immedia
by Etac

CE
MD

EU Declaration of Conformity

Manufacturer	**Etac Immedia A/S** Parallelvej 3 DK-8751 Gedved Denmark www.etac.com
SRN	DK-MF-000019241
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	57080126001MM
Device description	Special Solutions
EMDN	Y1299
Intended purpose	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.
Device name(s)	SlingOn
Risk class of the device	Class I, rule I
Harmonized/Established Standards	Separate list available upon request
Place	Gedved, Denmark
Date of issue	15. January 2024
Name and function	Michael Bruun, Senior Vice President

Signature, on behalf of Etac Immedia A/S