UK DECLARATION OF CONFORMITY

In accordance with UK Government Guidance

Manufacturer	Etac A/S Parallelvej 3, 8751 Gedved, Denmark Phone no. + 45 79685833 www.etac.com	UK Responsible Person	Etac Ltd Unit 60, Hartlebury Trading Estate, Hartlebury, Kidderminster, DY10 4JD Email enquiries.uk@etac.com Web www.etac.com/uk Phone +44 (0)121 561 2222

Statement Basic UDI-DI	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	570799517TQ
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 EvoSling FlexiStrap Molift UnoSling LimbLift
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