For recognized legal manufacturer, refer to product label.

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VICRYL™

ar خيوط جراحية | bg КОНЕЦ | cs SICÍ MATERIÁL | da SUTUR | de NAHTMATERIAL | el ΡΑΜΑΚΑ | en SUTURE | es SUTURA | et ÕMBLUSAMATERJAL | fi OMMELLANE | fr FIL DE SUTURE | hu VARRÓANYAG | it SUTURA | lt ŠILKA | lv VĪTNIŅŠĶASU DIEGS | mt ħjieħ tad-draħ | nl HECHTDRAAD | no SUTUR | pl NICI CHIRURGICZNE | pt FIO DE SUTURA | ro FIR DE SUTURĂ | ru ШОВНЫЙ МАТЕРИАЛ | sk CHIRURGICKÁ NIť | sv SUTUR | tr SÜTÜR | zh-cn 缝线 | zh-tw 縫合線

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02/2021

CE 2797

Instructions for use

en

VICRYL™
POLYGLACTIN 910
STERILE SYNTHETIC ABSORBABLE SURGICAL SUTURE

DESCRIPTION
VICRYL™ Suture is a sterile, synthetic, absorbable, surgical suture composed of a copolymer made of 90% glycolide and 10% L-lactide. The empirical formula of the copolymer is $(C_2H_2O_2)_x(C_3H_4O_3)_y$. VICRYL™ Suture is available as a coated or an uncoated suture.
VICRYL™ Suture is coated with a mixture composed of equal parts of copolymer of glycolide and lactide (Polyglactin 370) and calcium stearate. Polyglactin 910 copolymer and Polyglactin 370 with calcium stearate have been found to be nonantigenic, nonpyrogenic and elicits only a slight tissue reaction during absorption.
VICRYL™ Suture is available undyed and dyed with D&C Violet #2 (Color Index 60725) to enhance visibility in the surgical field.
VICRYL™ Suture is available in a range of gauge sizes and lengths, types and sizes, and in presentations non-mounted or attached to needles of various types and sizes as described in the HOW SUPPLIED section.
VICRYL™ Suture complies with the requirements of the European Pharmacopoeia (Ph. Eur.) for Sterile Synthetic Absorbable Braided Sutures and United States Pharmacopeia (USP) for Absorbable Surgical Suture, except for a slight oversize in some gauges as shown in the table below.

USP Suture Size	Metric Suture Size	Maximum Oversize
9-0 (uncoated braid only)	0.3	0.005 mm
6-0	0.7	0.008 mm
5-0	1	0.016 mm
4-0	1.5	0.017 mm
3-0	2	0.018 mm
2-0	3	0.004 mm
0	3.5	0.022 mm

The European Pharmacopoeia recognizes units of measure Metric and Ph. Eur. Sizes as equivalent, which is reflected on the labeling.

INDICATIONS
VICRYL™ Suture is indicated for use in general soft tissue approximation and/or ligation including use in ophthalmic surgery, peripheral nerve anastomosis and microsurgery for vessels less than 2 mm diameter. The safety and effectiveness of VICRYL™ Suture in cardiovascular tissue has not been established.

APPLICATIONS
Sutures should be selected and implanted depending on patient condition, surgical experience, surgical technique and wound size.

PERFORMANCE / ACTIONS
VICRYL™ Suture elicits an initial, minimal inflammatory reaction in tissues and ingrowth of fibrous connective tissue. Progressive loss of tensile strength and eventual absorption occurs by means of hydrolysis, where the copolymer degrades to glycolic and lactic acids which are subsequently absorbed and metabolized in the body. Absorption begins as a loss of tensile strength followed by a loss of mass. All of the original tensile strength is lost by five weeks post implantation. Absorption is essentially complete between 56 and 70 days.

Days Implantation	Approximate % Original Strength Remaining
14 Days	75%
21 Days USP 6-0 (Metric 0.7) and larger	50%
21 Days USP 7-0 (Metric 0.5) and smaller	40%
28 Days USP 6-0 (Metric 0.7) and larger	25%

CONTRAINDICATIONS
These sutures, being absorbable, should not be used where extended approximation of tissues under stress is required.

