Box size: 2.25 x 5
Overall size: 18 x 25.5
Folds to: 2.25 x 4.25

COVIDIEN

Maxon™ and Maxon™ CV

Monofilament Absorbable Suture

1300372

EN

BEFORE USING PRODUCT, READ THE FOLLOWING INFORMATION THOROUGHLY.

DESCRIPTION

Maxon™ and Maxon™ CV monofilament polyglyconate synthetic absorbable sutures are prepared from a copolymer of glycolic acid and trimethylene carbonate. Maxon™ and Maxon™ CV synthetic absorbable sutures are dyed with D&C Green No. 6 to increase visibility and are also available clear (undyed).

Maxon™ and Maxon™ CV synthetic absorbable sutures meet all requirements established by the United States Pharmacopeia (USP) and the European Pharmacopeia (EP) for synthetic absorbable surgical sutures, except for minor variations in suture diameter.

Such variations are:

Maximum Suture Oversize in Diameter (mm) from USP & EP		
USP Size	USP Size Designation (mm)	Maximum Overage (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0.20-0.249	0.055
2-0	0.30-0.339	0.055
0	0.35-0.399	0.070
1	0.40-0.499	0.070

INDICATIONS

Maxon™ and Maxon™ CV sutures are indicated for use as absorbable sutures in general soft tissue approximation and/or ligation, including use in pediatric cardiovascular tissue, where growth is expected to occur, and in peripheral vascular tissue.

Maxon™ and Maxon™ CV sutures are not indicated for use in adult cardiovascular tissue, ophthalmic surgery, microsurgery, and neural tissue.

ACTIONS

Maxon™ and Maxon™ CV synthetic absorbable sutures elicit a minimal acute inflammatory reaction in tissues, which is followed by gradual encapsulation of the suture by fibrous connective tissue. Progressive loss of tensile strength and eventual absorption of Maxon™ and Maxon™ CV synthetic absorbable sutures occurs by means of hydrolysis where the suture is broken down and subsequently absorbed and metabolized by the body.

Studies indicate that tensile strength for Maxon™ and Maxon™ CV sutures is approximately 80% of the original suture strength at one week post-implant. Approximately 75% of the original suture strength remains at two weeks, 65% at three weeks, 50% at four weeks and 25% at six weeks post-implant. Absorption of Maxon™ and Maxon™ CV synthetic absorbable sutures is minimal until about the 60th day after implantation. Absorption is substantially complete within six months.

CONTRAINDICATIONS

The use of this suture is contraindicated in patients with known sensitivities or allergies to its components.

This suture, being absorbable, should not be used where extended approximation of tissue is required or fixation of permanent cardiovascular prostheses or synthetic grafts.

WARNINGS

Do not resterilize. Sterile unless packaging has been opened or damaged. Discard open, unused sutures. Store at room temperature. Avoid prolonged exposure to elevated temperature.

In surgery of the urinary and biliary tracts, care should be taken to avoid prolonged contact of this, or any other, suture with salt solutions, as calculus formation may result.

Users should be familiar with surgical procedures and techniques involving absorbable sutures before employing Maxon™ and Maxon™ CV synthetic absorbable sutures for wound closure, as the risk of wound dehiscence may vary with the site of application and the suture material used.

Acceptable surgical practice must be followed with respect to drainage and closure of contaminated or infected wounds.

The use of this suture may be inappropriate for patients with any conditions which, in the opinion of the surgeon, may cause or contribute to delayed wound healing.

The use of supplemental nonabsorbable sutures should be considered by the surgeon in the closure of sites subject to expansion, stretching or distention, or requiring additional support.

PRECAUTIONS

Under some circumstances, notably orthopedic procedures, immobilization by external support may be employed at the discretion of the surgeon.

In handling this or any other suture material care should be taken to avoid damage from handling. Avoid crushing or crimping damage due to application of surgical instruments such as forceps or needle holders.

Adequate knot security requires the accepted surgical technique of flat, square ties, with additional throws as warranted be surgical circumstances and the experience of the surgeon.

The use of additional throws may be particularly appropriate when knotting monofilaments.

Care should be exercised when knotting sutures used in continuous mass or layered closure of the abdomen, i.e. non-slip locking knots should be used at both ends of the continuous closure.

