Fastening Vest

INTENDED USE

The **Fastening Vest** is designed to provide maximum safety and comfort to the patient.

Especially suitable for hospital areas where patients are constantly posts and lifted out of bed or wheelchair, also used in geriatrics and intensive care units..

COMPOSITION

Vest body: Polyamide 100%
Straps: non woven Polyester 100%

INSTRUCTIONS FOR USE

- After choosing the appropriate size, put the vest on the patient. Remember that the V-shaped opening should be in front of the vest.
- Cross the vest over the front and insert the left belt through the right groove.
- Tie the ribbons at the neck and make sure that the vest is positioned correctly.
- Position the patient in bed or in a wheelchair and tie the straps on opposite sides.

PRODUCT CHARACTERISTICS

Versatility in implementation, adapting to each patient. Comfort for nurses and patients.
Easy to apply, adjustable. Adapting to each patient.
The back safety straps allows an additional patient adjustment. Increase security.
The body vest allows air circulation. Fresh and comfortable.
Extra-long straps to reach the frame of the bed. Flexibility in implementation.
Polyester straps with additional sewn: strong product, durability. Reliable product.

PRECAUTIONS AND RECOMMENDATIONS

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

STORAGE AND SHELF-LIFE

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

UNIT PRODUCT MEASUREMENTS

REFERENCE	MODEL	L (cm)	W (cm)	STRAP L (cm)
1015670	LARGE (BLUE)	56	50	135
1015660	MEDIUM (GREEN)	47	48	156
1015650	SMALL (WHITE)	41	48	150

CONFIGURATION			
REFERENCE	**UNITS BOX**	**MEASUREMENTS BOX (cm)**	**WEIGHT BOX* (kg)**
1015670	10 un/box	31x21x30	2.7
1015660	10 un/box	31x21x30	2.2
1015650	10 un/box	31x21x30	1.7

TOLERANCE: Weight ±10%

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.