Declaration of Conformity

1. Legal Manufacturer

Easy Roller AS
Hagebyveien 40
3734 Skien
Norway

2. Production Site

ROTO GROUP d.o.o.
Gorička 150, Černelavci
9000 Murska Sobota
Slovenia

3. Device

- EasyRoller — REF: ER
  Basic UDI-DI: 709006211EASYROLLER2R
  - Intended use:
    By being metal free, the EasyRoller wheelchair will not affect nor be affected by magnetic resonance imaging´s magnetic fields or metal detectors. Using EasyRoller porters will be able to transport PRM throughout the entire hospital environment without any restrictions. Passing of security gates is done without any intimidating manual control of the person with reduced mobility.

- EasyRoller Pool — REF: ERP
  Basic UDI-DI: 709006211EASYROLLER2R
  - Intended use:
    The person with reduced mobility can be transported from the reception area through showers and with the chair submerged in the swimming pool the person with reduced mobility can do exercising in water with the wheelchair at hand. The EasyRoller is not affected by any use in a Sauna environment.

4. Classification

EasyRoller and EasyRoller Pool are classified as Class 1 Medical Device according to MDR Regulation (EU) 2017/745 Annex VIII, Rule 1.

5. Conformity Assessment

MDR, Regulation (EU) 2017/745, Art. 19, Annex IV & V.

6. EC Representative

N/A

7. Single Registration Number

NO-MF-000010301

8. GMDN Code

58086

9. EORI Number

DE1807838

We, the manufacturer, declare herewith under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices. All supporting documentation is retained on the premises of the manufacturer.

Place:	Date:	Signature of Chief Executive Officer:
Skien	25/11/2025	EasyRoller