EN

Multi Arm Sling

Instructions for Use



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TalarMade Ltd is a UK-based manufacturer of Class I medical devices, committed to providing high-quality, safe, and effective products designed to support healthcare professionals and patients.

This Instruction for Use (IFU) document provides essential information for the correct and safe use of TalarMade products, in compliance with the requirements of the UK Medical Devices Regulations 2002 (as amended) and the EU Medical Device Regulation (EU) 2017/745 (MDR), as applicable.

All TalarMade medical devices are designed, manufactured, and distributed under a certified Quality Management System in accordance with ISO 13485:2016, ensuring that they consistently meet both regulatory and customer requirements for safety, performance, and reliability.

Prescribers and users are advised to read this document carefully before use. Proper understanding and adherence to the instructions, warnings, and precautions outlined

herein are essential to ensure optimal product performance and to minimise the risk of misuse or adverse events.

Product Details

Product Name: Multi Arm Sling

GMDN: 12101 – Shoulder immobilizer/orthosis, reusable

BASIC UDI-DI (if applicable): 50552043ULOSEWHO4Z

Packaging contents: 1 x Multi Arm Sling

Manufacturer Details

Manufacturers Registered Address

TalarMade Ltd. Edinburgh House, Millennium Way, Chesterfield, Derbyshire, S41 8ND

Tel: +44 (0)1246 268 456 | talarmade.com | Email: info@talarmade.com

Device Description

- Simple arm sling made from brushed velour with fully bound edges
- Fully adjustable shoulder strap and broad full circumference torso strap
- Easy touch-and-close fastenings
- Universal sizing fitting both left and right
- Sold singly

Intended Use

The Multi Arm Sling is designed to provide immobilisation of the arm/elbow/shoulder.

Target users are healthcare professionals and laypeople, both in hospital/healthcare environments and at home.

The Multi Arm Sling should be worn/used under the guidance of a healthcare professional and as outlined in the instructions for use.

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Indications

The Multi Arm Sling is indicated for use in the following situations:

- Immobilisation of the forearm, elbow and arm
- Post upper limb trauma/surgery
- Weakness /injury to the shoulder capsule
- Paralysis of the arm

Contraindications

The Multi Arm Sling should not be used:

- For treatment of individuals with severe fixed deformity
- Over open wounds that may come into contact with the device

Warnings

Caution	Hazard	Potential Harm	Risk-Control / IFU Warning
⚠ **Warning**	Application Over Open Wounds	Delayed Healing or Infection	Do not use over open wounds without appropriate dressings and medical supervision.
⚠ **Warning**	Incorrect Application	Ineffective function	Application should be performed by healthcare professionals or trained caregivers. If the device is incorrectly fitted or positioned, function may be reduced, and the risk of pressure related issues may increase.
⚠ **Warning**	Skin Monitoring	Patient Injury	The patient's skin must be inspected regularly for signs of redness or irritation. If irritation or skin damage occurs, discontinue use and consult a healthcare professional.

⚠ **Infection Control**	Cross-contamination, loss of performance	Infection transmission	The device is intended for single patient use only. The device may be cleaned during use according to the cleaning instructions provided.
⚠ **Do Not Modify**	Device Condition	Ineffective function, increased injury risk	Do not cut, pierce, or modify the device, as this may affect safety, performance, and function.
⚠ **Fire Hazard**	Fire	Fire, Patient injury	Keep the device away from open flames, lit cigarettes, or other sources of ignition.
⚠ **Keep dry / moisture hazard**	Use of wet or contaminated device	Skin maceration, microbial growth, reduced device performance	Ensure device is dry before use and is cleaned regularly

Precautions

- This device is for **single patient use only**. Re-use by another individual may increase the risk of cross-contamination, improper fit, or compromised device performance
- The device should be worn as instructed by the prescriber
- Inspect the device integrity before use
- Remove brace and cease use if the brace causes pain, redness or skin breakdown

Instructions for Use

Preparation

- Remove packaging before inspecting brace, making sure all parts are intact and without damage

Application

1. Gently place the injured arm into the sling ensuring that the support is snug around the elbow. Adjust for length by turning the sling inwards from the

patient's fingers to the desired position.

2. Close the sling over the hand by attaching/fastening the D-ring strap
3. Close the sling around the elbow by attaching/fastening the sling closure strap
4. The arm can now be supported in the desired position by bringing the shoulder strap (fixed at the elbow) over the contralateral shoulder and fixing through the top D-ring near hand using touch-and-close fastening.
   The affected hand and forearm should be elevated above the level of the elbow as this helps to prevent blood and fluid from pooling in the wrist/hand.
5. To immobilise the shoulder, attach one end of the torso strap through the D-ring at the front of the forearm before taking around the body and attaching in the final remaining D-ring near the hand.

Removal

1. Release touch-and-close straps
2. Remove Multi Arm Sling
3. Inspect skin, checking for any signs of irritation, redness, or pressure marks.

During Use

- Wearing schedule:
  - Follow the duration and daily wear time recommended by your healthcare professional or medical care plan
- Monitoring:
  - Periodically check skin condition and comfort.
  - Adjust positioning of brace as needed

Cleaning & Maintenance

- Close touch-and-close straps
- Multi Arm Sling can be hand washed in cool water with mild detergent
- Do not use bleach or chlorine-based cleaning products
- Not for use in washing machine or tumble dryer
- Brace should be left to dry naturally before re-applying

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Storage Conditions

- Store in a cool clean dry place and away from direct sunlight or heat to prevent discolouration or deformation
- Do not place any objects on top of the device

Disposal Instructions

Dispose of the device and its packaging in accordance with applicable local, regional, or national regulations.

Regulatory Compliance

EU REP

EUROPEAN AUTHORIZED REPRESENTATIVE

MedNet EC-REP GmbH, Borkstrasse 10, 48163 Muenster, Germany

SRN number: DE-AR-000000002

CH REP

SWISS AUTHORIZED REPRESENTATIVE

MedNet SWISS GmbH, D4 Platz 4, 6039 Root D4, Switzerland

CHRN: CHRN-AR-20000730

Warranty Information

The device is supplied under the manufacturer's warranty as outlined in the current Terms and Conditions available on our website. Users are advised to review these terms prior to use.

Contact Information

We are committed to continuously improving our products.
Please scan the QR code to share your feedback with us.

No camera? Click here or on the QR code to open the product feedback form.

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