



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 57080121011LM

Intended purpose The Multiglide can be used anywhere where it is beneficial to reduce friction at pressure points during manual handling: Turning users in bed, pulling them higher up in the bed, getting in and out of bed, etc.

Product / device name Immedia MultiGlide SG

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