EU Declaration of conformity no. 200828-025

Product Name:	**Vacuum Splint Kit AS 190 FW**
Intended use:	**The Vacuum Splint Kit are primarily intended to be used in prehospital and hospital environment, by professional trained emergency and hospital personnel to safely stabilise injured patient extremities during transport. The Vacuum Splints are suited for fixation of patients with hand, arm, ankle, leg injuries.** **Including: Vacuum Splint AS 100** **Vacuum Splint AS 120** **Vacuum Splint AS 140** **Vacuum Hand pump** **Repair Kit** **Bag for Splints** **Directions 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: Björn Holmqvist
Date: 2021-05-25

Sign: 

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