
STECCOBENDE IN NEOPRENE
NEOPRENE SPLINTS
ATTELLES EN NÉOPRÈNE
FÉRULAS DE NEOPRENO
TALAS DE NEOPRENE
NEOPRENSKE UDLAGE
ATELE DE NEOPREN

È necessario segnalare qualsiasi incidente grave verificatosi in relazione al dispositivo medico da noi fornito al fabbricante e all'autorità competente dello Stato membro in cui si ha sede.

All serious accidents concerning the medical device supplied by us must be reported to the manufacturer and competent authority of the member state where your registered office is located.

Il est nécessaire de signaler tout accident grave survenu et lié au dispositif médical que nous avons livré au fabricant et à l'autorité compétente de l'état membre où on a le siège social.

Es necesario informar al fabricante y a la autoridad competente del Estado miembro en el que se encuentra la sede sobre cualquier incidente grave que haya ocurrido en relación con el producto sanitario que le hemos suministrado.

É necessário notificar ao fabricante e às autoridades competentes do Estado-membro onde ele está sediado qualquer acidente grave verificado em relação ao dispositivo médico fornecido por nós

Orice accident grav produs, privitor la dispozitivul medical fabricat de firma noastră, trebuie semnalat producătorului și autorității competente în statul membru pe teritoriul căruia își are sediul utilizatorul

Potrebno je prijaviti svaku ozbiljnu nezgodu koja se dogodila u vezi s isporučenim medicinskim proizvodaču i nadležnom tijelu države članice u kojoj se nalazi

REF 04SN0040 (34680) - 04SN0020 (34681)
04SN0010 (34682) - 04SN0050 (34683)
04SN0030 (34684) - 04KSBRO5 (34685)



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Made in Italy



COMPOSITION

Neoprene - Acrylic Velcro fasteners
The device materials are latex free.

INDICATIONS AND INTENDED USE

Support, protection and immobility of the limb or of the area to which it is applied. In case of long-term care / use, it is recommended to remove or loosen the device for short periods in order to aerate the affected part.

INSTRUCTIONS

Apply the device and secure it with Velcro, ensure that the application has been correctly performed (the end with the Velcro of a different colour should be up), the metal core could be folded or removed depending on the requirement.

Non-sterile device, do not apply in direct contact with damaged skin.



Warnings

Use extreme caution when applying.

Application by medical specialists or according to medical personnel's indications is recommended. Before applying, inform the doctor, who prescribes the device, about possible disorders or health problems.

If you should experience any allergic reaction, that may be expected to result from the use of this product, consult a doctor.

Not sterile device.

CARE AND CLEANING

Store in a cool, dry and clean place, away from direct sunlight and heat sources.

Reusable device, hand wash not above 30°C, do not dry clean, do not bleach, do not tumble dry, do not iron.

Lot number and manufacture date are on the label.

Class I Medical Device

Dispose of according to local laws in force.

Symbols

	Product code		Manufacturer		Date of manufacture
	Lot number		Consult instructions for use		Expiration date
	Medical Device compliant with Regulation (EU) 2017/745		Medical Device		Non-sterile

GIMA WARRANTY TERMS

The Gima 12-month standard B2B warranty applies.