Lastodur® light

Long-stretch bandage

HARTMANN

EN Instructions for use

Product description of the device
The long-stretch compression bandages are narrow woven. They are provided with an end finder. The bandages themselves are made of different natural (cotton, viscose) and synthetic (polyamide, elastane) materials in various proportions. Lastodur light is non-active, non-sterile and is a white colored, permanently elastic long-stretch bandage with a yellow line in the center to guide through the application process.

Intended purpose
The devices are single-use, non-sterile long-stretch compression bandages which can be used for long-term compression therapy for varicose symptoms; as prophylaxis for thrombosis on immobile patients; as a support and relief bandage in musculo-skeletal injuries. They can be used by healthcare professionals in clinical and home environments in combination with padding bandages and inner lining material.

Properties
- Extensibility: approx. 170%.

Indications
The devices can be used for the treatment of varicose veins, thrombosis / thrombophlebitis, chronic venous insufficiency (CVI), primary and secondary prevention of leg ulcer venosum / venous insufficiency, oedemata of other origin (ankle brachial pressure index - ABPI 0.8 - 1.3). The devices can also be used as support and relief bandages in musculoskeletal injuries. They are suitable for use as support and relief bandages in the management of limbs following amputation. There is an additional usage as support bandages for distorsions, contusions, tendon injuries, thoracic injuries and dislocations (bone).

Composition
83% viscose, 12% polyamide, elastane balanced to viscose and polyamide.

Contraindications
The devices must not be used in cases of advanced peripheral arterial occlusive disease (PAOD), decompensated cardiac insufficiency, phlegmasia cerulea dolens, septic phlebitis and in case of ankle-brachial pressure index (ABPI) > 1.3 and ≤ 0.5.

Method of application
Apply the bandage from the base of the toes in a figure of 8 technique to just below the knee. Use the yellow line as a guide to determine one half the width of the bandage. While wrapping, stretch the bandage to 50% of its maximum extensibility. Secure the end with tape.

Further Remarks
- If the bandage is applied in a figure of eight with 50% stretch and 50% overlap the pressure achieved will be higher than if it is applied in a spiral with 50% stretch and 50% overlap.
- Do not use the bandage as a primary dressing or directly on open wounds. The bandage should be used in conjunction with an undercast padding bandage to protect the leg and pad out bony prominences.
- The bandage is recommended to be removed overnight.
- Do not let the bandage get wet during showering / bathing.
- A slipping of the bandage indicates a loss of compression. In this case, a medically trained person has to reapply the bandage.
- Reusing a single-use medical device is dangerous. Reprocessing devices, in order to reuse them, may seriously damage their integrity and their performance. Information available on request.

Precautions
- Special precautionary measures are required in the event of pronounced sensory disorders of the extremities (e.g. in the case of advanced diabetic peripheral polyneuropathy) compensated peripheral arterial occlusive disease (PAOD) with a reduced ankle-brachial pressure index (ABPI 0.6-0.8), chronic compensated cardiac insufficiency and florid infectious disease.
- If the foot pulse is not palpable, a Doppler ultrasound should be carried out to determine the ABPI prior to starting treatment with the medical device.
- Incorrect bandaging technique can lead to constrictions, undesired pressure points or other damage.
- If the patient develops severe pain, numbness, tingling or noticeable discoloration of the toes at any time during treatment with the medical device, a medical professional must be contacted immediately and the bandage must be removed.

Incident reporting
For a patient/user/third party in the European Union and in countries with identical regulatory regime (Regulation 2017/745/EU on Medical Devices); if, during the use of this device or as a result of its use, a serious incident has occurred, please report it to the manufacturer and/or its authorised representative and to your national authority.

Product disposal
In order to minimize the risk of potential infection hazards, or environmental pollution, disposable components of medical device should follow disposal procedures according to applicable and local laws, rules, regulations and infection prevention standards. Dispose of the medical device with regular hospital waste.

Date of revision of the text: 2020-08-26

AU – PAUL HARTMANN Pty. Ltd. · Macquarie Park, NSW 2113

Special instructions

MD	Medical Device
	Manufacturer
	Date of manufacture
	Use-by date
LOT	Batch code
REF	Catalogue number
	Consult instructions for use
	Do not re-use
	Keep dry
	Do not cut
UDI	Unique Device Identifier

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