



## 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: ROLLATOR AT51029

Basic UDI-DI: 59015714AT5102945

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

**Intended purpose:** rollators are designed for disable people to enable them to move. User rests on the rollator, makes a step forward and then pushes the rollator.

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 ISO 11199-2:2005

PN-EN 20417:2021

PN-EN ISO 10993-1:2021

PN-EN ISO 14971:2020

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

PN-EN ISO 13485:2016

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