GIMA
PROFESSIONAL MEDICAL PRODUCTS

ENGLISH

ANTI-DECUBITUS LINE IN SILICONE HOLLOW FIBRE

WARNINGS

Before carrying out any operation, visually check the integrity of the device. If the product shows any abnormalities, immediately contact the point of sale.

Erregi S.a.s. is not liable in case of improper use of the device.

In the event of a serious accident involving the use of the device, please report the incident to the manufacturer and to the competent authority of the member state where the user and/or patient lives.

DESCRIPTION

Cushion for the prevention of decubitus in 100% cotton, with padding from silicone hollow fibre. Some devices are with Velcro fasteners. The high resilience and breathability allow an effective action in the prevention of the onset of pressure ulcers.

INDICATION

Decubitus prevention for wheelchair-bound patients in order to decrease pressure peaks, reduce shear and friction forces and promote air circulation.

PRECAUTIONS FOR USE

Medical device indicated for the prevention of decubitus, to be used following a clinical evaluation carried out by a specialist doctor. Place the cushion on the wheelchair seat, making sure that the Velcro fasteners, where present, are positioned on the backrest side. Then secure the cushion to a rigid wheelchair structure using the Velcro fasteners. Although the device is indicated for the prevention of decubitus, it cannot control all the factors that favour its onset. It is therefore recommended to regularly monitor the affected area, to move the patient frequently, and to pay particular attention to hygiene. For this purpose, it is advisable to consult the protocols and guidelines for the prevention of pressure ulcers. Always wear personal protective clothing when using the device. Device not fireproof

CONTRAINDICATIONS

No contraindications were found except in cases of ascertained sensitivity to the component materials

MATERIAL

Padding: Silicone hollow fibre
Cover: 100% cotton

WASHING INSTRUCTIONS

The manufacturer is not liable for any changes in the technical features of the aid in the event of failure to observe the washing methods.

DISINFECTION

Use medical surgical products.

STORAGE

Store away from sources of light, heat, humidity.

DISPOSAL

Dispose of the device in compliance with the current regulations on environmental protection and recycling.

REF	Product code	MD	Medical Device
LOT	Lot number		Manufacturer
CE	Medical Device compliant with Regulation (EU) 2017/745		Consult instructions for use
	Keep away from sunlight		Keep in a cool, dry place
	Date of manufacture	LATEX	Not made with natural rubber latex
UDI	Unique identifier		

DO NOT remove the label from the Medical Device, as it allows tracing the supplied product (Manufacturer, UDI, batch number, product code).

GIMA WARRANTY TERMS

The Gima 12-month standard B2B warranty applies.

REF **08/13 (GIMA 28555)**
13/11 (GIMA 28556)
13/04 (GIMA 28557)

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MD CE

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