

GB - USER MANUAL FOR REUSABLE SURGICAL INSTRUMENTS

These instruments are intended to be used by medical practitioners who are specially trained on how to use and care of them. The incorrect use, poor or inappropriate maintenance can rapidly lead to deterioration of the instruments. The first time, and after every use it is recommended to clean, dry and sterilise the instruments. The instruments must always be cleaned prior to sterilisation. For automated cleaning use only washing equipment with approved and certified detergents. For manual cleaning use approved and certified detergents, brush and running water. Always follow instructions on how to use the detergent; clean the instruments both when

open and closed; rinse for 3 minutes and check that water also enters and exits the blind holes several times. Use completely demineralised water in the final rinse phase. Instruments that are not dried could suffer damage by corrosion. Always dry the instruments. After cleaning, and before sterilisation it is recommended that you treat the instruments with physiologically safe oil, especially the tips, connectors, terminals and all moving parts. Also make sure that the product does not come into contact with acids or other aggressive disinfectants that could corrode it. The instruments must be sterilized in autoclave by moist heat sterilization method at 132°C for 15 minutes prior to use in surgery. It is recommended that sterilization temperatures should not exceed 137°C. The process of steam sterilisation must take place in accordance with EN ISO 17664. In the context of validating the sterilisation process, check the suitability of the specific measures for drying. The humidity in the container can cause the instruments to rust. Often bad, and insufficient drying, is due to the incorrect positioning of the load and the use of unsuitable types of cloths for drying. There is no advice about maximum number of sterilisation cycles, this depends largely on the state of the product. Instruments that show signs of corrosion must be discarded immediately. Always perform a visual inspection for damage or signs of wear: sharp edges must be free of dents and with continuous edges; there should be no distortion of instruments with long parts; the instruments that are part of a larger assembly, must be checked together with other assembly components; always check the rotating movement of hinges, which must not have excessive play; always check that the locking systems are working. Whilst the instruments are under warranty, repairs and/or replacement of all defective parts, for reasons that are accepted by the manufacturer, are free, with the exception of costs for labour, travel, transport, packing, etc. Damage caused by improper use of the product is not covered by the warranty.

GIMA WARRANTY TERMS

The Gima 12-month standard B2B warranty applies.

WARNINGS

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.

Simboli / Symbols / Symboles / Symbole / Simbolos / Símbolos / Σύμβολα / حرف

	IT Fabbricante GB Manufacturer FR Fabricant ES Fabricante PT Fabricante DE Hersteller GR Παραγωγός SA الشركة المصنعة
	IT Data di fabbricazione GB Date of manufacture FR Date de fabrication ES Fecha de fabricación PT Data de fabrico DE Herstellungsdatum GR Ημερομηνία παραγωγής SA تاريخ التصنيع
	IT Non sterile GB Non-sterile FR Pas stérile ES No estéril PT Não estéril DE Nicht steril GR Όχι αποστειρωμένο SA ليس معقم
	IT Attenzione: Leggere e seguire attentamente le istruzioni (avvertenze) per l'uso GB Caution: read instructions (warnings) carefully FR Attention: lisez attentivement les instructions (avertissements) ES Precaución: lea las instrucciones (advertencias) cuidadosamente PT Cuidado: leia as instruções (avisos) cuidadosamente DE Achtung: Anweisungen (Warnungen) sorgfältig lesen GR Προσοχή: διαβάστε προσεκτικά τις οδηγίες (ενστάσεις) SA الحذر: قراءة التعليمات (التحذيرات) بعناية
	IT Leggere le istruzioni per l'uso GB Consult instructions for use FR Consulter les instructions d'utilisation ES Consultar las instrucciones de uso PT Con- sulte as instruções de uso DE Gebrauchsanweisung beachten GR Διαβάστε προσεκτικά τις οδηγίες χρήσης SA اقرأ بدقة وحرص تعليمات الاستخدام
	IT Conservare al riparo dalla luce solare GB Keep away from sunlight FR À conserver à l'abri de la lumière du soleil ES Conservar al amparo de la luz solar PT Guardar ao abrigo da luz solar DE Vor Sonneneinstrahlung geschützt lagern GR Κρατήστε το μακριά από ηλιακή ακτινοβολία SA يحفظ بعيدا عن أشعة الشمس
	IT Conservare in luogo fresco ed asciutto GB Keep in a cool, dry place FR À conserver dans un endroit frais et sec ES Conservar en un lugar fresco y seco PT Armazenar em local fresco e seco DE An einem kühlen und trockenen Ort lagern GR Διατηρείται σε δροσερό και στεγνό περιβάλλον SA يحفظ في مكان بارد وجاف