WARNINGS
Users should be familiar with surgical procedures and techniques involving absorbable sutures before employing VICRYL™ Suture for wound closure, as risk of wound dehiscence may vary with the site of application and the suture material used. Medical professionals should consider the *in vivo* performance (under the PERFORMANCE / ACTIONS section) when selecting a suture.
Acceptable surgical practice should be followed for the management of contaminated or infected wounds.
As this is an absorbable suture material, the use of supplemental non-absorbable sutures should be considered by the surgeon in the closure of the sites which may undergo expansion, stretching or distension, or which may require additional support.
Do not resterilize/reuse. Reuse of this device (or portions of this device) may create a risk of product degradation, which may result in device failure and/or cross-contamination, which may lead to infection or transmission of blood-borne pathogens to patients and users. As an absorbable, this suture may act transiently as a foreign body. Like all foreign bodies, this product may potentiate infection.

PRECAUTIONS
In handling this or any other suture material, care should be taken to avoid damage. Avoid crushing or crimping damage due to application of surgical instruments such as forceps or needle holders.
Skin sutures which must remain in place longer than 7 days may cause localized irritation and should be snipped off or removed as indicated.
Under some circumstances, notably orthopedic procedures, immobilization of joints by external support may be employed at the discretion of the surgeon.
Consideration should be taken in the use of absorbable sutures in tissues with poor blood supply as suture extrusion and delayed absorption may occur.
Subcutaneous sutures should be placed as deeply as possible to minimize the erythema and induration normally associated with the absorption process.
This suture may be inappropriate in elderly, malnourished or debilitated patients, or in patients suffering from conditions which may delay wound healing.
As with any suture material, adequate knot security requires the standard surgical technique of flat, square ties with additional throws as warranted by surgical circumstance and the experience of the surgeon. The use of additional throws may be particularly appropriate when knotting monofilament sutures.
Care should be taken to avoid damage when handling surgical needles. Grasp the needle in an area one-third (1/3) to one-half (1/2) of the distance from the attachment end to the point. Grasping in the point area could impair the penetration performance and cause fracture of the needle. Grasping at the attachment end could cause bending or breakage. Reshaping needles may cause them to lose strength and be less resistant to bending and breaking.
Users should exercise caution when handling surgical needles to avoid inadvertent needle stick injury. Broken needles may result in extended or additional surgeries or residual foreign bodies. Inadvertent needle sticks with contaminated surgical needles may result in the transmission of blood-borne pathogens. Discard used needles in "sharps" containers.

ADVERSE REACTIONS
Adverse reactions associated with the use of this device include wound dehiscence, calculi formation in urinary or biliary tracts when prolonged contact with salt solutions such as urine or bile occurs, transient inflammatory tissue reaction, and transient local irritation at the wound site.

STERILITY
VICRYL™ Suture is sterilized by ethylene oxide gas. Do not resterilize. Do not use if package is opened or damaged. Discard opened, unused sutures.

STORAGE
No special storage conditions required. Do not use after expiry date.

HOW SUPPLIED
Please note that not all sizes are available in all markets. Please contact your local sales representative for size availability.
VICRYL™ Suture is available as sterile strands in sizes 10-0 through 6 (metric sizes 0.2-4.0), in a variety of lengths, with and without permanently attached needles.
VICRYL™ Suture is also available as a monofilament in sizes 9-0 and 10-0 (metric sizes 0.3 and 0.2).
VICRYL™ Suture is also available as sterile strands attached to CONTROL RELEASE™ removable needles which enable the needles to be pulled off instead of being cut off.
VICRYL™ Suture is available in one, two or three dozen units per box.

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ar تجاه استخدامها | bg За употребата | cs Před použitím | da Inden brug | de Vor Gebrauch | el Πριν τη χρήση | en Prior to use | es Antes de usar | et Kasutada enne | fi Ennen käyttöä | fr Avant utilisation | hr Prije uporabe | hu Használat előtt | it Prima dell'uso | lt Prieš naudojimą | lv Pirms lietošanas | mt Qabel tużu | nl Vóór gebruik | no Før bruk | pl Przed użyciem | pt Antes de utilizar | ro Înainte de folosire | ru Перед использованием | sk Pred použitím | sl Pred uporabo | sv Före användning | tr Kullanımdan önce | zh-cn 请勿重复使用 | zh-tw 不得重複使用

