Dotted Anti-Bedsore Pillow

INTENDED USE

The **Dotted Anti-Bedsore Pillow** is a medical device, which is used to prevent bedsores in patients who have to stay long in chair

In addition, the cushion can be placed in a waterproof cover to prevent it from being damaged by liquid spills.

COMPOSITION

The pillow is formed by cotton 100% fireproof and filling with silicone hollow fibre.
Waterproof cover polyamide 100%.

INSTRUCTIONS FOR USE

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

PRODUCT CHARACTERISTICS

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

1) Recovery and immediate reaction of the fibres 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 fibres 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)
5020100	DOTTED ANTI-BEDSORE PILLOW	48	48
5020200	COVER	49	51

CONFIGURATION			
REFERENCE	**UNITS BAG**	**MEASUREMENTS BOX (cm)**	**WEIGHT BOX* (kg)**
5020100	1 un/bag	49x50x7	0.9
5020200	1 un/bag	35x37	0.1

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.

Sistema de Gestión ISO 9001:2015 ISO 22716:2007 www.tuv.com ID 9105078921