EU Declaration of Conformity

The manufacturer:

Company: Pam Mobility s.r.l.

Address: Via Verdi, 39 - 42043 Gattatico (RE) - Italy

SRN: IT-MF-000027951

Declares, under its own and exclusive responsibility, that the device(s)

Code	Model	ID BD/RDM	BASIC UDI-DI
82700002	Rod lifts patient with trapezium	2322661/R	80557742082700002HS

Intended use: The device have been designed and built to be installed on PAM Mobility beds to help autonomous patient lift.
Environment of use: within healthcare and health facilities.
The device cannot be used in a potentially explosive or flammable atmosphere.
Personnel intended for use of the product: specialist operators and doctors.

Risk class: Class I (in accordance with Rule 1, Annex VIII of Regulation (EU) 2017/745)

It complies with the following European Union legislative acts:

(EU) 2017/745 — Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) no. 178/2002 and Regulation (EC) no 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC

The device is subject to the conformity assessment procedure provided for in article 52, section 7 of Regulation (EU) 2017/745

Gattatico,
27 april 2023

Managing Director
Andrea Muzzini

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