# UK DECLARATION OF CONFORMITY

*In accordance with UK Government Guidance*

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

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| Statement<br>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 | 570799525TP |
| 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 RgoSling StandUp Padded<br>Molift RgoSling StandUp w/Support Padded<br>Molift RgoSling Active<br>Molift RgoSling Active Slim<br>Etac Nova StandUp<br>Molift UnoSling Standup |
| 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 | 08. September 2025 |
| Name and function | Sampavi Rajkumar, Quality Specialist |

Signature, on behalf of Etac A/S