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 1 (EU AUTHORIZED REPRESENTATIVE)	Ferno S.r.l. via B.Zallone, n.26,40066 Pieve di Cento, Bologna, Italy +39.051.6860028 www.ferno.it
Authorized Representative 2 (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:	8588008129F2LockingNT
Name of the Device (s):	F2 LOCKING DEVICE
Product code:	60-0150-003
Classification:	Accessories
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: EN 1789 Class I: N/A – Accessories

This declaration of conformity is issued under the sole responsibility of Ferno Slovakia s.r.o. We hereby declare that he medical device(s) specified above meet the provision 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, 27.7.2022

Attachment nr.1 to declaration of conformity F2 LOCKING DEVICE

F2 LOCKING DEVICE

Part Number	Product description	GTIN (UDI-DI)
60-0150-003	F2 LOCKING DEVICE	8588008621721