ar تاريخ انتهاء الصلاحية | bg Да се употребя до / cs Datum použitelnosti / da Anvendes inden / de Verwendbar bis / el Ημερομηνία λήξης / en Use by / es Usar antes de / et Kasutada enne / fi Käytettävä viimeisen käyttöpäivän / fr À utiliser avant / hu Felhasználási idő / it Utilizzare entro / lt Naudojamas iki / lv Izlietojams līdz / mt Użu sal-/ nl Vóór de vervaldatum / no Brukes innen / pl Wykorzystać do / pt Prazo de validade / ro A se utiliza până la / ru Использовать до / sk Spotrebuje do / sl Uporabi do / sv Använd före / tr Son kullanma tarihi / zh-cn 有效期至 / zh-tw 有效期限

LOT ar رقم الدفعة | bg Партиден код / cs Číslo šarže / da Batchkode / de Chargenbezeichnung / el Κωδικός παρτίδας / en Batch code / es Código de lote / et Partii kood / fi Eräkoodi / fr Numéro de lot / hu Tételszám / it Numero lotto / lt Gamybinės partijos kodas / lv Partijas kods / mt Partiju kodu / nl Lotnummer / no Partinummer / pl Kod partii / pt Código de lote / ro Cod lot / ru Номер партии / sk Kód šarže / sv Satskod / tr Parti kodu / zh-cn 批号 / zh-tw 批號

STERILE EO ar تم تعقيمها بإستخدام أكسيد الإيثيلين / bg Стерилизирано с етиленов оксид / cs Sterilizováno ethylenoxidem / da Steriliseret med ethylenoxid / de Sterilisation mit Ethylenoxid / el Αποστειρωμένο με οξείδιο του αιθυλενίου / en Sterilized using ethylene oxide / es Esterilizado usando óxido de etileno / et Steriliseeritud etüleenoksiidiga / fi Sterilisoitu etyleenioksidilla / fr Stérilisé à l'oxyde d'éthylène / hu Etilénoxiddal sterilizált / it Sterilizzato a ossido di etilene / lt Sterilizuojamas etileno oksidu / lv Sterilizēts ar etilēnoksīdu / mt Isterilizzat bil-użu ta' ossidu ta' etilene / nl Steriel door middel van ethylenoxide / no Sterilisert med etylenoksid / pl Wyjałowiono tlenkiem etylenu / pt Esterilização por óxido de etileno / ro Sterilizat cu ajutorul unui oxid de etilenă / ru Стерилизовано этиленоксидом / sk Sterilizované etylénoxidom / sv Steriliserad med etylenoxid / tr Etilen oksit kullanılarak sterilize edilmiştir / zh-cn 使用环氧乙烷灭菌 / zh-tw 使用環氧乙烷滅菌

REF ar رقم الكتالوج | bg Каталожен номер / cs Katalogové číslo / da Varenummer / de Katalognummer / el Κωδικός καταλόγου / en Catalogue Number / es Número de catálogo / et Kataloogi number / fi Tuotenumero / fr Numéro de référence au catalogue / hu Katalógusszám / it Numero di catalogo / lt Katalogo numeris / lv Katalogā numurs / mt Numru tal-katalogu / nl Catalogusnummer / no Katalognummer / pl Numer katalogowy / pt Número de catálogo / ro Numărul de catalog / ru Номер по каталогу / sk Katalógové číslo / sv Katalognummer / tr Katalog numarası / zh-cn 目录编号 / zh-tw 目錄型號

⚠ ar تحذير | bg ВНИМАНИЕ / cs UPOZORNĚNÍ / da BEMÆRK / de ACHTUNG / el ΠΡΟΣΟΧΗ / en CAUTION / es ATENCIÓN / et ETTEVAATUSE / fi HUOMIO / fr ATTENTION / hu MIGYELEM / it ATTENZIONE / lt DĖMESIO / lv UZMANĪBU / mt ATTENZJONI / nl LET OP / no OBS / pl UWAGA / pt ATENÇÃO / ro ATENȚIE / ru ВНИМАНИЕ / sk UPOZORNENIE / sv FÖRSIKTIGHET / tr DIKKAT / zh-cn 注意 / zh-tw 注意

