



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	
FAST TOURNIQUET - green	25728	80232790000V90030000000VS
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, 27/07/2026

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.