

Upper Limb

Paediatric Cotton Sling

201/PES	202/PES	201/PES.RD	202/PES.RD
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Description

The paediatric Cotton Sling is constructed from a comfortable and lightweight cotton webbing material, offering support for the elbow, wrist and hand. The lightweight design has a large interior pocket, which provides comfortable support with plenty of room for a cast. The sling features a secure and adjustable hook and loop closure system, making it easy to find the most comfortable angle and position for recovery. The soft Velfoam padded shoulder strap reduces pain and discomfort that can often occur around the back of neck. The fully adjustable shoulder strap makes it easy to find the most comfortable angle and position for recovery.

Intended Purpose

Provides support for injuries and conditions including: Paralysis, Postoperative, Oedema control, Immobilisation following injury or surgery.

Sizing

Measure the length from the elbow point to the start of the little finger

▲ 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. Place the injured arm into the sling, ensuring the elbow and forearm are supported securely to avoid discomfort
2. Run the shoulder strap up from the injured elbow, round the neck and over the uninjured shoulder. Feed it through the D-ring on the outside of the sling and secure with the hook and loop fastening **See Fig.1.**
3. Attach the immobilisation strap on to the side of the sling **See Fig.2.**
4. Feed the immobilisation strap through the D-ring (as shown) and secure with the hook and loop fastening **See Fig.3.**
5. The soft thumb loop maintains optimal position and supports the hand **See Fig.4.**

Fig.1.

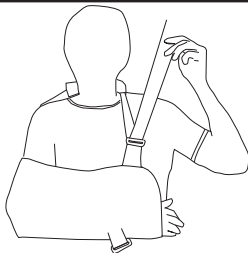


Fig.2.

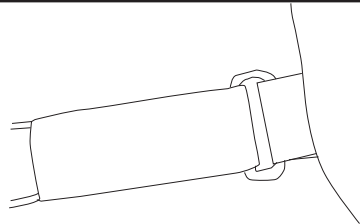


Fig.3.

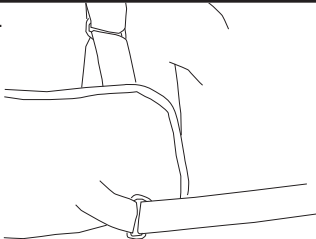
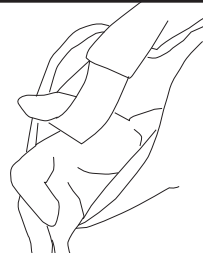


Fig.4.



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 re-use 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

Cotton, Hook and Loop, Plastic Buckle

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



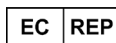
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This Medical Device conforms to the requirements of Regulation (EU) 2017/745 of the European Parliament and of the council of 5 April 2017



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