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UK DECLARATION OF CONFORMITY

In accordance with UK Government Guidance

Manufacturer

UK Responsible Person

Etac Ltd

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Statement

This declaration of conformity is issued under the sole responsibility of the manufacturer.

The device(s) covered by this declaration of conformity is/are in conformity with EU

Regulation 2017/745 on medical devices.

Basic UDI-DI

570799524TM

Device description

Patient Slings

Intended Purpose

The Sling is an assistive device intended for alleviation of or compensation for a functional impairment due to an injury or disability. The device is designed for an individual lacking the ability to stand up and transfer oneself over shorter distances to another sitting position to/from a bed, a wheelchair, a chair, a toilet or similar, due to

reduced mobility or physical strength.

Device name(s)

Molift UnoSling Ambulating Vest Molift EvoSling Embrace Vest Molift RgoSling Ambulating Vest

Parts covered in this

declaration

Separate list available upon request.

Risk class of the device

Class I. Rule 1

Harmonized/Established

Standards

Separate list available upon request

Place

Gedved, Denmark

Date of issue

11. November 2025

Name and function

Sampavi Rajkumar, Quality Specialist

Signature, on behalf of Etac A/S