

## 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

### **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

<b>pelvi.loc</b> Basic UDI-DI: <b>426072548 01650120229 D2</b>			<b>torso.loc</b> Basic UDI-DI: <b>426072548 20015018019015 Z5</b>	<b>foot.loc</b> Basic UDI-DI: <b>426072548 6015015020 AX</b>
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 **01.01.2025**

Altusried, 25.05.2024

*Armin Janusch*

Armin Janusch  
CEO A2J GmbH

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

Version 1.1	Erstellt von: TC	Freigegeben von: AJ – 25.05.2024	Qualitätsmanagementsystem nach EN ISO 13485		
Datei: A2J CE KE-EN PL 05-24.docx		Anlage: 11.03.2021	Stand: 25.05.2024	Seite 1 von 1	
Firma A2J GmbH – Am Fuchsberg 13 – D 87452 Altusried			© Castner Consulting – Medizinische Systemberatung		