EU – Declaration of Conformity

We, A2J GmbH, Am Fuchsberg 13 - D 87452 Altusried, represented by Armin Janusch, CEO, declare under our sole responsibility that the medical device group

Crazy Horse Active Power wheelchair pulling aid

Basic UDI-DI: **426072548 1302090225 8N**

complies with the relevant provisions of Regulation (EU) 2017/745 (MDR) on medical devices and, where applicable, other relevant Union legislation

The product group **Crazy Horse Active Power wheelchair pulling aid** includes the following products	
Crazy Horse Active Power	all model versions

The intended use of the **Crazy Horse Active Power wheelchair pulling aid** is:
Manual wheelchair pulling aid for outdoor activities

According to Annex VIII, Rule 1 MDR, all devices of the **Crazy Horse Active Power wheelchair pulling aid** product group are class I medical devices and the applicable essential safety and performance requirements according to Annex I MDR are fulfilled. Technical documentation in accordance with Annex II & III MDR is maintained.

Applicable harmonized standards, national standards, or other regulatory documents:

DIN EN ISO 13485 – Medical devices – Quality management systems – Requirements for regulatory purposes
DIN EN ISO 14971 – Medical devices – Application of risk management to medical devices
DIN EN 12183 – Manual wheelchairs – Requirements and test methods
DIN EN ISO 10993-1 – Biological evaluation of medical devices - **DIN EN ISO 10993-5** – In vitro Cytotoxicity

This EU Declaration of Conformity applies in conjunction with the delivered product

Altusried, the **27.04.2025**

Armin Janusch
CEO A2J GmbH

Manufacturers SRN: **DE-MF-000008341**