GIMA S.p.A.
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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 | Intended purpose | Rule |
|---|---|---|---|---|
| PROBE - 15 cm | 26701 | 802327900L0311995200000ZE | intended to be used to explore cavities, wounds or fistulas during abdominal, ENT and/or orthopedic surgery | 1 |
| BUTTERFLY PROBE- 14.5 cm | 26702 | 802327900L0311995200000ZE | Used in dissections to protect organs from the scalpel blade or scissors | 1 |
| UNNA OVAL - 14 cm | 26820 | 80232790000L24995200000UK | Skin probe | 6 |
| UNNA ROUND - 14 cm | 26821 | 80232790000L24995200000UK | Skin probe | 6 |
| FORCEPS ROUND - 9 cm | 26823 | 8023279L0313040200000002W | Skin clamp | 1 |
| KATSCH - 14 cm | 26825 | 80232790000L24995200000UK | Skin probe | 6 |
| GILLES HOOK - 13 cm | 26826 | 8023279L0314010552000006L | Skin probe | 5 |

risk class I (not sterile), in accordance with the rules 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, 30/06/2026

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

