DATA SHEET WESTMED® Universal FlexSplint

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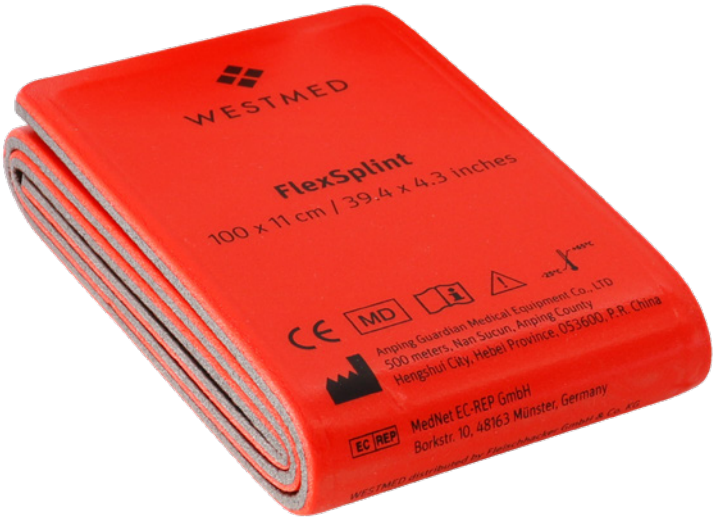

Order number
5810321

Description

Our WESTMED® Universal FlexSplints are suitable for splinting and immobilizing extremities. The easy-to-apply splint is ideal in an emergency.

The FlexSplints are made of a plastic foam (IPXE) and aluminum mixture and can be easily cut to the desired length with scissors. They can be shaped to suit almost any application. The liquid-repellent material makes them easy to disinfect and reuse after use.

Technical information	
Reference number	AZ-ST100
Intended use	The FlexSplint Universal Splints are used exclusively for immobilization and splinting of fractures and sports injuries such as ligament sprains.
Dimensions	100 cm x 11 cm
Wheight	160 g
Material	Cover: Polyethylene foam (IXPE) Core: Aluminum
Storage and transport conditions	–25 °C to 65 °C
Minimum durability	5 years from date of production
Packaging unit	1 piece per pack 50 packs per box 40 boxes per pallet

Cleaning and disinfection	The FlexSplint Universal Splint can be reused. The splint must be cleaned and disinfected after each use in accordance with the relevant local hygiene guidelines. FlexSplint can be cleaned with a mild cleaner or a surface disinfectant. Please follow the manufacturer‘s instructions for the cleaner or disinfectant. After cleaning or disinfection, the splint must be rinsed with clear water to remove the remaining cleaner or disinfectant. Allow the splint to dry afterwards.

General information	
Safety instructions	• These instructions for use do not replace detailed instruction or training of the user. • The FlexSplint universal splints may only be used for the purpose described in these instructions for use. • For prolonged use (> 1 hour), the splint should be padded on the patient side. • When cutting the universal splint to size, sharp edges may occur which must be covered or folded. • Open wounds must be covered with sterile dressings before using the splint. • The splint is not suitable for use in fractures of the spine and pelvis. • When securing the splint, it is essential to ensure that the blood supply is not interrupted. The tips of the fingers or toes must be exposed to check the blood supply.
Regulatory information	
Classification	Class I medical device According to Annex VIII, Regulation (EU) 2017/745

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CE	The CE marking is short for „Conformité Européenne“, which means „European Conformity“. It symbolizes the conformity of the product with the applicable requirements that the European Community places on the manufacturer.
	Manufacturer
	Date of production
	Observe instructions for use
EC REP	Authorized representative Europe
REF	Reference number/article number
	Temperature limit
MD	Medical device
	Caution/Attention
	Distributor
	Importer

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EC REP

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