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EU DECLARATION OF CONFORMITY

Manufacturer	Etac Immedia A/S Egeskovvej 12 DK-8700 Horsens Denmark
SRN	NA
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	57080126007MZ
Intended purpose	Intended use Emergency evacuations. Transfer out of bed, and slide out to safety.
Product / device name	Immedia RescueSheet

Brand	Immedia
Risk class of the device	Class I

Place	Horsens, Denmark
Date of issue	15 April 2021
Name and function	Michael Bruun, Senior Vice President

Signature, on behalf of Etac Immedia A/S