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ar تعقيم بأشعة أكسيد الإيثيلين / bg Да не се стерилизира повторно / cs Neresterilizujte / da Må ikke resteriliseres / de Nicht erneut sterilisieren / el Μην επαναποστειρώνετε / en Do not resterilize / es No resterilizar / et Mitte resteriliseerida / fi Ei saa sterilisoida uudelleen / fr Ne pas restériliser / hu Nem sterilizálja újra / it Non risterilizzare / lt Nesterilizuoti / lv Nesterilizēt atkārtoti / mt Ma tisterilizzax qiegħ / nl Niet opnieuw steriliseren / no Skal ikke resteriliseres / pl Nie wyjałowiać ponownie / pt Não resterilizar / ro A nu se resteriliza / ru Не использовать повторно / sk Nesterilizujte opakovane / sv Får ej återsterilisera / tr Tekrar kullanmayın / zh-cn 请勿重复使用 / zh-tw 不得重複使用

ar لا تستخدم إذا كانت العبوة تالفة | bg Да не се използва, ако опаковката е повредена / cs Nepoužívejte, je-li obal poškozený / da Må ikke anvendes, hvis pakningen er beskadiget / de Nicht verwenden, wenn Verpackung beschädigt / el Μη χρησιμοποιείτε εάν η συσκευασία έχει υποστεί ζημιά / en Do not use if package is damaged / es No usar si el envase está dañado / et Mitte kasutada, kui pakend on kahjustatud / fi Ei saa käyttää, jos pakkaus on vaurioitunut / fr Ne pas utiliser si l'emballage est endommagé / hu Ne használja fel, ha a csomagolás megsérült / hr Non usare se la confezione è danneggiata / it Non usare se la confezione è danneggiata / lt Nenaudoti, jeigu pakinėta pažeista / lv Neizlietot, ja iepakojums ir bojāts / mt Ħież użu jekk l-imballaġ ikun ħsarat / nl Niet gebruiken als de verpakking beschadigd is / no Ikke bruker hvis pakningen er skadet / pl Nie używać, jeśli opakowanie jest uszkodzone / pt Não utilizar se a embalagem estiver danificada / ro Nu se utilizează dacă ambalajul este deteriorat / ru Не использовать, если упаковка повреждена / sk Nepoužívajte, ak bolo balenie poškodené / sv Får ej användas om förpackningen är skadad / tr Ambalajı hasarlıysa kullanmayın / zh-cn 如果包装有破损，请勿使用 / zh-tw 如果包裝有破損，請勿使用

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ar صنعة للتصدير | bg Производител / cs Výrobce / da Producent / de Hersteller / el Κατασκευαστής / en Manufacturer / es Fabricante / et Tootja / fi Valmistaja / fr Fabricant / hu Gyártó / lt Pagamintojas / lv Ražotājs / mt Fabrikant / no Produsent / pl Producent / pt Fabricante / ro Producător / ru Производитель / sk Výrobca / sv Tillverkare / tr Imalatçı / zh-cn 制造商 / zh-tw 製造商

EC REP ar الممثل المعتمد في المجتمع الأوروبي | bg Упълномощен представител в Европейската общност / cs Autorizovaný zástupce v Evropském společenství / da Autoriseret repræsentant (EU) / de Bevollmächtigter in der Europäischen Gemeinschaft / el Εξουσιοδοτημένος αντιπρόσωπος στην Ευρωπαϊκή Κοινότητα / en Authorized Representative in the European Community / es Representante autorizado en la Comunidad Europea / et Ametlik esindaja Euroopa Ühenduses / fi Valtuutettu edustaja Euroopan yhteisössä / fr Représentant autorisé dans la Communauté européenne / hr Ovlašteni predstavnik u Europskoj zajednici / hu Meghatalmazott képviselet az Európai Közösségben / lt Atstovaujantis asmuo Europos Bendrijoje / lv Pilnvarotais pārstāvis Eiropas Kopienā / mt Rappresentant awtorizzat fil-Komunità Ewropea / nl Gemachtigd vertegenwoordiger in de Europese Unie / no Autorisert representant (EU) / pl Autoryzowany przedstawiciel we Wspólnocie Europejskiej / pt Representante autorizado na Comunidade Europeia / ro Reprezentant autorizat în Uniunea Europeană / ru Уполномоченный представитель в ЕС / sk Autorizovaný zástupca v Európskom spoločenstve / sv Auktoriserad representant i Europeiska gemenskapen / tr Avrupa Topluluğu Yetkili Temsilcisi / zh-cn 欧洲共同体授权代理 / zh-tw 歐洲共同體授權代表

