



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
DECONTAMINATING MAT 45x90 cm - 30 layers - blue	36685	802327900T03050160000003E
DECONTAMINATING MAT 60x90 cm - 30 layers - blue	36686	802327900T03050160000003E
DECONTAMINATING MAT 45x115 cm - 30 layers - blue	36693	802327900T03050160000003E
DECONTAMINATING MAT 60x115 cm - 30 layers - blue	36694	802327900T03050160000003E
DECONTAMINATING MAT 90x115 cm - 30 layers - blue	36695	802327900T03050160000003E
DECONTAMINATING MAT 45x115 cm - 30 layers - white	36696	802327900T03050160000003E

intended purpose: protecting the patient from the risk of contamination and pathologies

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, 24/09/2025

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.