Anti-decubitus cushion

INTENDED USE

The **Anti-decubitus cushions** are a medical device, which is used to prevent bedsores in patients who have to stay long in chair.

COMPOSITION

Filling with silicone hollow polyester fibre.
Synthetic upper part and bottom 100% cotton flame retardant.

INSTRUCTIONS FOR USE

- Remove the cushion from the packaging and always hold it on the side.
- Place the cushion in the seat area.

PRODUCT CHARACTERISTICS

The filling of silicon polyester fiber is designed to achieve three basic objectives:

1) Recovery and immediate reaction of the fibers to any pressure applied to them, thus achieving an optimal distribution of weight.
2) Its light weight makes it very handy to use and transfer.
3) The set of fibers provide constant ventilation avoiding accumulation and moisture absorption.

The hollow central part allows transpiration and prevents pressure on the affected part when the patient must remain seated.
Hand wash with cold water and non-abrasive products. After washing let dry horizontally.

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)	DIAMETER (cm)
5010000	SQUARED	44	44	-
5010001	HORSESHOE	44	44	-
5010002	SQUARED WITH ORIFICE	44	44	-
5010003	RING	-	-	49.5

CONFIGURATION			
REFERENCE	**UNITS BAG**	**MEASUREMENTS BOX (cm)**	**WEIGHT BOX* (kg)**
5010000	1 un/bag	52x52x12	0.8
5010001	1 un/bag	52x52x12	0.6
5010002	1 un/bag	52x52x12	0.6
5010003	1 un/bag	52x52x12	0.6

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.