Grey Sling

Nursing Care

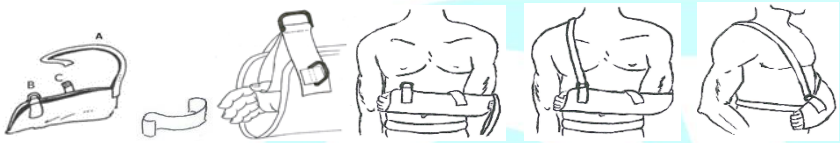
INTENDED USE

The **Grey Sling** is a medical device which is intended to be used for the immobilization of the arm/shoulder by injuries or fractures.

COMPOSITION

Product composed of three-layer polyamide and polyurethane material, with polypropylene rings. Polyester finger-support.

INSTRUCTIONS FOR USE



- Deploy the immobilizer deploying the strips.
- Put the arm inside. Support the thumb and adjust the tape B.
- The patient's arm must rest on the same square.
- Pass the tape fastening A above the arm and adjust the measure through the ring.
- Pass the subject strap D around the chest, immobilizing the arm.

PRODUCT CHARACTERISTICS

Designed for easy application and adjustment.
It has velcro system for closing and adjusting.
It supports the forearm and wrist. Incorporates support strip for the thumb.

Presents a vertical adjustable strap with hooks, its function is to distribute the weight of the arm across his back and shoulder for great comfort. The strap C facilities the adjustment of the arm sling.

PRECAUTIONS AND RECOMMENDATIONS

The use by more than a patient may provoke a cross-infection.
Do not use the product on injured skin.
Do not use the product if it is polluted, dirty or evidently deteriorated.

STORAGE AND SHELF-LIFE

Keep away from sunlight. Keep dry.
Shelf-life: 5 years.

PRODUCT DIMMENSIONS

In order to choose the correct size, measure the length of the forearm from the elbow to the knuckles.

REFERENCE	MODEL	L (cm)	W (cm)	L. Strap (cm)	W. Strap (cm)
1000720	XL	49	19	Chest: 140 Neck: 110	5
1000721	L	46	17	Chest: 140 Neck: 110	5
1000722	M	42	16	Chest: 140 Neck: 110	5
1000723	S	31	14	Chest: 75 Neck: 75	5
1000724	XS	26	11	Chest: 75 Neck: 75	5

CONFIGURATION			
REFERENCIA	UNITIES BOX	MEASUREMENTS BOX (cm)	WEIGHT BOX* (kg)
1000720	10 un/box	31x21x23	1.14
1000721	10 un/box	31x21x23	0.97
1000722	10 un/box	31x21x23	0.77
1000723	10 un/box	30x22x15	0.7
1000724	10 un/box	30x22x15	0.7

* TOLERANCES: Peso ± 0.2 Kg

IN COMPLIANCE WITH:

- Directive 93/42/EEC of the Council, Directive 2007/47/EC of the European Parliament and of the Council and Regulation (EU) 2017/745 of the European Parliament and of the Council.
- Class I, non-sterile (according to Annex IX of Directive 93/42 and Annex VIII of Regulation 745/2017).
- Enterprise certified by ISO 13485 and ISO 9001.
- Medical Devices Manufacturer Operating License No.: 3689-PS.