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EU DECLARATION OF CONFORMITY

MANUFACTURER				
Name of Company and Address		EUDAMED SRN / Application ID		
www.ferno.it	FERNO S.r.I Via B. Zallone n.26 40066 Pieve di Cento (BO) Italy +39 (051) 6860028	IT-MF-000031330 / APP000027477		
SWISS AUTHORIZED REPRESENTATIVE AND IMPORTER				
Name of Company and Address		Swiss Single Registration Number (CHRN)		
CH REP www.ferno-schweiz.ch	FERNO S.r.I Pieve di Cento, succursale di Savosa Via Tesserete, 67 6942 Savosa, Switzerland +41 (41) 259 60 00	CHRN-AR-20002332 - AUTHORIZED REPRESENTATIVE CHRN-IM-20002288 - IMPORTER		

The manufacturer declares under its own responsibility that the medical device(s):

PRODUCT IDENTIFICATION				
Product Brand Name		Photo		
FERNO, TSL Expander		* CEED		
EMDN		FENILOFI		
V0880 - MEDICAL SUPPORT EQUIPMENT - ACCESSORIES			Cana.	
Intended Purpose		ERNILLOFE		
TSL Expander is an accessory to the ScoopEXL stretcher for the		Scoop Expander Kit		
movement and transport of large sized patients.		OFENO		
REF (Item / Catalog)	Item Description	GTIN (UDI-DI)	GMN (Basic UDI-DI)	
25-0030-002	TSL Expander for ScoopEXL (MANU00090)	08051380870310	805138087V0880TSLCN	
25-0030-003	TSL Expander for ScoopEXL 1 piece	08051380870327	805138087V0880TSLCN	
RISK CLASS FOR MEDICAL DEVICES				
Device Classification	Common Specifications			
Class I Rule 1	Not applicable			

according to:

HARMONIZED AND NON-HARMONIZED STANDARDS		
Item	Description	
EN 1865-1:2010+A1:2015	Patient handling equipment used in road ambulances - Part 1: General stretcher systems and patient handling	
	equipment.	
EN ISO 14971:2019/A11:2021	Medical devices - Application of risk management to medical devices (ISO 14971:2019)	
EN ISO 15223-1:2021	Medical devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General	
	requirements (ISO 15223-1:2021)	
EN ISO 13485:2016+A11:2021	Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)	
EN ISO 9001:2015	Quality management systems - Requirements (ISO 9001:2015)	

complies with the essential requirements listed in Annex I of the Regulation (EU) 2017/745 concerning Medical Devices.

Pieve di Cento, July 22, 2023

Signature Enrico Carletti - Managing Director, PRRC

This document is compiled in accordance with Annex IV - EU declaration of conformity

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