

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.

Basic UDI-DI: 7340210004003B

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 | 1 |
| 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 | 1 |

Common Specifications applied: No applicable CS available

Solna 2023-03-13

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Person Responsible for Regulatory Compliance

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