

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: 73402100680059

EMDN (CND) code: M04010102

GMDN code: 44990

Intended purpose: The product is intended to be used as a mechanical barrier and for absorption of exudates from minor wounds.

REF	Name of product	Risk class
510003320	Salvequick X-Ray Detectable Plasters	I

Common Specifications applied:
No common specification available.

Solna 2024-04-22

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Johanna Brinck

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Person Responsible for Regulatory Compliance

Orkla Wound Care AB