EU Declaration of conformity no. 200828-025

Product Name:	Vacuum Splint Kit AS 190 FW
Intended use:	The vacuum splints are intended for use in pre-hospital and hospital settings by professionally trained emergency and healthcare personnel to provide stabilisation and support of injured patient extremities during patient handling and transport. The vacuum splints are intended for use in patients with hand, arm, ankle and leg injuries. Including: Vacuum Splint AS 100 Vacuum Splint AS 120 Vacuum Splint AS 140 Vacuum Hand pump Repair Kit Bag for Splints Instructions for use
SRN: **Basic UDI-DI:** **UDI DI:** **Germa Article No:**	SE-MF-000003932 735001959P02VACSPLIGQ 07350019591925 23005002210
Manufacturer: **Visiting address:** **Phone:** **Email:** **Web:**	AB Germa Industrigatan 54-56, SE-29136 Kristianstad +46 (0)44 123030 info@germa.se www.germa.se
UKCA Representative UK CA	Ferno (UK) Limited Ferno House, Stubs Beck Lane, Cleckheaton, West Yorkshire BD19 4TZ, England Telephone: +44 (0) 1274 851999, www.ferno.co.uk
Product class:	Class I according to rule 1 in Annex VIII in MDR 2017/745
Conformity procedure:	Self-certification according to Annex IV in MDR 2017/745
Identification:	All products with serial numbers issued from; LOT number: 517470 Date: 2021-05-25 (yyyy-mm-dd).

Declaration statement:
This EU declaration of conformity is issued under the sole responsibility of the manufacturer AB Germa. The devices covered by this declaration is in conformity with the requirements in the European Medical Device Regulation 2017/745.

Signed in Kristianstad, Sweden on behalf of AB Germa by:

Position: Managing Director
Name: Patrik Tornström
Date: 2026-03-17
Sign: 

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