



# 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** 57080121013LR

**Intended purpose** The Multiglide SinglePatientUse 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.  
Multiglide SinglePatientUse can only be used at one patient but several times until it gets soiled

**Product / device name** Immedia MultiGlide Single Patient Use

**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