AB GERMA®

EU Declaration of conformity no. 200828-002

Product Name:	**Vacuum splint AS 120**
Intended use:	The vacuum splint 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 splint are suited for fixation of patients with arm injuries.
SRN: **Basic UDI-DI:** **UDI DI:** **Germa Article No:**	**SE-MF-000003932** **735001959P02VACSPLIGQ** **07350019591130** **14005002013**
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: 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: 

Postadress/Postal address	Adress/Goods address	Telefon/Phone	Telefax/Fax	E-mail/Web page	VAT No.	Bankgiro Bank account
AB GERMA Box 2127 SE-291 02 KRISTIANSTAD SWEDEN	Industrigatan 54-56 SE-291 36 Kristianstad	+46 (0)44-123030	+46 (0)44-10 31 79	info@germa.se www.germa.se	SE556086941301	481-4125