

## SAM® Splint Declaration of Conformity

EUDOC-0001-D

Valid through: 2026-05-18

Page 1 of 3

## EU DECLARATION OF CONFORMITY AS PER ANNEX IV OF THE REGULATION (EU) MDR 2017/745

Manufacturer:



SAM® Medical Products

12200 SW Tualatin Road, Suite 200, Tualatin, OR 97062, USA

Tel: + 1 (503) 639-5474 | Fax: +1 (503) 639-5425

quality@sammedical.com

Single Registration Number (SRN):

US-MF-000002589

EU Authorized Representative:



**Emergo Europe** 

Westervoortsedijk 60, 6827 AT Arnhem, The Netherlands

Tel: +31 (0)70 345 8570 emergoeurope@ul.com

Single Registration Number (SRN):

NL-AR-000000116

Product Family Name

SAM® Splint

Basic UDI-DI:

0822045SP01US (see details in Table 1 attached)

Device(s) concerned:

This Declaration applies to all devices and variants included within the SAM® Splint Product Family (see details in Table 1 attached).

Intended Purpose:

The SAM Splint is intended to temporarily immobilize extremities following traumatic injury.

Risk Class per Annex VIII:

Class I (non-sterile) as per Rule 1

**GMDN** Code

63273 (First aid limb splint, mouldable, reusable)

**EMDN** Code

M03050202 (Synthetic Cast Bandages and Splints)

**Notified Body:** 

Not applicable.

Class I (non-sterile, non-measuring) devices are not reviewed by a Notified body.

Conformity Assessment Route:

SAM Medical<sup>®</sup> Products utilizes Annex II and Annex III Technical Documentation (including PMS) for Class I EU medical devices and issues a Declaration of Conformity (self-certification).

Applicable CE Certificate(s):

Not applicable - Class I (non-sterile) devices are self-certified.

Standards and Common Specifications (CS):

This certificate further declares that the products covered herein also comply with the applicable requirements of relevant standards and Common Specifications specified in Table 2.

This declaration of conformity is issued under the sole responsibility of SAM® Medical Products. We hereby declare that the medical devices specified above meet the applicable provisions of the Medical Devices Regulation (EU) MDR 2017/745. All supporting documentation is retained at the premises of the manufacturer.

Person authorized to sign on behalf

Signature & date:

2023-06-26

sign on beh of SAM® Medical Products:

