# 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 |
|---|---|---|---|
| 7285RT50 | Variable height stretcher | 2322457/R | 8055774207285RT50YC |
| 7285RT52 | Variable height stretcher | 2322460/R | 8055774207285RT52YG |
| 7285RT51 | Variable height stretcher | 2322462/R | 8055774207285RT51YE |
| 7285RT60 | Variable height stretcher | 2322464/R | 8055774207285RT60YF |
| 7285RT61 | Variable height stretcher | 2322466/R | 8055774207285RT61YH |

Intended use: The device is intended to be used exclusively as a stretcher for the transportation, diagnosis, treatment and monitoring of patients under the close supervision and surveillance of medical personnel.
The device cannot be used for inpatient purposes
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

PAM MOBILITY SRL
Via Verdi, 39
42043 GATTATICO (RE)
P.IVA 02429390350 - Tel. 0522 473859
e-mail: info@pammobility.com