GIMA
PROFESSIONAL MEDICAL PRODUCTS

ENGLISH

CONTAINMENT BELT

WARNINGS
Before performing any operation, visually check the integrity of the device. If the product shows any anomalies, immediately contact the point of sale where the purchase was made.
Erregi sas assumes no responsibility for improper use of the device.
In the event of a serious incident related to the use of the device, please report the incident to the manufacturer and the competent authority of the member state where the user and/or patient is located.
DESCRIPTION
Bed restraint belts are devices designed to ensure the safety of patients who require therapy and do not actively cooperate. Made from highquality cotton straps, these belts use a locking system with a magnetic lock, which can be opened with a magnetic key to ensure easy and safe management.
Double bed belt: consists of two straps; the inner strap should be placed around the patient's waist and fastened with a magnetic lock, while the outer strap is anchored to the bedrails using ribbons and magnetic lock.
This model allows the patient to maintain a supine or lateral position, depending on the therapeutic needs.
Single bed belt: composed of a single strap to be placed around the patient's waist, fastened with a locking block. The two straps with hookand- loop fasteners allow the belt to be anchored to the bed's bedrails. The single belt is designed to allow only the supine position for the patient.
Double bed belt with inguinal strap: composed of three straps, the inguinal strap and the inner strap are placed around the patient's waist, while the outer strap is fastened to the bedrails. As in the previous models, the closing is done with a magnetic lock, which can be opened with a magnetic key. This model allows the patient to maintain both supine and lateral positions.
INTENDED USE
Devices used for the restraint of bedridden, non-cooperative adult patients.
INDICATIONS AND PRECAUTIONS FOR USE
The device should only be applied by medical and paramedical personnel.
The belt should not be applied to the pelvis or chest.
Devices with a magnetic closure system should not be used on patients who have a heart pacemaker, as it could lead to tachycardia.
Ensure the belt is properly fitted to the patient; a flat hand should be able to pass between the patient and the belt. Always check that the closures are securely fastened; if this does not happen, the patient's safety may be compromised. If the closures do not open, the ribbons can be cut with cutters or scissors.
Always use the device while the patient is wearing protective clothing. The device is not fireproof.
CONTRAINDICATIONS
No contraindications have been detected, except in cases of proven sensitivity to the component materials.
COMPOSITION
Cotton
WASHING INSTRUCTIONS

40°			
WASH AT MAX 40°C	DO NOT BLEACH	DO NOT IRON	DO NOT TUMBLEDRY
P	USE ANY SOLVENT EXCEPT TETRACHLORETHYLENE		

Attention: remove keys and blocks before washing, as contact with liquids may compromise their functionality The manufacturer is not responsible for any changes in the technical specifications of the device in case of non-compliance with the washing instructions.

DISINFECTION
Use products for medical-surgical use.
STORAGE
Store in a place protected from light, heat, and moisture.
DISPOSAL
Dispose of the device in accordance with the applicable environmental protection regulations and separate waste collection.

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
CH REP	Authorized Representative in Switzerland		Distributed by
UDI	Unique Device Identifier		

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

GIMA WARRANTY TERMS
The Gima 12-month standard B2B warranty applies.

REF	Produktkod	MD	Medicinteknisk produkt
LOT	Batchnummer		Tillverkare
CE	Medicinteknisk produkt som uppfyller kraven i förordning (EU) 2017/745		Läs bruksanvisningen
	Förvaras skyddat från solljus		Förvaras svalt och torrt.
	Tillverkningsdatum	LATEX	Inte tillverkad av naturgummilatex.
CH REP	Auktoriserad representant i Schweiz		Distribueras av
UDI	Unik enhetsidentifierare		

Ta INTE bort etiketten från den medicintekniska produkten, eftersom den tillåter spårbarheten av den levererade produkten (Tillverkare, UDI, satsnummer, produktkod).

GARANTIVILLKOR GIMA
Man tillämpar standard garanti B2B Gima på 12 månader.

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REF **24/07/01 - 24/07** S-M **- 24/07** L-XL

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MD CE

LATEX

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