

Spinal

Dual Pull Lumbo Support

DPLS/03	DPLS/05	DPLS/07	DPLS/08
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Description

An orthosis designed to support the spine by providing stability for lower back pain and sacroiliac joint pain. The brace provides added support by facilitating good posture, preventing strain on the pain-producing structures of the lumbar area. The brace features flexible non-removable metal stays to improve posture. The brace can be adjusted from gentle to maximum compression by pulling the wings with little to maximum force. Once the brace has been adjusted to the desired level of compression using the anterior touch and close fastenings, the belt acts as a support to reinforce the spine and torso. The contoured shape of the brace has an anterior depth of 180mm and a posterior depth of 240mm.

Intended Purpose

To provide back support for an array of injuries and conditions, including Pre or Postoperative treatment for: Sciatica, Lumbosacral strains, Degenerative Disc Disease, General posture support.

Sizing

Measure the circumference of the waist.

△ 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. Align the belt to midline of lower back, making sure the flexible steels are equidistant apart from either side of the spine **See Fig.1.**
2. Wrap the brace around the front and secure the hook and loop fastening **See Fig.2.**
3. For additional support, pull the side straps and secure the hook and loop fastening **See Fig.3.**
4. Adjust as necessary to ensure optimum fit **See 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
- This product contains Natural Rubber Latex which may cause allergic reactions
- 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

Elastic, Aluminium, Hook and Loop, Latex

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



Contains natural rubber latex



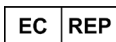
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This Medical Device conforms to the requirements of Regulation (EU) 2017/745 of the European Parliament and of the council of 5 April 2017



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