EU DECLARATION OF CONFORMITY

METRICA SPA (SRN code: IT-MF-000028794) with operational headquarters at Via Grandi 18, 20097 San Donato Milanese (MI), registered office at Viale Vicenza 40, 36071 Arzignano (VI) and production site at Via Santo Stefano 16, 32040 Danta di Cadore (BL), as the manufacturer of medical devices:

Trade names / Models	Items	Base UDI-DI
Analogue Baby Measuring rod with caliper	10455	8001066 CALNEO FZ
Analogue Orthopaedic caliper	10460	8001066 CALORTO JG
Analogue Craniometer	70410	8001066 CRANIOM KD
Analogue Skinfold Caliper	70200 70209	8001066 PLICANALOG N6
Analogue Device for Lymphedema	70420	8001066 DMLINF L2

intended for measuring anatomical parameters for anthropometric assessments by a physician; risk class Im with measuring function, in accordance with Rule 1 of Annex VIII of EU Regulation 2017/745,

declares under its sole responsibility that these devices:

- comply with the general safety and performance requirements and the provisions of Regulation (EU) 2017/745 as per the Technical File filed with the body;
- no Common Specifications have been used for the conformity of the devices;
- are manufactured in accordance with Technical File no. DT 002 ME, which meets the requirements of Part XI A of Regulation (EU) 2017/745, as per Certificate no. 0425-MDR-030059-00 issued on 12/12/2023 by ICIM S.p.a. Sesto San Giovanni (MI) in Piazza Don Enrico Mapelli, 75, Notified Body no. 0425.

Arzignano (VI),
26/02/2026

METRICA SPA
Legal Representative
(Mario Doriguzzi Bozzo)