Name: Jeff Lipps

Position: Director RA/QA, SAM® Medical Products

Place of Issue: 12200 SW Tualatin Road, Suite 200, Tualatin, OR 97062, USA



## SAM® Splint Declaration of Conformity

Page 2 of 3

EUDOC-0001-D Valid through: 2026-05-18

Table 1: Medical devices and variants included in the SAM® Splint Product Family

		ical devices and variants included in the SAM® Splii	Pookering Level	SKU
Basic UDI-DI	GTIN	Product	Packaging Level	SKU
	00822045428737	SAM WRIST SPLINT 9IN FLAT CHARCOAL	Each	SP500-CH-EN
	10822045428734		Case	
	00822045428720	SAM WRIST SPLINT 9IN FLAT	Each	SP500-OB-EN
	10822045428727	ORANGE/BLUE	Case	
	00822045428928	SAM JUNIOR SPLINT 18IN FLAT CHARCOAL	Each	SP502-CH-EN
	10822045428925	SAM SOMICIC OF EIGHT TOHAT EACH OF MICOCALE	Case	
	00822045000988	SAM JUNIOR SPLINT 18IN FLAT	Each	SP502-OB-DE
	10822045000985	ORANGE/BLUE	Case	0,002 05 52
	00822045428713	SAM JUNIOR SPLINT 18IN FLAT	Each	SP502-OB-EN
	10822045428710	ORANGE/BLUE	Case	
	00822045000995	WALLEST OF THE CONTROL OF THE CONTRO	Each	CDEO4 OD EN
	10822045000992	INDUSTRIAL SPLINT 24IN FLAT OR	Case	SP504-OB-EN
	00822045001022		Each	00500 011 05
	10822045001029	SAM SPLINT 36IN ROLL CHARCOAL	Case	SP506-CH-DE
	00822045001023		Each	
	10822045428680	SAM SPLINT 36IN ROLL CHARCOAL	Case	SP506-CH-EN
			Each	
	00822045001053	SAM SPLINT 36IN ROLL ORANGE/BLUE	Case	SP506-OB-DE
	10822045001050		Each	
	00822045428669	SAM SPLINT 36IN ROLL ORANGE/BLUE	Case	SP506-OB-EN
	10822045428666			
	00822045001305	SAM SPLINT 36IN ROLL OD	Each	SP506-OD-DE
	10822045001302		Case	
	00822045001299	SAM SPLINT 36IN ROLL OD	Each	SP506-OD-EN
	10822045001296	GAIN OF EIN COUNTROLL OF	Case	
	00822045001077	SAM SPLINT 36IN FLAT CHARCOAL	Each	SP507-CH-DE
	10822045001074	SAW SPEINT SOINT EAT CHARGONE	Case	0,00,01,02
	00822045428690	SAM SPLINT 36IN FLAT CHARCOAL	Each	SP507-CH-EN
	10822045428697	SAM SPLINT SOIN FLAT CHARCOAL	Case	
	00822045001091	CANAGE INT COIN ELAT OF ANOE /PLUE	Each	SP507-OB-DE
0822045SP01US	10822045001098	SAM SPLINT 36IN FLAT ORANGE/BLUE	Case	
	00822045428676		Each	SP507-OB-EN
	10822045428673	SAM SPLINT 36IN FLAT ORANGE/BLUE	Case	
	00822045001312		Each	SP507-OD-DE
	10822045001319	SAM SPLINT 36IN FLAT OD	Case	
	00822045001313		Each	SP507-OD-EN
	10822045428918	SAM SPLINT 36IN FLAT INT'L OD	Case	
	00822045428904		Each	
		SAM SPLINT XL 36IN FLAT CHARCOAL	Case	SP508-CH-EN
	10822045428901		Each	
	00822045001114	SAM SPLINT XL 36IN FLAT ORANGE/BLUE	Case	SP508-OB-DE
	10822045001111		Each	
	00822045428652	SAM SPLINT XL 36IN FLAT ORANGE/BLUE		SP508-OB-EN
	10822045428659		Case	
	00822045428744	SAM FINGER SPLINT ORANGE/BLUE	Each	SP510-OB-EN
	30822045428745		Case	
	00822045428973	ORANGE/BLUE SPLINT KIT 1 36IN, 1 18IN, 2	Each	SP512-OB-DE
	10822045428970	FINGERS	Case	
	00822045000858	OMS-ZOLL SPLINT 18IN FLAT	Each	SP522-OB-CZ-759
	10822045000855	ORANGE/BLUE	Case	5. 522 55 52 755
	00822045000834	HELBIG SPLINT 18IN FLAT ORANGE/BLUE	Each	SP522-OB-DE-572
	10822045000831	HELBIG SPLINT TOIN FEAT ONANGE/BEGE	Case	01 022-00-DL-012
	00822045000636	ONAC ZOLL CRUNT SCIN DOLL CHARCOAL	Each	SP526-CH-CZ-784
	10822045000633	OMS-ZOLL SPLINT 36IN ROLL CHARCOAL	Case	01 020-011-02-704
	00822045000674	RORACO SPLINT 36IN ROLL ORANGE/BLUE	Each	SP526-OB-DE-512
	10822045000671		Case	
	00822045000704	TINOVAMED SPLINT 36IN ROLL	Each	CDE26 OD DE 742
	10822045000701	ORANGE/BLUE	Case	SP526-OB-DE-713
	00822045000701	SAMARITER SPLINT 36IN ROLL	Each	SP526-OB-DE-714
	10822045000711		Case	
	00822045000718		Each	00500 05 55 55
		1	Case	SP526-OB-DE-794
	10822045000770	ORANGE/BLUE	Lase	I



