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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	**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	Gedved, Denmark
Date of issue	1 June 2022
Name and function	Michael Bruun, Senior Vice President

Signature, on behalf of Etac Immedia A/S