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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	**570799516TN**
Device description	Patient slings
Intended purpose	For users who are in need of support to be lifted for a short time; e.g. people with limited ability to move and who require body support during transfers. Not an exhaustive list. The user group for the device is based on individual health and mobility function, and not on a specific diagnosis or age.
Device name(s)	Molift RgoSling Ampu MediumBack Padded Molift RgoSling Ampu HighBack Padded Molift EvoSling Ampu MediumBack Net Padded
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