



**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:**

Designed to stretch over the opposite shoe or boot to equalise leg discrepancy caused by wearing an orthosis such as a walker boot or post-op shoe

**Indications For Use:**

- For use with walker boots or post-op footwear to balance leg length

**Product Characteristics / Features:**

- Designed to stretch over the opposite shoe or boot to equalise leg discrepancy caused by wearing an orthosis such as a walker boot or post-op shoe
- Lightweight design and height adjustable with removable heel to accommodate various types of footwear
- Unique insole with 3 layer split design allowing for greater range of adjustment
- Simply add or remove an insole layer to provide 15mm, 24mm or 30mm of height
- Improves gait and relieves back and hip pain.
- Suitable for both indoor and outdoor use
- Fits both left and right

**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:**

This product is to be used under the supervision of a medical professional. If you experience any pain, swelling, sensation changes, or any unusual effects while using this product, consult your medical professional immediately. Not intended for use during high-impact activities. Check fit regularly; improper adjustment can cause discomfort or imbalance.

**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:**

- Refer to the size chart to select the appropriate size based on your usual shoe size
- Place the Equaliser under the shoe on the unaffected side e.g. on the opposite foot from your walker boot or post-op shoe
- Add or remove insole layers to achieve the desired "matching" lift
- Adjust the flexible strap around your shoe to provide a secure and comfortable fit
- Test your gait by walking on a flat surface and adjust height as needed.

## 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	Shoe Size		Code
	UK	EU	
Small	3 - 5	35 - 38	EQL-S
Medium	5 - 8½	38 - 42	EQL-M
Large	9 - 11	43 - 46	EQL-L
X Large	11½+	47+	EQL-XL

**EXPLANATION OF 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

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