

Warsaw, 25.05.2021

EU DECLARATION OF CONFORMITY MEDICAL DEVICE

Irena Groniecka - Tarnkowska, Andrzej Tarnkowski "ANTAR" Spółka Jawna ul. Zawiślańska 43 03-068 Warszawa

acting as Manufacturer registered in Eudamed, SRN number: PL-MF-000001583, according to Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017, herein declare that medical devices:

ORTHOPAEDIC SEPARATION PILLOW: AT03018

The Basic UDI-DI: not issued

have been classifed as medical device class I, rule 1.

Intended purpose: orthopedic pillow between the knees helps to ensure the correct, anatomical positioning of the hips and pelvis while lying on the side, e.g. in the process of rehabilitation after surgery. It reduces pain caused by incorrect positioning of the hips when lying on your side. It reduces the tension in the knee joints.

We herein declare that the medical devices are in conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017. The assessment of conformity has been proceeded in compliance with the requirements of the above Regulation.

Applied standards:

PN-EN ISO 9001 (current edition)

PN-EN ISO 13485 (current edition)

PN-EN ISO 15223 (current edition)

PN-EN ISO 14971 (current edition)

PN-EN ISO 10993 (current edition)

The EU declaration of conformity is issued under the sole responsibility of the Manufacturer.

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co-owner independent reprezentation of the company based on the Company Register





