منع الحركة مع حماية اليد

Immobilizer with Hand Protection

INTENDED USE

The **Immobilizer with Hand Protection** is indicated for patients undergoing hand surgery, resuscitation, and every time that is required a strong subject of the hands, given an elevated level of comfort and safety for the patient.

مثبت الحركة مع حماية اليد للمرضى الذين يخضعون لجراحة اليد والإنعاش ، مع توفير مستوى عالٍ من الراحة والأمان للمريض.

COMPOSITON

Product composed of polyester mesh, with velour velcro strips and velcro system. It has polyester filling and three-layer material in polyamide, polyether and cotton base. Polypropylene ring.

المنتج يتكون من شبكة بوليستر ، مع أشرطة قطيفة ونظام فيلكرو. تحتوي على حشوة بوليستر ومادة ثلاثية الطبقات من مادة البولي أميد والبولي إيثر والقطن. حلقة من مادة البولي بروبيلين.

INSTRUCTIONS FOR USE

- Put the patient's hand in order to keep the fingers separated in each space. The mesh should cover the entire hand.
- Fix, first, the velcro strips on the metacarpal area.
- Fix the soft foam strip around the whist using the velcro lock.
- Wrap the strip around the protector and fix firmly.
- Tie the strip to the bed or another fixed part.

PRODUCT CHARACTERISTICS

Provided with a padded envelope with thick surface based on: synthetic fibres that prevent mould and bed sores. Fire-resistant. The stuffing not returns to his original form, for better protection of the palm of your hand.
Elaborated with special breathable tissue, that allows the transpiration of the skin.

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	LENGHT (cm)	WIDTH (cm)	STRIP LENGHT (cm)
1090300	25	17	168

CONFIGURATION			
REFERENCE	UNITS BOX	MESUREMENTS BOX (cm)	WEIGHT BOX* (kg)
1090300	16 un/box	38x21x35	2.23

TOLERANCIA: ± 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.

Sistema de Gestión ISO 9001:2015 ISO 22716:2007