



EU DECLARATION OF CONFORMITY

ANTAR MEDICAL Limited Liability Company
ul. Zawisłańska 43; 03-068 Warsaw, Poland
SRN: PL-MF-000001583

We hereby declare, under our sole responsibility, that the medical device covered by this declaration of conformity complies with all applicable requirements of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices (MDR).

MEDICAL DEVICES:

AT51134 ROLLATOR - Basic UDI-DI: 59015714AT5113443

AT51135 ROLLATOR - Basic UDI-DI: 59015714AT5113545

AT51136 ROLLATOR - Basic UDI-DI: 59015714AT5113647

AT51137 ROLLATOR - Basic UDI-DI: 59015714AT5113749

The device has been classified in accordance with Annex VIII of the MDR as a **Class I medical device, Rule 1**

Intended purpose: the product is intended to provide support for people with limited walking ability during movement. The rehabilitation rollator is designed for one person.

The conformity assessment was carried out according to Annex II and III of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, with the application of the following standards:

PN-EN ISO 15223-1:2022

PN-EN 20417:2021

PN-EN ISO 14971:2020

PN-EN ISO 13485:2016

PN-EN ISO 11199-2:2005

PN-EN ISO 10993-1:2021

PN-EN 62366-1:2015-07/A1:2021

The EU Declaration of Conformity has been issued under the sole responsibility of the manufacturer.



Andrzej Tarnkowski
President of the Management Board

Warsaw, 24.03.2026