REF	IT Codice prodotto GB Product code FR Code produit ES Código producto PT Código produto DE Erzeugniscode GR Κωδικός προϊόντος SA كود المنتج
LOT	IT Numero di lotto GB Lot number FR Numéro de lot ES Número de lote PT Número de lote DE Chargennummer GR Αριθμός παρτίδας SA رقم الدفعة
MD	IT Dispositivo medico GB Medical Device FR Dispositif médical ES Producto sanitario PT Dispositivo médico DE Medizinprodukt GR Ιατροτεχνολογικό προϊόν SA جهاز طبي
CE	IT Dispositivo medico conforme alla Direttiva 93/42/CEE; in accordo alla MDCG 2020-2, sarà reso conforme al Regolamento (UE) 2017/745, per cambio classe. GB Medical Device complies with Directive 93/42/EEC; in accordance with MDCG 2020-2, they shall be made compliant with Regulation (EU) 2017/745, for change of class. FR Dispositif médical conforme à la directive 93/42 / CEE; conformément au MDCG 2020-2, ils seront mis en conformité avec le règlement (UE) 2017/745, par changement de classe. ES Dispositivo médico según a la Directiva 93/42 / CEE; en conformidad con la MDCG 2020-2, se adecuarán al Reglamento (UE) 2017/745 para el cambio de clase. PT Dispositivo médico em conformidade com a Diretiva 93/42/ CEE, de acordo com a MDCG 2020-2, deverão estar em conformidade com o Regulamento (UE) 2017/745 devido a mudança de classe. DE Medizinprodukt gemäß Richtlinie 93/42/CEE; sie werden in Übereinstimmung mit der MDCG 2020-2 konform mit der Verordnung (EU) 2017/745 für den Wechsel der Klasse gestaltet. GR Ιατρική συσκευή σύμφωνα με την οδηγία 93/42 / CEE; Σύμφωνα με το MDCG 2020-2, θα έχουν συμμορφωθεί με τον Κανονισμό (ΕΕ) 2017/745, λόγω αλλαγής κατηγορίας. SA جهاز طبي يتوافق مع التوجيه 93/42/CEE
UDI	IT Identificatore univoco del dispositivo GB Unique device identifier FR Identifiant unique de l'appareil ES Identificador de dispositivo único PT Identificador exclusivo do dispositivo DE Unique Device Identifier (Eindeutige Kennung des Geräts) GR Μοναδικό αναγνωριστικό συσκευής SA - معرّف فريد للجهاز
CH REP	IT - Rappresentante autorizzato in Svizzera GB - Authorized Representative in Swiss FR - Représentant autorisé au Suisse ES - Representante autorizado en Suizo PT - Representante autorizado no Suíço DE - Autorisierter Vertreter im Schweizer GR - Εξουσιοδοτημένος Αντιπρόσωπος στο Ελβετός SA - الممثل المعتمد في سويسرا
UK REP	IT Rappresentante autorizzato nel Regno Unito GB Authorized Representative in the UK FR Représentant autorisé au Royaume-Uni ES Representante autorizado en el Reino Unido PT Representante autorizado no Reino Unido DE Autorisierter Vertreter im Vereinigten Königreich GR Εξουσιοδοτημένος Αντιπρόσωπος στο Η.Β SA الممثل المعتمد في المملكة المتحدة



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