



The following should be read thoroughly before use:

**Important Notes:**

This document is designed to provide instructions on how this product/s is used. It is not a reference to clinical techniques, and does not replace local or national guidance or protocols.

**Intended User:**

For use by both suitably trained healthcare professionals in a healthcare establishment and 'Lay Users' in a home use environment.

**Intended Purpose:**

Provides foot protection without total immobilization. Adjustable to accommodate bandages

**Indications For Use:**

- Any forefoot conditions e.g. Hammer Toes, Bunions, Neuroma etc
- Mid-foot procedures
- Osteotomies

**Product Characteristics / Features:**

Temporary Footwear

Key functional elements include:

- Padded collar to reduce slippage
- Hook and loop straps with D-ring for secure fastening
- Lightweight sole with padded breathable upper
- Removable tongue for optimum placement and comfort
- Sold singularly
- Latex free

**How it's supplied:**

It is supplied non-sterile and ready for single patient use.

**Devices Required For Use, And Interoperability**

**Requirements:**

None

**Storage Conditions:**

Store in box, away from direct sunlight and heat sources in a cool and dry environment.

**Care and Maintenance:**

- Wash periodically by hand with a damp cloth and a mild soap
- Use a towel to absorb most of the dampness and then allow to dry at room temperature
- Do not dry clean or tumble dry - allow to dry naturally away from direct heat sources
- Do not expose to alcohol, ointments or solvents
- The device should be safety checked by the providing service and/or user every 6 months.

**Contraindications:**

None

**Precautions / Warnings / Residual Risks:**

If allergic skin rashes occur, please consult your healthcare professional

**Possible Adverse Reactions:**

None

**Single Patient Use Precaution:**

This device is designed and sold for single patient use only. The effects of any unauthorised reuse can result in the following complications:  
Risk of Cross contamination;  
Risk of mechanical fatigue, and/or associated failure.

**Directions For Use / Fitting Instructions:**

- Open all the straps as fully as possible
- Carefully insert the foot into the footwear making sure the heel is at the back of the shoe
- Close the straps securely but comfortably

**Disposal**

Discard after single patient use and dispose of as common waste according to national guidelines.
















**Serious Incident Reporting**

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

### Sizes & REF / Codes

Size	Sole Length (CM)	Shoe Size		Code
		UK	EU	
XX Small	24	1 - 3	33 - 36	POS-F-XXS
X Small	27	3 - 5	36 - 38	POS-F-XS
Small	28.5	5 - 6½	38 - 40	POS-F-S
Medium	29	6½ - 8	40 - 42	POS-F-M
Large	30	8 - 9½	42 - 44	POS-F-L
X Large	32	9½ - 11	44 - 46	POS-F-XL
XX Large	33.5	11 - 13½	46 - 48	POS-F-XXL

**LABELLING SYMBOLOGY**  
EN ISO 15223-1:2021, unless otherwise stated

	Manufacturer		Consult Instructions for Use		Unique Device Identifier
	Authorised representative in the European Community		Caution		Single Patient Use
	Medical Device		Non-Sterile		Quantity - Abbreviation
	Catalogue Number		CE Mark - EU Regulation 2017/745		Contains Latex
	Batch Code		UKCA Mark - UK Regulation SI 2002 No 618		Date of Manufacture

POS-F IFU - Revision 6 - September 2024

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