



Declaration of Conformity - CE

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER			
Name of Company	Address	SRN	Telephone/Email
Ferno-Washington, Inc. 	70 Weil Way, Wilmington, OH 45177-9371 U.S.A.	US-MF-000041603	+1 (937) 382-1451 fernoorders@ferno.com

EU AUTHORIZED REPRESENTATIVE			
Name of Company	Address	SRN	Telephone/Email
Ferno S.r.l. 	Via B. Zallone n.26 40066 Pieve di Cento (BO) Italy	IT-AR-000031265	+39 (051) 6860028 www.ferno.it

PRODUCT IDENTIFICATION	
Product Brand Name	Photo
POWER F2	
EMDN	
V0880	
MEDICAL SUPPORT EQUIPMENT - ACCESSORIES	
Intended Purpose	
An accessory to a Wheeled Stretcher is designed to assist in loading, unloading and securing compatible stretchers inside a ground-based ambulance.	

REF (Item / Catalog #)	Item Description	GTIN (UDI-DI)	GMN (Basic UDI-DI)
0567091	POWER F2 WITH POWERDRIVE (configuration of 0567093 Carriage, 0567095 Rail, 0567096 Connect)	00190790006570	0190790V0880K5

Related Accessories (if applicable):

Item / Catalog #	Item Description	GTIN (UDI-DI)	GMN (Basic UDI-DI)
n/a	n/a	n/a	n/a

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 14971:2019	Medical devices – Application of risk management to medical devices
ISO 15223-1:2021	Medical devices – Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements
ISO 13485:2016	Medical devices – Quality management systems – Requirements for regulatory purposes
EN 1865-2:2024	Patient Handling Equipment used in Road Ambulances – Part 2: Power Assisted Stretcher
EN 1865-3:2012+A1:2015	Patient Handling Equipment used in Road Ambulances – Part 3: Heavy Duty Stretcher
BS EN 1865-5:2012	Patient Handling Equipment used in Road Ambulances – Part 5: Stretcher Support
EN 1789:2020+A1:2023	Medical Vehicles and their Equipment – Road Ambulances

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

COMPANY REPRESENTATIVE:

DAWN M. GREENE

TITLE: DIR, REGULATORY

PLACE: Wilmington, OH, USA

SIGNATURE:

DATE: July 15, 2026

DoC Revision: 01

Record History:

Revision	Summary of Changes	Author	Effective Date
00	Initial issue	D. Greene	12/31/2025
01	Addition of EN 1865-5 compliance	D. Greene	07/15/2026