CE 2797 ar علامة الجماعة المطابقة لمنتج الموجهات للسوق الأوروبية لقطار الأوروبي يوفر التعريف من قبل مجموعة جهة فريق معادل بها من الاتحاد الأوروبي وهذا يعني أن الدفع مطابق للشروط الأساسية لتلك الموجهات تحقيقًا لالتزامها الكلية المقررة من السوق الأوروبية لقطارات يوفر / EECA2793

bg CE маркировка и идентифициращият номер на нотифицираното органа. Продуктът отговаря на съществените изисквания на Директива 93/42/ЕИО, за медицинските изделия / cs Označení CE a identifikační číslo notifikované osoby. Výrobek vyhovuje základním požadavkům směrnice 93/42/EHS o zdravotnických prostředcích / da CE-mærkning og identifikationsnummer for bemyndiget organ. Produktet opfylder de væsentlige krav i Direktiv 93/42/EØF om medicinsk udstyr / de CE-Kennzeichnung und Kennnummer der Benannten Stelle. Das Produkt entspricht den grundlegenden Anforderungen der Richtlinie 93/42/EWG des Rates über Medizinprodukte / el Σήμανση CE και αριθμός ανεγκρίσεως του κοινοποιημένου οργανισμού. Το προϊόν πληροί τις ουσιώδεις απαιτήσεις της οδηγίας 93/42/ΟΚ περί ιατροτεχνολογικών προϊόντων / en CE-mark and Identification number of Notified Body. The product meets the essential requirements of Medical Device Directive 93/42/EEC / es Marca CE + número de identificación del organismo notificado. Este producto cumple los requisitos esenciales de la directiva de productos sanitarios 93/42/CEE / et CE-tähis ja teavitatud asutuse registreerimisnumber. Toode vastab meditsiinisadmete direktiivi 93/42/EMÜ põhinõuetele / fi CE-merkintä ja ilmoitetun laitoksen tunnusnumero. Tuote noudattaa lääkinnällisiä laitteita annetun direktiivin 93/42/ETY säännöksiä / fr Marquage CE et numéro d'identification de l'organisme notifié. Le produit est conforme aux exigences essentielles de la Directive sur les dispositifs médicaux 93/42/CEE. / hr CE je ili a brojevi grupi testiše izdana. A temnik megfelel az orvostechnikai eszközök biztonságosságról szóló 93/42/EGK számú irányelv előírásainak / ik Ahachos CE e numero di identificazione dell'organismo notificato. Il prodotto è conforme ai requisiti essenziali della direttiva 93/42/CEE sui dispositivi medici / lt CE – ženklas ir atsakingo kūrybinės organizacijos identifikacinis numeris. Šis gaminys atitinka Medicininės priemonės, žvaigždės, žvaigždės 93/42/EB/EB Direktyvos esminius reikalavimus / lv CE zīme un paziņotās iestādes identifikācijas numurs. Šo izstrādājumu atbilst medicīniskās ierīces direktīvas 93/42/EEK pamatprasībām / mt CE-marking u-identifikazzjonimumer tan-ġarantiġ ħaġġ. Ħaġġ. �митвенсалимын нигеҗ талаптарына жавап бирелә / ka ᲙᲝᲓᲘᲡ და ᲛᲘᲜᲘᲨᲜᲘᲡ ᲜᲘᲨᲜᲘᲡ 93/42/EEC ᲛᲝᲗᲮᲝᲕᲜᲘᲡ ᲫᲘᲠᲘᲗᲐᲓᲘ ᲛᲝᲗᲮᲝᲕᲜᲔᲑᲘ

