



## 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
SET OF UTERINE SUCTION CANNULAS (7 cannulas+1 handle)	28260	8023279A0601010312CC000SD
S/S BIERER ASPIRATION TUBE diam. 8 mm	28262	
S/S BIERER ASPIRATION TUBE diam. 12 mm	28264	

intended purpose: allow the suction or drainage of biological fluids from the uterus

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, 15/07/2025

**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.