



DECLARATION OF CONFORMITY Regulation (EU) 2017/745

Manufacturer: FERNO S.r.l.
Manufacturer's address: Via B. Zallone n.26 – 40066 Pieve di Cento (BO) - Italy
Single Registration Number: (available when the managing system will be implemented by the
(Art.31(2)) European Commission)

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

Product code	Name of Device	Class (Annex VIII)
SLAM-26	SLAM for stretchers 26-S and 26P	I
SLAM-26+	SLAM for stretcher 26-S (with update)	I
SLAM-26/PAS	SLAM for stretcher 26-S on PAS plan	I
SLAM-26/PAS+	SLAM for stretcher 26-S on PAS plan (with update)	I
SLAM-26P/PAS	SLAM for stretcher 26-P on PAS plan	I
SLAM-5026/EFX	SLAM for stretchers 5026, EFX, CUSTOM-M	I
SLAM-4050	SLAM for stretchers 4051 e 4052	I
21-0180-006	Multi Chair Fastener	I
21-0086-001	VENICECLACK	I
25-1000-020-UK	VENICE SIMPLE CLACK	I
65-0430-001	EZCLACK fastener for chair 59-T	I
561-O	Horizontal fastener for stretcher Scoop 65EXL	I
561-V	Vertical fastener for stretcher Scoop 65EXL	I

Annex applied for the CE marking: Annex II and Annex III
Basic UDI-DI 805138087BLCG00678
Intended use: Locking system for devices in ambulance

In accordance with the provisions of harmonized and non-harmonized standards:


EN 1789:2007 + A2:2014 Medical vehicles and their equipment – Road ambulances



FERNO

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complies with the essential requirements listed in Annex I of the European Regulation 2017/745 concerning Medical Devices

Ferno maintains a Quality Management System that fulfills the requirements of ISO 13485:2016. Copies of Ferno's ISO 13485:2016 certificate issued by DNV is available upon request.

Pieve di Cento, May 28th 2021

Signature
Enrico Carletti - Managing Director

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

V: 2021.04_ENG