



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 57080126002MP

Intended purpose Transfer in bed. Transfer between 2 beds or operational theatre. Helping patients up from the floor.
The MiniBoard can be used for turning in bed, moving upwards in bed, and transfer from bed to operation table as well as for lifting from floor to bed/chair. It is supplied with head support which helps lifting or carrying the client in recumbent position.

Product / device name Immedia MiniBoard

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