



# EU DECLARATION OF CONFORMITY

<b>Manufacturer</b>	<b>Etac A/S</b> Parallevej 3 DK-8751 Gedved Denmark
<b>SRN</b>	DK-MF-000017724
<b>Statement</b>	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.
<b>Basic UDI-DI</b>	<b>570799510TA</b>
<b>Intended purpose</b>	Patient Lifting System intended for sitting and horizontal transfers as well as standing and gait training situations.
<b>Product / device name</b>	Molift AIR 200 Molift AIR 200 (Litium battery) Molift AIR 350 Molift AIR 350 Life Molift AIR 350 Propulsion Molift AIR 350 IRC Propulsion Molift AIR 350 IRC Molift AIR 350 Tilt
<b>Parts covered by this declaration</b>	Parts and components listed in the Instructions For Use for the mentioned devices.  As for the compatibility with other rail systems: Trolley 2510000 / 2520000 / 2520007 / 2520020 2520001 2520004 2520005 2520008 2520009 2520010 2520011 2520012 2520013 2520014 2520018 2520019  Rail manufacturer Molift Guldmann ABC / Invacare Liko Merivaara & Nova Ergolet BMH Hoyer /Guldmann mini Nova Freeway Oxford Voyager Spektra Chiltern Horcher
<b>Brand</b>	Molift
<b>Risk class of the device</b>	Class I
<b>Place</b>	Gedved, Denmark
<b>Date of issue</b>	11. January 2022
<b>Name and function</b>	Michael Bruun, Senior Vice President

Signature, on behalf of Etac A/S