GIMA S.p.A.
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ITALIAN DIVISION
gima@gimaitaly.com
EXPORT DIVISION
export@gimaitaly.com

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
KINESIOLOGY TAPE 5 m x 5 cm - black	34745	8023279M0304029969000000L4
KINESIOLOGY TAPE 5 m x 5 cm - blue	34746	8023279M0304029969000000L4
KINESIOLOGY TAPE 5 m x 5 cm - red	34747	8023279M0304029969000000L4
KINESIOLOGY TAPE 5 m x 5 cm - green	34749	8023279M0304029969000000L4
KINESIOLOGY TAPE 5 m x 5 cm - pink	34750	8023279M0304029969000000L4
KINESIOLOGY TAPE 5 m x 5 cm - skin	34751	8023279M0304029969000000L4
KINESIOLOGY TAPES 5 m x 5 cm - mix colours	34752	8023279M0304029969000000L4
KINESIOLOGY TAPE PRO 5 m x 5 cm - skin	34757	8023279M0304029969000000L4

Intended purpose: used to treat muscular symptoms, to stimulate blood circulation and in cases of nerve and lymph shunts.

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, 30/06/2025

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