GIMA S.p.A.
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ITALIAN DIVISION
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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
KILLIAN NOSE FORCEPS 50 mm, 14 cm	26776	802327900L1402020000000MP
FIBRE OPTIC KILLIAN NOSE FORCEPS 75 mm, 14 cm	26777	802327900L1402020000000MP
KILLIAN NOSE FORCEPS 22 mm/14 cm	26778	802327900L1402020000000MP

intended purpose: Used to keep the nostrils open, allowing the nasal cavity to be examined and other instruments inserted for rhinoscopy.

risk class I (not sterile), in accordance with the rule 5 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, 23/01/2025

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