



EU Declaration of Conformity

Manufacturer

Etac A/S

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SRN

DK-MF-000017724

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

570799526TR

Device description

Accessories for patient slings

EMDN

Y1299

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 RgoSling Groin Straps Molift RgoSling Extension loops Molift EvoSling Soft padding hygiene

Molift Evosling Soft Padding Molift EvoSling Head support Molift EvoSling Extension Loops

Molift Strap Marker

Risk class of the device

Class I, Rule I

Harmonized/Established

Standards

Separate list available upon request

Place

Gedved, Denmark

Date of issue

20.March 2025

Name and function

Sampavi Rajkumar, Quality Specialist

Signature, on behalf of Etac A/S

nm. R.