



AB GERMA®

EU Declaration of conformity no. 200828-001

Product Name:	Vacuum splint AS 100
Intended use:	The vacuum splint 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 splint is suited for fixation of patients with hand, arm injuries.
SRN: Basic UDI-DI: UDI DI: Germa Article No:	SE-MF-000003932 735001959P02VACSPILIGQ 07350019591123 14005002012
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

AB GERMA
Box 2127
SE-291 02 KRISTIANSTAD
SWEDEN

Adress/Goods address

Industrigatan 54-56
SE-291 36 Kristianstad

Telefon/Phone

+46 (0)44-123030

Telefax/Fax

+46 (0)44-10 31 79

E-mail/Web page

info@germa.se
www.germa.se

VAT No.

SE556086941301

Bankgiro

Bank account

481-4125