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EU DECLARATION OF CONFORMITY

Manufacturer	**Etac A/S** Parallelvej 3 DK-8751 Gedved Denmark
SRN	DK-MF-000017724
Statement	This EU declaration of conformity is issued under the sole responsibility of the manufacturer. The device(s) covered by present declaration is/are in conformity with EU Regulation 2017/745 on medical devices.
Basic UDI-DI	**570799517TQ**
Device description	Patient slings
Intended purpose	The slings 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 lift a body part for a short time during examination, reposition, dressing, undressing or similar, due to reduced mobility or physical strength.
Device name(s)	Molift EvoSling FlexiStrap Molift UnoSling LimbLift
Brand	Molift
Risk class of the device	Class I, rule I
Place	Gedved, Denmark
Date of issue	24. August 2023
Name and function	Michael Bruun, Senior Vice President

Signature, on behalf of Etac A/S