EU Declaration of Conformity

according to the REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL

Doc. No.: RX-T02-008-01

On the basis of the referenced test report(s), sample(s) tested of the below product have been found to comply with the standards harmonized with the directives listed on this verification at the time the tests were carried out. Other standards and Directives may be relevant to the product. This verification is part of the full test report(s) and should be read in conjunction with it <them>.

Manufacturer:		[redacted]
Address:		[redacted]
Single Registration Number (SRN) of the Manufacturer:		--
Authorised representative (AR):		Armyshop Herzogenbuchsee GmbH
Address:		Lagerstrasse 19, 3360 Herzogenbuchsee
Single Registration Number (SRN) of AR:		DE-AR-000005132

We, the manufacturer, declare under our sole responsibility that:

the medical device(S)	**Product Name:**	Military Tourniquet
	Type/model, identification of product allowing traceability (Where applicable):	95*3.8cm
	EMDN Code:	M03040201
	Intended Purpose:	The Military Tourniquet is intended use to stop massive haemorrhage of limbs on field conditions or in normal pre-hospital emergency.
	Classification: (Annex VIII of the MDR)	Class I Medical Device according to Rule 1, Annex VIII of Regulation (EU) 2017/74
	Basic UDI-DI:	--
Conformity assessment procedure:		Annex IX of Regulation (EU) 2017/745
Applied harmonized standards and Common Specification:		*Refer to the Appendix I for details.*