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EU DECLARATION OF CONFORMITY

Manufacturer	**Etac Immedia A/S** Parallelvej 3 DK-8751 Gedved Denmark
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	**57080125001ME**
Intended purpose	Transfer between wheelchair, chair, bed, car and similar. In/out of bed. Legs in/out of bed The Sling is an extended arm with handles. The Sling is used for moving the client higher up in the bed, back to a chair, to stand up and transfer between two sitting areas. The Sling has handles in each side.
Product / device name	Immedia Sling

Brand	Immedia
Risk class of the device	Class I
Place	Gedved, Denmark
Date of issue	1 June 2022
Name and function	Michael Bruun, Senior Vice President

Signature, on behalf of Etac Immedia A/S