Symbol Definitions in IFU ar تعريفات الرموز في تعليمات الاستخدام / bg Дефиниции на символите в Инструкциите за употреба / cs Definice symbolů v návodu / da Forklaring på symboler i brugsanvisningen / de Erklärung der Symbole in der Gebrauchsanleitung / el Ορισμοί συμβόλων στις Οδηγίες χρήσης / en Symbol Definitions in IFU / es Definición de los símbolos en las instrucciones de uso / et Tähendused definitsioonid kasutusjuhendis / fi Tuotesoitteisten symbolien merkitykset / fr Définitions des symboles utilisés dans le mode d'emploi / hu A tájékoztatóban előforduló jelek magyarázata / ik Legenda dei simboli nelle Istruzioni per l'uso / lt Žymėjimai naudojamo instrukcijoje / lv Apzīmējumu definīcijas lietošanas instrukcijā / mt Definitie van symbolen in de gebruiksaanwijzing / no Symbolforklaring i bruksanvisning / pl Definicje symboli użytych w instrukcji użytkowania / pt Definições dos símbolos nas IDU / ro Definiții de simboluri din cadrul instrucțiunilor de utilizare / ru Значения символов в инструкции / sk Definície symbolov použitých v NIoP / sv Symboldefinitioner i bruksanvisningen / tr Kullanma Talimatındaki Sembol Açıklamaları / zh-cn 和IFU的符号的定义 / zh-tw 使用說明中的標誌定義

CE 2797 lt CE ženklas ir notifikuotosios įstaigos identifikavimo numeris. Gaminys atitinka pagrindinius medicininių prietaisų 93/42/EB direktyvos reikalavimus / hr CE oznake i identifikacijski broj ovlaštene ustanove. Ovaj proizvod ispunjava zahtjeve medicinske direktive 93/42/EEZ / pl Oznaczenie CE i numer identyfikacyjny jednostki notyfikowanej. Produkt spełnia wymagania zasadnicze dyrektywy o wyrobach medycznych 93/42/EWG / pt Marcação CE e Número de identificação do organismo notificado. O produto cumpre os princípios orientadores da diretiva sobre dispositivos médicos 93/42/CEE / ro Marcajul CE și numărul de identificare a organismului notificat. Acest produs corespunde cerințelor esențiale ale Directivei privind dispozitivele medicale 93/42/CEE / ru Знак соответствия «CE» и идентификационный номер уведомленного органа. Данное изделие отвечает основным требованиям директивы о медицинском оборудовании 93/42/ЕЭС / sk Označenie CE a identifikačné číslo notifikovaného orgánu. Produkt spĺňa základné požiadavky smernice 93/42/EHS o zdravotných pomôckach / sv CE-märke och identifikationsnummer för ansvarsfullt organ. Produkten uppfyller de väsentliga kraven i direktivet för medicinsk utrustning, 93/42/EEG / tr CE işareti ve Onaylanmış Kuruluş kimlik numarası. Ürün, 93/42/AET sayılı Tıbbi Cihazlar Direktifi'nin başlıca gerekliliklerine uygundur / zh-cn CE标志和证明机构的识别号。产品符合医疗设备指令93/42/EEC的基本要求 / zh-tw CE標誌和認證機構的識別號。產品符合醫療設備指令93/42/EEC的基本要求

ar عدد الوحدات | bg Брой единици / cs Počet kusů / da Antal enheder / de Stückzahl / el Αριθμός συσκευών / en Number of units / es Número de unidades / et Seadmete arv / fi Yksikköjen lukumäärä / fr Nombre d'unités / hu Az egységek száma / ik Numero di unità / lt Gaminių skaičius / lv Vienību skaits / mt Aantal eenheden / no Antall enheter / pl Liczba sztuk / pt Número de unidades / ro Numărul de unități / ru Число единиц / sk Počet kusov / sv Antal enheter / tr Ünite sayısı / zh-cn 件数 / zh-tw 數量