The commonly used looped-to-stand knot during termination of a running suture is a configuration, which may lead to slippage. Extreme care must be utilized to ensure that multiple square knots are well secured.

Sutures in vaginal mucosa or skin that remain in place for extended periods may be associated with localized irritation and should be removed as indicated.

Since any foreign material in the presence of bacterial contamination may enhance bacterial infectivity, standard surgical practice should be followed with respect to drainage and closure of infected wounds.

Dispose of contaminated suture and packaging materials utilizing standard hospital procedures and universal precautions for biohazardous waste.

ADVERSE REACTIONS

Adverse effects, which may be associated with the use of this product, include: wound dehiscence, failure to provide adequate wound support in sites where expansion, stretching or distension occur; failure to provide adequate wound support in patients with conditions which may delay wound healing; tissue granulation or fibrosis; wound suppuration and bleeding, as well as sinus formation; localized irritation when skin sutures are left in place for 7 or more days; calculi formation when prolonged contact with salt solutions occurs; enhanced bacterial infectivity; minimal acute inflammatory reaction; and pain, edema, and erythema at the wound site.

HOW SUPPLIED

Maxon™ sutures are available in USP and E.P sizes 1 (4 Metric) through 7-0 (0.5 Metric). Maxon™ CV sutures are available only in USP and EP sizes 4-0 (1.5 Metric) through 7-0 (0.5 Metric). Maxon™ and Maxon™ CV sutures are dyed with D&C Green No. 6. Maxon™ sutures are also available in clear (Undyed). The sutures are supplied sterile, in pre-cut lengths; non-needled or affixed to various needle types, using both permanent and removable needle attachment techniques. The sutures are available in box quantities of one, two and three dozen.

FR

Suture résorbable monofilament

AVANT D'UTILISER CE PRODUIT, LIRE ATTENTIVEMENT LES INFORMATIONS CI-DESSOUS.

Dépassement maximal du diamètre de la suture (en mm) établi par l'USP et l'EP.		
Taille USP	Désignation de taille USP (mm)	Dépassement maximum (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30 à 0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

DE

Monofiles resorbierbares Nahtmaterial

VOR GEBRAUCH DES PRODUKTES SIND DIE FOLGENDEN INFORMATIONEN SORGFÄLTIG DURCHZULESEN.

Maximale Übergröße des Nahtmaterial-Durchmessers (mm) in Bezug auf USP und EP & EP		
USP-Format	USP-Fadenstärke Nennngröße (mm)	Maximale Übergröße (mm)
7-0	0,050-0,069	0.015
6-0	0,070-0,099	0.025
5-0	0,10-0,149	0.035
4-0	0,15-0,199	0.040
3-0	0,20-0,249	0.055
2-0	0,30-0,339	0.055
0	0,35-0,399	0.070
1	0,40-0,499	0.070

IT

Sutura assorbibile in monofilamento

PRIMA DELL'USO LEGGERE ATTENTAMENTE LE SEGUENTI INFORMAZIONI.

Massimo calibro di sutura sovradimensionato rispetto a quanto stabilito dalla USP (diametro mm) e dalla EP.		
Calibro USP	Classificazione del calibro secondo l'USP (mm)	Eccedenza massima (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

ES

Sutura reabsorbible monofilamento

ANTES DE USAR EL PRODUCTO, LEA DETENIDAMENTE LA SIGUIENTE INFORMACIÓN.

Excedente máximo del diámetro (mm) de la sutura respecto a lo indicado por USP y EP		
Tamaño USP	Designación de tamaño de USP (mm)	Diferencia máxima (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0.20-0.249	0.055
2-0	0.30-0.339	0.055
0	0.35-0.399	0.070
1	0.40-0.499	0.070

PT

Sutura absorvível monofilamentosa

ANTES DE USAR O PRODUTO, LEIA ATENTAMENTE ESTAS INFORMAÇÕES.

Tamanho máximo da sutura em diâmetro (mm) de acordo com a USP e EP		
Tamanho USP	Designação de tamanho USP (mm)	Excesso máximo (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

NL

Absorbeerbare monofilamenthechtingen

LEES DE VOLGENDE INFORMATIE ZORGVULDIG DOOR VOORDAT U HET PRODUCT GEBRUIKT.

