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:	8588009265NATOSTRETCHER2Q
Name of the Device (s):	STANAG 2040 4-FOLD NATO STRETCHER w/TW
Product code:	60-0122-002
Classification:	Class I
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. Ferno Slovakia s.r.o. declares the products conform with STANAG 2040 (except chapter 2 road-worthiness). Class I: EU conformity declaration according to Annex IV + Annex VIII of MDR 2017/745 This DoC is issued based on the ISO 13485 Certification.

This declaration of conformity is issued by Ferno Slovakia s.r.o.
We hereby declare that the medical device(s) 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, 8.7.2024

Attachment nr.1 to declaration of conformity NATO STRETCHER

STANAG 2040 4-FOLD NATO STRETCHER

Part Number	Product description	GTIN (UDI-DI)
60-0122-002	STANAG 2040 4-FOLD NATO STRETCHER w/TW	8588009265559

Related Accessories (if applicable):

Part Number	Product description	GTIN (UDI-DI)
65-0440-001	WHEELED STRETCHER UNDERCARRIAGE	8588009265542