



EU Declaration of Conformity

Manufacturer Etac Immedia A/S
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Denmark
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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 57080125003MJ

Device description Support

EMDN Y1299

Intended purpose The assistive device is intended for alleviation of, or compensation for, a functional impairment due to an injury or disability. The device is designed for an individual lacking the ability to transfer or position themselves due to reduced mobility or physical strength.

Device name(s) SmartMove

Risk class of the device Class I, rule I

Harmonized/Established Standards Separate list available upon request

Place Gedved, Denmark

Date of issue 15. January 2024

Name and function Michael Bruun, Senior Vice President

A handwritten signature in blue ink, appearing to be 'Michael Bruun', is written over a horizontal line.

Signature, on behalf of Etac Immedia A/S