Maximale diameterafwijking (mm) met betrekking tot USP & EP		
Vereiste USP-afmeting	Aanduiding USP-afmeting (mm)	Maximale overmaat (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

SV

Monofil resorberbar sutur

LÄS NEDANSTÅENDE INFORMATION NOGA INNAN PRODUKTEN TAS I BRUK.

Maximal diameteravvikelse (mm) från USP och Eur. Ph.		
USP-storlek	USP-storleksangivelse (mm)	Maximal avvikelse (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

DA

Resorberbar monofilamentsutur

FØLGENDE ANVISNING SKAL LÆSES GRUNDIGT, INDEN PRODUKTET TAGES I BRUG.

Maksimal overstørrelse af suturdiameter (mm) i forhold til USP & EP		
USP-størrelse	USP-størrelsesbetegnelse (mm)	Maksimal overstørrelse (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

FI

Yksisäikeinen resorboituva ommelaine

SEURAAVAT OHJEET ON LUETTAVA HUOLELLISESTI ENNEN TUOTTEEN KÄYTTÖÄ.

Ommelaineen läpimitan enimmäisvaihtelu (mm) USP- ja EP-kokoon verrattuna		
USP-koko	USP-koko Nimitys (mm)	Enimmäisyli-ikä (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

EL

Μονόκλωνο, απορροφήσιμο ράμμα

ΠΡΙΝ ΧΡΗΣΙΜΟΠΟΙΗΣΕΤΕ ΤΟ ΠΡΟΪΟΝ, ΔΙΑΒΑΣΤΕ ΠΡΟΣΕΚΤΙΚΑ ΟΛΕΣ ΤΙΣ ΑΚΟΛΟΥΘΕΣ ΠΛΗΡΟΦΟΡΙΕΣ.

Μέγιστη υπέρβαση διαμέτρου ράμματος(σε mm) από τη USP & την EP		
Μέγεθος κατά USP	Υπεύθυνη μεγέθους κατά USP (mm)	Μέγιστη υπέρβαση μεγέθους (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,55
2-0	0,30-0,339	0,55
0	0,35-0,399	0,070
1	0,40-0,499	0,070

PL

Wchłanialne monofilamentowe nici chirurgiczne

PRZED ROZPOCZĘCIEM KORZYSTANIA Z PRODUKTU NALEŻY UWAŻNIE PRZECZYTAĆ PONIŻSZE INFORMACJE.

Maksymalny nadwymiar średnicy nici chirurgicznych (mm) z USP i EP		
Rozmiar USP	Oznaczenie rozmiaru USP (mm)	Maksymalny Nadwymiar (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0,20 – 0,249	0.055
2-0	0,30 – 0,339	0.055
0	0,35 – 0,399	0.070
1	0.40-0.499	0.070

TR

Monofilament Emilebilir Dikiş İpliği

ÜRÜNÜ KULLANMADAN ÖNCE, AŞAĞIDAKİ BİLGİLERİ DİKKATLİ ŞEKİLDE OKUYUN.

USP ve EP'ye göre Maksimum İplik Çapı Fazlalığı		
USP Boyutu	USP Boyutu Tanımı (mm)	Maksimum Büyük Boyut (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20 - 0,249	0,055
2-0	0,30 - 0,339	0,055
0	0,35 - 0,399	0,070
1	0,40-0,499	0,070

RU

Нить из абсорбируемого моноволокна

ПЕРЕД ТЕМ КАК ПРИСТУПИТЬ К ИСПОЛЬЗОВАНИЮ ЭТОГО ИЗДЕЛИЯ, ВНИМАТЕЛЬНО ОЗНАКОМЬТЕСЬ СО СЛЕДУЮЩЕЙ ИНФОРМАЦИЕЙ.

Максимальное превышение по диаметру (мм) согласно USP и EP		
Размер по USP	Обозначение размера USP (мм)	Максимальное превышение (мм)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20 - 0,249	0,055
2-0	0,30 - 0,339	0,055
0	0,35 - 0,399	0,070
1	0,40-0,499	0,070

CS

Jednovláknový vstřebatelný šicí materiál

PŘED POUŽITÍM VÝROBKU SI DŮKLADNĚ PŘEČTĚTE NÁSLEDUJÍCÍ INFORMACE.

