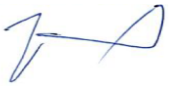
EU Declaration of Conformity

Manufacturers Name:	Ferno Slovakia s.r.o.
Manufacturers Address:	Bošáca 893 / 913 07 / Slovakia
SRN (Single Registration Number):	SK-MF-000001474
Authorized Representative (UKCA REPRESENTATIVE)	Ferno House Stubs Beck Lane, Cleckheaton, West Yorkshire, BD19 4TZ United Kingdom +44 (0) 1274 851999 www.ferno.co.uk
Basic UDI-DI:	8588008621LIFTERFASTXH
Name of the Device (s):	FAST-LIFT FAST-LIFT X
Product code:	60-0254-001 60-0254-002
Classification:	Medical Device Accessory
Notified Body name:	N/A
Notified Body Address:	N/A
Notified Body Identification number:	N/A
Conformity assessment route:	Ferno Slovakia s.r.o. uses the following procedures for the CE-labeling of their products according the Regulation MDR 2017/745. Class I: EU conformity declaration according to Annex IV + Annex VIII of MDR 2017/745 This DoC is supported by a quality management system certified according to ISO 13485. This Declaration of Conformity is issued by Ferno Slovakia s.r.o.

We hereby declare that the medical device accessory(ies) specified above meet the provisions of the Regulation (EU) MDR 2017/745 for medical devices.

All supporting documentation is retained at the premises of the manufacturer.

Signature:



Silvia Vančová
Managing Director

Place and date of issue:

Bošáca, 2.4.2026

Attachment nr.1 to declaration of conformity FAST-LIFT

FAST-LIFT

Product Number	Product description	GTIN (UDI-DI)
60-0254-001	FAST-LIFT X	8588008621912
60-0254-002	FAST-LIFT	8588008621943