




# UK DECLARATION OF CONFORMITY

In accordance with UK Government Guidance

<b>Manufacturer</b>	<b>Etac Immedia A/S</b> Parallevej 3 DK-8751 Gedved Denmark  <a href="http://www.etac.com">www.etac.com</a>	<b>UK Responsible Person</b>	<b>Etac Ltd</b> Unit 60, Hartlebury Trading Estate, Hartlebury, Kidderminster, DY10 4JD  Email <a href="mailto:enquiries.uk@etac.com">enquiries.uk@etac.com</a> Web <a href="http://www.etac.com/uk">www.etac.com/uk</a> Phone +44 (0)121 561 2222
<b>Statement</b>	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 UK MDR 2002 (Medical Device Regulations 2002)		
<b>Basic UDI-DI</b>	57080126001MM		
<b>Device description</b>	Special Solutions		
<b>GMDN</b>	37162		
<b>Intended purpose</b>	The assistive device is 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 transfer or position themselves due to reduced mobility or physical strength.		
<b>Device name(s)</b>	SlingOn		
<b>Risk class of the device</b>	Class I, rule I		
<b>Harmonized/Established Standards</b>	Separate list available upon request		
<b>Place</b>	Gedved, Denmark		
<b>Date of issue</b>	15. January 2024		
<b>Name and function</b>	Michael Bruun, Senior Vice President		

  
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Signature, on behalf of Etac Immedia A/S