

Upper Limb

Sub Lux Support			
SLS/03	SLS/05	SLS/07	SLS/08

Description

An orthosis that provides effective unilateral anti-subluxation support. The support positions the scapula in external rotation and enhances the natural gait pattern. The dynamic straps allow patient to utilise returning muscles and perform rehabilitation exercises.

Intended Purpose

To provide upper limb support for an array of injuries and conditions, including Pre or Postoperative treatment for: Post stroke, Soft tissue injury, Ligament strain, Glenohumeral subluxation.

Sizing

Measure the circumference at the largest part of the upper arm.

⚠ Fitting Instructions

We recommend the initial fitting of the device be conducted by a suitably qualified Healthcare Professional who will advise fitting, removal and period of use.

1. With the buckles on the upper side of the cuff, wrap the cuff around the upper arm making sure that you have two fingers width between the top edge of the cuff and the axilla. **See Fig.1.**
2. Secure the cuff using the hook and loop fastenings.
3. Once the cuff is securely fastened, attached the strap to the back buckle **See Fig.2.**
4. Take the strap across the back and over the opposite shoulder. **See Fig.3.**
5. Once over the shoulder, bring the strap down and under the axilla (of the non-affected shoulder) and across the back and over the affected shoulder. The straps are now "crossed" at the back. **See Fig.4.**
6. Locate the end of the strap in the anterior buckle on the cuff and secure. **See Fig.5.**
7. Draw through the tension on the posterior buckle until the affected shoulder lifts approximately in line with the non-affected shoulder and/or relieves the patient discomfort. Secure the bar buckle.



Contraindications

- Must not be used by individuals for whom compression is contraindicated
- Do not use on open wounds
- Any known allergy and/or hypersensitivity to any material listed in the product composition
- Caution should be used if prescribed to patients with diabetes, vascular deficiency and neuropathy

Warnings and Precautions

- Carefully read all instructions and warnings before use
- Follow all instructions to ensure proper performance of the brace
- Do not use if liniments, ointments, gels, creams or any other substances have been applied to the affected area
- Do not reuse for another patient, doing so risks cross-infection and can compromise product integrity
- To ensure the straps are not over tightened, perform regular skin and circulation checks, especially for patients with diabetes, vascular deficiencies and neurological conditions
- Should any adverse reactions occur, please remove, do not use and contact your healthcare professional or provider
- Check brace and all components for wear and tear
- The hook and loop fasteners must always be fastened when the device is not being worn or when being washed
- The durability of the device may be compromised by certain factors, e.g. objects with sharp edges or damage to the hook and loop fasteners

Washing and care instructions

- Cover all the hook and loop straps
- Hand wash the brace in cool water using a mild detergent
- Rinse and air dry flat
- Do not spin or tumble dry
- Do not use fabric conditioner at any time
- Check for any degradation that may affect use
- Periodically check the hook and loop straps are operating securely

Product composition

Velfoam, Hook and Loop, Webbing, Nylon, Nickel

Storage and transport conditions

Store in a cool, dry place out of direct sunlight, in the original packaging.

Recycling and disposal

Packaging and constituent parts should be recycled or safely disposed of in accordance with local or national laws.

Serious incident

Report any serious incident to the manufacturer and the competent authority of the EU Member State, or Country in which you reside.

Wash symbols



Hand wash



Dry flat



Do not tumble dry



Do not bleach



Do not dry clean



Do not iron



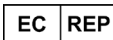
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This Medical Device conforms to the requirements of Regulation (EU) 2017/745 of the European Parliament and of the council of 5 April 2017



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