CR/X ar عدد الخيوط في كل خطاف (X) CONTROL RELEASE™ | bg CONTROL RELEASE™ (X=броят конци в едрия опаковка) / cs CONTROL RELEASE™ (X=počet vláken v jednom balení) / da CONTROL RELEASE™ (X=Antal tråde pr. pakke) / de CONTROL RELEASE™ (X=Anzahl der Stränge pro Paket) / el CONTROL RELEASE™ (X=Ο αριθμός κλώνων ανά συσκευή) / en CONTROL RELEASE™ (X=The number of strands per packet) / es CONTROL RELEASE™ (X=número de hebras por envase) / et CONTROL RELEASE™ (X=pakendis olevate kuidude arv) / fi CONTROL RELEASE™ (X=Fäkkaisessa olevien lankojen lukumäärä) / fr CONTROL RELEASE™ (X=nombre de brins par boîte) / hu CONTROL RELEASE™ (X=A kartonizsák szálma csomagonként) / hr CONTROL RELEASE™ (X=broj niti u jednom balenju) / ik CONTROL RELEASE™ (X=Numero di fili per confezione) / lt CONTROL RELEASE™ (X=Ritės skaičius paketėje) / lv CONTROL RELEASE™ (X=Vītņu skaitlis paketē) / mt CONTROL RELEASE™ (X=Ħajt ħaġa ħaġa fil-pakkett) / nl CONTROL RELEASE™ (X=Aantal draden per verpakking) / no CONTROL RELEASE™ (X=Antall tråder i hver pakke) / pl CONTROL RELEASE™ (X=Liczba nici w opakowaniu) / pt CONTROL RELEASE™ (X=numéro de filamento per pacote) / ro CONTROL RELEASE™ (X=numărul de filamente per pachet) / ru CONTROL RELEASE™ (X=число нитей в упаковке) / sk CONTROL RELEASE™ (X=počet vlákien v jednom balení) / sv CONTROL RELEASE™ (X=antalet suturer per förpackning) / tr CONTROL RELEASE™ (X=Paket başına iplik sayısı) / zh-cn CONTROL RELEASE™ （X=每包胶线的数目）/ zh-tw CONTROL RELEASE™ （X=每包胶囊量）

MS/X ar عدد الخيوط في كل خطاف (X) | bg Многоконечно влакно (X=броят влакна в едрия опаковка) / cs Multifilament (X=počet vláken v jednom balení) / da Multitråd (X=Antal tråde pr. pakke) / de Mehrfachstrang (X=Anzahl der Stränge pro Paket) / el Πολλαπλός κλώνος (X=Ο αριθμός κλώνων ανά συσκευή) / en Multistrand (X=The number of strands per packet) / es Multihebra (X=número de hebras por envase) / et Multistrand (X=pakendis olevate kuidude arv) / fi Monisäikeinen (X=Pakkaussa olevien lankojen lukumäärä) / fr Multibrins (X=nombre de brins par boîte) / hu Multistrand (Többszálú) (X=A kartonizsák száma csomagonként)

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ik Multifilo (X=numero di fili per confezione) / kk Көп түйін (X=Оқшау орауыштағы түйіндердің санағы) / ko 다중 가닥(X=패킹당 가닥 수) / lt Kelios ritės (X=ritės skaičius paketėje) / lv Daudzpavediens (X=pavediens skaits iepakojumā) / mt Mewtiene doubes (X = ħut santal doubes per wrappking) / no Flertråder (X = antall tråder i hver pakke) / pl Wielopasmowy (X=Liczba pasm w opakowaniu) / pt Multifilamentos (X=número de filamento per pacote) / ro Multifilament (X=numărul de filamente per pachet) / ru Многоволоконный (X=число нитей в пачке) / sk Viacvláknové (X=počet vlákien v jednom balení) / sv Multistrand (X = antalet suturer per förpackning) / tr Çok ipliklı (X=Paket başına iplik sayısı) / zh-cn 多股 （X=每包胶线的数目）/ zh-tw 多股 （X=每包胶囊量）