## SAM® Splint Declaration of Conformity

EUDOC-0001-D Valid through: 2026-05-18

Page 3 of 3

Basic UDI-DI	GTIN	Product	Packaging Level	SKU
	00822045000780	ORK SPLINT 36IN ROLLED ORANGE/BLUE	Each	SP526-OB-DE-795
	10822045000787	ORK SPLINT 30IN ROLLED ORANGE/BLUE	Case	
	00822045000766	ALLENSPACH SPLINT 36IN ROLL	Each	SP526-OB-EN-780
	10822045000763	ORANGE/BLUE	Case	
	00822045001275	DAV SPLINT 36IN ROLL ORANGE/BLUE	Each	SP526-OB-DE-811
	10822045001272		Case	
	00822045000797	SCHWEIZER SPLINT 36IN ROLL	Each	SP526-OB-EN-779
	10822045000794	ORANGE/BLUE	Case	
	00822045000766	ALLENSPACH SPLINT 36IN ROLL	Each	SP526-OB-EN-780
	10822045000763	ORANGE/BLUE	Case	
	00822045000803	SP SERVICES SPLINT 36IN ROLL	Each	SP526-OB-EN-797
	10822045000800	ORANGE/BLUE	Case	
	00822045000445	ONAG ZOLL ODLINIT SCINI EL AT CHADCOAL	Each	SP527-CH-CZ-802
	10822045000442	OMS ZOLL SPLINT 36IN FLAT CHARCOAL	Case	
	00822045000469	OMS-ZOLL SPLINT 36IN FLAT	Each	SP527-OB-CZ-721
0822045SP01US	10822045000466	ORANGE/BLUE	Case	
	00822045000490	ALLENSPACH SPLINT 36IN FLAT	Each	SP527-OB-DE-781 - SP527-OB-DE-796
	10822045000497	ORANGE/BLUE	Case	
	00822045000513	HELBIG SPLINT 36IN FLAT ORANGE/BLUE	Each	
	10822045000510		Case	
	00822045001282	TINOVAMED SPLINT 36IN FLAT	Each	SP527-OB-DE-812
	10822045001289	ORANGE/BLUE	Case	31 321-0D-DE-012
	00822045000506	SP SERVICES SPLINT 36IN FLAT	Each	SP527-OB-EN-792
	10822045000503	ORANGE/BLUE	Case	
	00822045001251	ALLENSPACH SPLINT 36IN FLAT	Each	SP527-OB-EN-808
	10822045001258	ORANGE/BLUE	Case	
	00822045001268	SCHWEIZER SPLINT 36IN FLAT	Each	SP527-OB-EN-809 SP528-CH-CZ-789
	10822045001265	ORANGE/BLUE	Case	
	00822045000568	OMS ZOLL XL 36IN FLAT CHARCOAL	Each	
	10822045000565		Case	
	00822045000582	OMS-ZOLL SPLINT XL 36IN FLAT	Each	SP528-OB-CZ-760
	10822045000589	ORANGE/BLUE	Case	

Table 2: Standards and Common Specifications (CS) applied

Standard #	Title	Year / Version
otaridard #	Applied Standards	
EN ISO 10993-1	Biological evaluation of medical devices — Part 1: Evaluation and testing within a risk	2020
	management process	
EN ISO 10993-18	Biological evaluation of medical devices — Part 18: Chemical characterization of medical	2020
	device materials within a risk management process	
EN ISO 13485	Medical devices - Quality management systems - Requirements for regulatory purposes	2016+A11:2021
EN ISO 14971	Medical Devices - Application of Risk Management to Medical Devices	2019+A11:2021
EN ISO 15223-1	Medical devices - Symbols to be used with medical device labels, labelling and information to	2021
	be supplied - Part 1: General requirements	
EN ISO 20417	Medical Devices - Information supplied by the manufacturer	2021
EN 62366-1	Medical devices – Part 1: Application of usability engineering to medical devices	2015+A1:2020
	Other relevant standards	
EN ISO 17100	Translation services — Requirements for translation services	2015+A1:2017
ASTM F2052	Standard test method for measurement of magnetically induced displacement force on	2021
	passive implants in the magnetic resonance environment	
ASTM F2503	Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic	2020
	Resonance Environment	
AAMI TIR 12	Designing, testing, and labeling reusable medical devices for reprocessing in health care	2010
	facilities: A guide for medical device manufacturers	
AAMI TIR 30	A compendium of processes, materials, test methods, and acceptance criteria for cleaning	2011(R)2016
	reusable medical devices	
	Common Specifications	
	No common specifications relevant to the device family have been published in OJ at this time.	