EU DECLARATION OF CONFORMITY

Manufacturer: PROMA REHA, s.r.o.

Headquarters: Riegrova 342, 552 03 Česká Skalice, Czech Republic

SRN: CZ-MF-000013860

declares under his sole responsibility that

Products:	**Lever beds**
Made in types:	**GENEO, GRANO, SUPERTA E, SUPERTA H, SUPERTA M, DOMEO E, DOMEO H, DOMEO M, ABELA, SPARO, VERTICO** with applicable additional equipment
with basic UDI-DI:	**859420232LP01**

are in comply:

- Regulation of the European Parliament and of the Council (EU) 2017/745.
- Act No. 375/2022 Coll. on Medical Devices and In Vitro Diagnostic Medical Devices.
- Government Regulation No. 481/2012 Coll., Restrictions on the Use of Certain Hazardous Substances in Electrical and Electronic Equipment.
- Act No. 22/1997 Coll. on Technical Requirements for Products and on Amendments to Certain Acts.

Medical device classification according to paragraph 4.1 of Chapter III, Annex VIII of Regulation (EU) 2017/745 of the European Parliament and of the Council is product **Class I**, non-sterile, without measuring function.

In Česká Skalice

Date of last revision 22. 12. 2022

Digitally signed by Botnaru Andrei
Date: 2026.03.23 11:57:12 EET
Reason: MoldSign Signature
Location: Moldova
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Radek Jakubský
Managing director

PROMA REHA, s.r.o.