



# AB GERMA®

## EU Declaration of conformity no. 210319-016

<b>Product Name:</b>	<b>Transfer Mattress 12 Handles FNO Black</b>
<b>Intended use:</b>	Germa Transfer mattresses are primarily intended to be used in prehospital, and hospital environment, by professional trained emergency and hospital personnel, as a comfortable mattress on the stretcher during transportation, or/ and safely facilitate patient handling and the transfer of the patient e.g. from one stretcher to another.
<b>SRN:</b> <b>Basic UDI-DI:</b> <b>UDI DI:</b> <b>Germa Article No:</b>	<b>SE-MF-000003932</b> <b>735001959P03TRAMATTL9</b> <b>07350019593837</b> <b>24271008800</b>
<b>Manufacturer:</b> <b>Visiting address:</b> <b>Phone:</b> <b>Email:</b> <b>Web:</b>	<b>AB Germa</b> <b>Industrigatan 54-56, SE-29136 Kristianstad</b> <b>+46 (0)44 123030</b> <b><a href="mailto:info@germa.se">info@germa.se</a></b> <b><a href="http://www.germa.se">www.germa.se</a></b>
<b>Product class:</b>	<b>Class I according to rule 1 in Annex VIII in MDR 2017/745</b>
<b>Conformity procedure:</b>	<b>Self-certification according to Annex IV in MDR 2017/745</b>
<b>Identification:</b>	<b>All products with serial numbers issued from;</b> <b>LOT number: 517470</b> <b>Date: 2021-05-25(yyyy-mm-dd).</b>

### 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: 2021-05-23

Sign: \_\_\_\_\_

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