



## EU DECLARATION OF CONFORMITY

We, undersigned GIMA S.p.A. (single registration number (SRN): IT-MF-000011004), with operational headquarters in Gessate (MI), Via Marconi 1, and registered office in Milan, Via Tommaso Grossi 2, acting as manufacturer of the medical device:

| Product and trade name                    | Product code | Basic UDI-DI              |
|-------------------------------------------|--------------|---------------------------|
| MILITARY TOURNIQUET - orange              | 25526        | 80232790000V90030000000VS |
| MILITARY TOURNIQUET - black               | 25527        |                           |
| GIMA TOURNIQUET – velcro                  | 25721        |                           |
| VELCRO TOURNIQUET                         | 25725        |                           |
| FAST TOURNIQUET - LATEX FREE - light blue | 25726        |                           |
| FAST TOURNIQUET – red                     | 25727        | 80232790000V900300CC0007Z |
| FAST TOURNIQUET - green                   | 25728        |                           |
| FAST TOURNIQUET – Daisy                   | 37500        |                           |
| FAST TOURNIQUET – Peace                   | 37501        |                           |
| FAST TOURNIQUET - Space                   | 37502        |                           |

intended purpose: intended to be used to temporarily block venous return during intravenous infusion or blood sampling

risk class I (not sterile), in accordance with the rule 1 set out in Annex VIII of the Regulation (EU) 2017/745 (MDR), declares, under its sole responsibility, that this device:

- complies with the Regulation (EU) 2017/745 (MDR);
- Common Specifications have not been used for the compliance of the above medical device.

Gessate, 19/09/2025

**GIMA S.p.A.**  
The legal Representative  
(Nicola Manzoni)

A handwritten signature in black ink, appearing to read 'N. Manzoni', is written over a horizontal line.