

Warsaw, 25.05.2021

EU DECLARATION OF CONFORMITY MEDICAL DEVICE

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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:

ANTI-BEDSORE MATTERES: AT03103

The Basic UDI-DI: not issued

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

Intended purpose: the anti-bedsore foam mattress allows the optimal distribution of pressure exerted by the body on the tissues adhering to the substrate and thus slows down the process of blood vessel clogging, leading to the formation of pressure ulcers. The mattress is recommended for people with severely limited mobility: spending most of their time lying down

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 representation of the company
based on the Company Register