applicable CS available

Solna 2026-01-07

Johanna Brinck
Head of Regulatory and Quality
Person Responsible for Regulatory Compliance
Orkla Wound Care AB

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