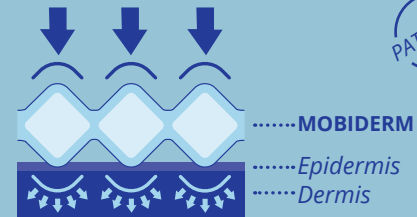


MOBIDERM Technology

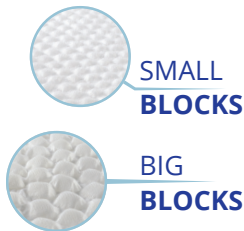
MOBIDERM is a medical device comprised of foam cubes encased between two non-woven bandages.

MOBIDERM Technology can be used under a reducing bandage (in its bandage or pad forms), or incorporated into mobilizing garments.

The pressure difference created between the support area of the blocks and the surrounding area stimulates the flow of lymphatic fluid, optimizing drainage efficacy.⁽³⁾



Two sizes of foam blocks adapt to the various needs of the affected area



- More effective in highly induced areas
- Better for small morphologies or body areas: pediatric patients, hands/foot, breast, and more.

- Better adapted along the limb

MOBIDERM autofit garments are recommend for patients with all stages of lymphedema, dysmorphic or regular morphology, progressive edema.

ORDERING INFORMATION

MOBIDERM autofit Thigh-High	Size	Ankle Circumference	Calf Circumference	Thigh Circumference	Short	Regular	Long	Xtra Long
		(at the narrowest point) (cB)	(at the widest point) (cC)	(at the widest point) (cC)	tG = 67-72 cm	tG = 72-77 cm	tG = 77-82 cm	tG = 82-87 cm
	1	16-24 cm	24-35 cm	46-57 cm	373503 20100231	373503 20100232	373503 20100233	373503 20100234
	2	20-28 cm	32-43 cm	53-65 cm	373503 20200231	373503 20200232	373503 20200233	373503 20200234
	3	24-32 cm	40-51 cm	61-72 cm	373503 20300231	373503 20300232	373503 20300233	373503 20300234
	4	28-36 cm	48-59 cm	68-80 cm	373503 20400231	373503 20400232	373503 20400233	373503 20400234
MOBIDERM autofit Below-Knee	Size	Ankle Circumference	Calf Circumference	Below-Knee Circumference	Short	Regular	Long	Xtra Long
		(at the narrowest point) (cB)	(at the widest point) (cC)	(on the head of the fibula) (cD)	tD = 38-40 cm	tD = 40-42 cm	tD = 42-44 cm	tD = 44-46 cm
	1	16-24 cm	24-35 cm	25-36 cm	373403 20100231	373403 20100232	373403 20100233	373403 20100234
	2	20-28 cm	32-43 cm	32-43 cm	373403 20200231	373403 20200232	373403 20200233	373403 20200234
	3	24-32 cm	40-51 cm	39-50 cm	373403 20300231	373403 20300232	373403 20300233	373403 20300234
	4	28-36 cm	48-59 cm	46-57 cm	373403 20400231	373403 20400232	373403 20400233	373403 20400234

(1) Mestre - Interest of an auto-adjustable night-time compression sleeve (MOBIDERM Autofit) in maintenance phase of upper limb lymphoedema: the MARILYN pilot RCT - Support Care Cancer (25:2455-2462) - 2017 - Page 7

(2) Quéré - Prospective multicenter observational study of lymphoedema therapy: POLIT study - Vascular Disease Journal - 2014 - Pages 4 and 5

(3) Todd - Mobiderm Autofit: an adjustable sleeve that enables patients to self-manage lymphoedema - Chronic oedema - April 2018 - Page 6

(4) Only in following countries: Algeria, Austria, Belgium, Canada, China, Czech Republic, Denmark, Finland, France, Germany, Hungary, India, Morocco, Italy, Netherlands, Poland, Portugal, Romania, Russian Federation, Slovakia, Slovenia, Spain, Sweden, Switzerland, Tunisia, United-Kingdom

Biflex - Indications: Venous insufficiency, simple and complicated varicose veins, reduction of venous and posttraumatic edema, dermatitis / lipodermatosclerosis, varicose ulcers, deep vein thrombosis (prevention and confirmed accident), post-thrombotic syndrome, following sclerotherapy, treatment of lymphedema. Contraindications: PAD (peripheral arterial disease), skin wounds, abscess, infected skin conditions, extra-anatomic bypass, uncontrolled heart failure.

BiflexIdeal® - Indications: Treatment of lymphedema, simple and complicated varicose veins, following sclerotherapy, following surgery, reduction of post-traumatic edema, varicose ulcers. Contraindications: Do not apply directly to an open wound.

Mobiderm Autofit and Standard - Indications: Maintenance of volume reduction in lymphedema. Contraindications: Skin infection of the limb or acute inflammation, known allergy to components used, septic thrombosis, severe peripheral neuropathy of the limb. Lower limb - PAD (peripheral arterial disease) of lower limbs with ABPI < 0.6, congestive heart failure, advanced diabetic microangiopathy, phlegmasia cerulea dolens, extra-anatomic bypass. Upper limb - pathology of brachial plexus, vasculitis of the extremities.

Availability of these products might vary from a given country or region to another, as a result of specific local regulatory approval or clearance requirements for sale in such country or region. All the medical devices mentioned on this document are CE marked according to the European council directive 93/42/EEC and its relatives or the Regulation 2017/745 on medical devices, unless specifically identified as "NOT CE MARKED". Products mentioned in this document are CE class I devices. Please contact Thuasne should you need any additional information on devices classification. Please read carefully the instructions for use, indications and contraindications of the product. Last revision date: 2020/07. Ref: 002



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MOBIDERM

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Patient's Last Name: _____

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Case No. for renewal: _____

1st Treatment

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⌀G

⌀D

