EU Declaration of conformity no. 240830-001

Product Name:	Vacuum Splint Set Blue Line FLAF
Intended use:	The Vacuum Splint Set is 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 arm, hand, leg, ankle and foot injuries. Including: Vacuum splint Full Leg Blue Line Vacuum Splint Arm Blue Line Vacuum Splint Forearm Blue Line Vacuum Hand pump Repair Kit Bag for Directions for use
SRN: **Basic UDI-DI:** **UDI DI:** **Germa Article No:**	SE-MF-000003932 735001959P02VACSPLIGQ 07350019597200 21272708000
Manufacturer: **Visiting address:** **Phone:** **Email:** **Web:**	AB Germa Industrigatan 54-56, SE-29136 Kristianstad +46 (0)44 123030 info@germa.se www.germa.se
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: 531078 Date: 2024-08-16 (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: 2024-08-30

Sign: _____

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