

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

PELVI.LOC® Positioning and Retraction System

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

The **PELVI.LOC®** System includes the following products

pelvi.loc Basic UDI-DI: 426072548 01650120229 D2			torso.loc Basic UDI-DI: 426072548 20015018019015 Z5	foot.loc Basic UDI-DI: 426072548 6015015020 AX
PL-2DB	PL-3DSR	PL-CSS	PL-W	PL-KG-KS
PL-2DS	PL-3DSRR	PL-UG	PL-WR	PL-KG-ZR
PL-2DA	PL-3DA	PL-UGR	PL-OPS	PL-KG-GO
PL-2DAS	PL-3DAS	PL-GF	PL-OPG-FLEX	PL-RR
PL-3DB	PL-3DASR	PL-3DWP	TL-3DSR	PL-RR-GO
PL-3DS	PL-ASY	RT-2DS	TL-W	

The intended use of the **PELVI.LOC®** System is: Positioning and retraction for persons with limited mobility e.g. wheelchair, seating shell, therapy chair, rehab buggy, sports equipment, standing device.

According to Annex VIII, Rule 1 MDR, all devices of the **PELVI.LOC®** System are class 1 medical devices and the applicable essential safety and performance requirements according to Annex I MDR are fulfilled. A conformity assessment procedure according to Annex II & III MDR has been carried out.

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 – 8.5

DIN EN 12184 – Electrically powered wheelchairs, scooters and their chargers – Requirements and test methods - 9.5

DIN EN ISO 10993-1 – Biological evaluation of medical devices - **DIN EN ISO 10993-5** – In vitro Cytotoxicity

This EU Declaration of Conformity is valid until **25.05.2024**

Altusried, the 20th of May 2023

Armin Janusch

Armin Janusch
CEO A2J GmbH

Manufacturers SRN: DE-MF-00008341

Version 1.1	Erstellt von: TC	Freigegeben von: AJ – 20.05.2023	Qualitätsmanagementsystem nach EN ISO 13485		
Datei: A2J CE KE-ENol PL 05-23		Anlage: 11.03.2021	Stand: 20.05.2023	Seite 1 von 1	
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