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ar جرعة واحدة لكل عبوة | bg ЕКЗЕМПЛЯР/ПОЛДЕНЕТЕ/ОТДЕЛЕТЕ ТУК / cs ZDE ODLOUPNUTI/ZOVINUTÍ/OTEVŘETÍ / da ÉRÅKK/LØFT/ÅBN HER / de HIER ABZIEHEN/ AUFREIßEN/ÖFFNEN / el ΑΠΟΚΟΛΛΗΣΤΕ/ΑΝΟΙΞΤΕ/ΑΝΟΙΓΤΕ ΕΔΩ / en PEEL/LIFT/OPEN HERE / es DESPRENDER/ELEVAR/ABRIR AQUÍ / et NÕMELA/TÕSTA/AVA SIIT / fi AVAA TÄSTÄ / fr DÉTACHER/SOULEVER/OUVRIR ICI / hr REZZA I LEMBLE BEJAN/TOLA FILI ITT / ik ÉRRAPPARE/SOLLEVARE/APRIRE QUI / kk ОССЫ ЖЕРДІ/ЕШ СЫҒЫП/ЕРІП ОСЫ ЖЕРДІ/ЕШ СЫҒЫП/ЕРІП / ko 여기를 떼어 내고/들어올려/개봉하십시오 / lt NULIUTI / PAKELTI / ATIDARYTI ČIA / lv ATLOBĪT/PAVILKT UZ AUGŠU/ATVērt ŠEIT / nl HIER OPENTREKKEN/OMHOOGTREKKEN/OPENEN / no FJERNELSES/ LØFT/ÅPNES HER / pl OTWIERAĆ/UNIESIĆ/OTWIERAĆ TUTAJ / pt DESTACAR/LEVANTAR/ABRIR AQUI / ro DECLANȘAȚI/RIDICAȚI/DESCHIDEȚI AICI / ru СОРВИТЬ/ПРИПОДНЯТЬ/ОТКРЫТЬ ЗДЕСЬ / sk TU ODLOUPNUT/NADVIHNUT/OTVORIŤ / sv DRA ALT/TOPPRA HÄR / tr BURADAN SOYUN/KALDIRIN/AÇIN / zh-cn 到开/提起/此处开启 / zh-tw 剝開/提起/此處開啟

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ar موقع هذا | bg СЕЗАРЕТЕ ТУК / cs ZDE ODTRHNUTÍ / da ÉRV HER / de HIER AUFREISSEN / el ΚΧΕΤΕ ΕΔΩ / en TEAR HERE / es RASGAR AQUÍ / et REBI SIIT / fi REVI TÄSTÄ / fr DÉCHIRER ICI / hu ZÁKATSA LE IT / ik ROMPERE QUI / kk ОСЫ ЖЕРДЕН ЖЫРТЫҢЫЗ/ЫҢ / ko 여기를 �faster 오십시오 / lt PLĖSTI ČIA / lv ATPLĒST ŠEIT / nl HIER AFSCHEUREN / no RIVES HER / pl UŻYJ ROZERWAĆ / pt RASGAR AQUI / ro RUPEȚI ACI / ru ОТОРВАТЬ ЗДЕСЬ / sk TU ODTRHNÚŤ / sv RIV HÄR / tr BURADAN YIRTIN / zh-cn 由此撕开 / zh-tw 在此撕開

ar مصطلحات الرموز في تعليمات الاستخدام / bg Дефиниции на символите в Инструкциите за употреба / cs Definice symbolů v návodu / da Forklaring på symboler i brugsanvisningen / de Erklärung der Symbole in der Gebrauchsanleitung / el Ορισμοί συμβόλων στις Οδηγίες χρήσης / en Symbol Definitions in IFU / es Definición de los símbolos en las instrucciones de uso / et Tähendused definitsioonid kasutusjuhendis / fi Tuotesoitteisten symbolien merkitykset / fr Définitions des symboles utilisés dans le mode d'emploi / hu A tájékoztatóban előforduló jelek magyarázata / ik Legenda dei simboli nelle Istruzioni per l'uso / kk ПАЙДАЛАНУ НҰСҚАУЛЫҒЫНДАҒЫ / ko 사용 설명서 내 기호 설명 / lt Žymėjimai naudojamo instrukcijoje / lv Apzīmējumu definīcijas lietošanas instrukcijā / mt Definitie van symbolen in de gebruiksaanwijzing / no Symbolforklaring i bruksanvisning / pl Definicje symboli użytych w instrukcji użytkowania / pt Definições dos símbolos nas IDU / ro Definiții de simboluri din cadrul instrucțiunilor de utilizare / ru Значения символов в инструкции / sk Definície symbolov použitých v NIoP / sv Symboldefinitioner i bruksanvisningen / tr Kullanma Talimatındaki Sembol Açıklamaları / zh-cn 和IFU的符号的定义 / zh-tw 使用說明中的標誌定義