

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 IIa medical devices and have been CE marked in accordance with Annex IX of MDR 2017/745.

Verified by Notified Body, Intertek Medical Notified body # 2862, certificate #: 28620123341.

Basic UDI-DI: 7340210102003C

EMDN (CND) code: M04040101

GMDN code: 61694

Intended purpose: The Wound cleanser is intended for mechanical cleansing of minor wounds in situations where clean tap water is not available.

REF	Name of product	Risk class
3239	Salvequick/Cederroth Wound Cleanser	IIa
323700	Salvequick/Cederroth Wound Cleanser 20	IIa
3227	Salvequick/Cederroth Wound Cleanser 40	IIa

Common Specifications applied: No applicable CS available.

Solna 2025-01-08

Johanna Brinck

[Johanna Brinck \(Jan 8, 2025 15:36 GMT+1\)](#)

Johanna Brinck

Head of Regulatory and Quality

Person Responsible for Regulatory Compliance

Orkla Wound Care AB

DoC Salvequick_Cederroth Wound Cleanser_MDR_EN

Final Audit Report

2025-01-08

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