EU DECLARATION OF CONFORMITY

MANUFACTURER	
Name of Company and Address	**EUDAMED SRN / Application ID**
FERNO Australia Pty Ltd 11 Johnstone Road, Brendale Queensland, 4500 Australia +61 7 3881 4999 www.ferno.com.au	AU-MF-000035726 / APP000052128
EU AUTHORIZED REPRESENTATIVE AND IMPORTER	
Name of Company and Address	**EUDAMED SRN / Application ID**
FERNO S.r.l Via B. Zallone n.26 40066 Pieve di Cento (BO) Italy +39 (051) 6860028 www.ferno.it	**IT-AR-000031265**/APP000028644
UK RESPONSIBLE PERSON AND IMPORTER	
Name of Company and Address	**MHRA Reference Number**
FERNO (UK) Ltd, Ferno House, Stubs Beck Lane, Cleckheaton, West Yorkshire, BD19 4TZ +44 (0) 1274 851999 www.ferno.co.uk	1270

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

PRODUCT IDENTIFICATION			
Product Brand Name		**Photo**	
KED Pro			
EMDN			
V08050101 GENERIC USE STRETCHERS			
NATO NUMBER (NSN)			
Intended Purpose			
The KED Pro is an emergency patient-handling device designed to aid in the immobilisation, short transfer movement and technical rescue of patients with suspected spinal/cervical injuries.			
REF (Item / Catalog)	**Item Description**	**GTIN (UDI-DI)**	**GMN (Basic UDI-DI)**
FWE125-2	Ferno #125 Ked PRO	09348498002153	93484980FWE125SF
BRB-KED	KED PRO Adjustable Lifting Bridle	09348498002221	93484980FWE125SF

RISK CLASS FOR MEDICAL DEVICES	
Device Classification	**Common Specifications**
Class I Rule 1	Not applicable

according to:

HARMONIZED AND NON-HARMONIZED STANDARDS	
Item	**Description**
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.

Robert Hall
National Quality and Compliance Manager
Perth, Australia
29/05/2023

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

Ferno Australia head Office | 11 Johnstone Road, Brendale, Queensland 4500, AUSTRALIA