



Data Sheet

THERMARMOUR XTRAS – HEAT WRAP

Product Code	OB900-91127---LBLX65
GTIN/UDI	5061035932327
Colour(s)	Light blue with darker blue borders
Size	91 x 127cm unfolded
Material	62% polypropylene (PP) / 38% polyethylene (PE), 72 gsm, 3-layer construction
Packing	10 pieces per carton, 31 x 34 x 30cm
Active	Iron Powder, Water, Vermiculite, Activated Carbon, Absorbent Polymer & Salt.
Care	Patient specific, single use only – do not reuse or launder
Key Features	- Individually vacuum packed for ease of storage - Featuring 8 oxygen-activated heat packs - No external power source required
Buy Online	www.interweavetextiles.com/products/patient-warming/

The Heat Wrap is intended to provide heat therapy for the temporary relief of injury or pain. It is applied to the required body area to provide heat therapy to the surface and/or underlying tissue.

Please read the full Instructions for Use, including all warnings, carefully before use and ensure they are readily accessible to users and/or appropriate specialist staff. This product is not a substitute for professional medical assessment or intervention. In cases of persistent discomfort, severe injury, or prolonged skin irritation, users should discontinue use and seek medical attention.

- Open the packaging and fully unfold the wrap.
- The wrap will begin to warm naturally; for best results, open 10-15 minutes before application.
- Apply to clothing or skin on the required body area.
- If access is needed, gently slide or lift aside — do not fold during use.
- Regularly check skin response and overall physical condition throughout.
- Remove and dispose of the wrap once the heat has dissipated.



Please note, Jiangsu Hanyao Medical Devices Co., Ltd. manufactures several variations of Heat Pack products. The Data Sheet & IFU applies to the non-adhesive THERMARMOUR Heat Wrap 91cm x 127cm only, distributed by Interweave Textiles Ltd. Interpretation and usage should be adapted accordingly. If in any doubt, please seek professional medical guidance. Any serious incidents that occur in relation to this device should be reported to the manufacturer and the competent authority of the patient/user.

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