Maximální rozměr přesahující standardní průměr šicího materiálu (mm) podle USP & EP		
Velikost podle USP	Označení velikosti podle USP (mm)	Maximální přesah (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20 – 0,249	0,055
2-0	0,30 – 0,339	0,055
0	0,35 – 0,399	0,070
1	0,40-0,499	0,070

HU

Egyszálas fonalú felszívódó fonalak

A TERMÉK HASZNÁLATA ELŐTT FIGYELMESEN OLVASSA EL AZ ALÁBBI TUDNIVALÓKAT.

Az USP & EP méretet meghaladó legnagyobb fonálátmérő (mm)		
USP Méret	USP méret kijelölése (mm)	Maximális többlet (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0,20 – 0,249	0.055
2-0	0,30 – 0,339	0.055
0	0,35 – 0,399	0.070
1	0.40-0.499	0.070

SK

Jednovláknové, syntetické, vstrebateľné nite

PRED POUŽITÍM PRODUKTU SI POZORNE PREČÍTAJTE NASLEDUJÚCE INFORMÁCIE.

Maximálny rozmer presahujúci štandard nite v priemere (mm) podľa USP a EP		
Veľkosť podľa USP	Popis veľkosti podľa USP (mm)	Maximálny presah (mm)
7-0	0,050 – 0,069	0,015
6-0	0,070 – 0,099	0,025
5-0	0,10 – 0,149	0,035
4-0	0,15 – 0,199	0,040
3-0	0,20 – 0,249	0,055
2-0	0,30 – 0,339	0,055
0	0,35 – 0,399	0,070
1	0,40 – 0,499	0,070

NO

Resorberbar monofilamentsutur

FØR PRODUKTET TAS I BRUK, MÅ FØLGENDE INFORMASJON LESES NØYE GJENNOM.

Maks. suturoverstørrelse i diameter (mm) fra USP & EP		
USP-størrelse	USP-størrelsebetegnelse (mm)	Maks. utvidelse (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

SL

Monofilamentna vpojna sutura

PRED UPORABO TEGA IZDELKA NATANČNO PREBERITE NASLEDNJE INFORMACIJE.

Največja čezmerna velikost premera suture (mm) po USP in EP		
Velikost USP	Oznaka velikosti USP (mm)	Največji čezmerni premer (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

ZH

单纤维可吸收缝合线

使用本产品之前，请务必先仔细阅读下面的信息。

比 USP和 EP 规定的缝合线直径超出的最大尺寸 (mm)		
USP 尺寸	USP 尺寸规格 (mm)	最大型号 (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0.20 - 0.249	0.055
2-0	0.30 - 0.339	0.055
0	0.35 - 0.399	0.070
1	0.40-0.499	0.070

ZH-T

單絲可吸收縫合線

使用本產品之前，請先讀完下列資訊。

縫合線最大直徑尺寸超過 USP 規定直徑尺寸 (mm) 以及 E.P 尺寸。		
USP 型號	USP 尺寸名稱 (mm)	最大超量 (mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0.20-0.249	0.055
2-0	0.30-0.339	0.055
0	0.35-0.399	0.070
1	0.40-0.499	0.070

KO

모노필라멘트 흡수성 봉합사

제품을 사용하기 전에 다음 내용을 자세히 읽으십시오.

USP 및 E.P의 봉합사 최대 직경(mm)		
USP 크기	USP 크기 지정 (mm)	최대 초과(mm)
7-0	0.050-0.069	0.015
6-0	0.070-0.099	0.025
5-0	0.10-0.149	0.035
4-0	0.15-0.199	0.040
3-0	0.20 - 0.249	0.055
2-0	0.30 - 0.339	0.055
0	0.35 - 0.399	0.070
1	0.40-0.499	0.070

BG

Еднонишков абсорбируем хирургически конец

ПРЕДИ ДА ИЗПОЛЗВАТЕ ТОЗИ ПРОДУКТ, ПРОЧЕТЕТЕ ВНИМАТЕЛНО СЛЕДВАЩАТА ИНФОРМАЦИЯ.

Макс. допустимо превишение на диаметъра (mm) според USP и EP		
USP размер	Желан USP размер (mm)	Макс. превишение (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

RO

Sutură absorbabilă cu monofilament

CITIŢI CU ATENŢIE INFORMAŢIILE DE MAI JOS ÎNAINTE DE A UTILIZA PRODUSUL.

Supradimensionarea maximă a suturii în diametru (mm) faţă de norma americană şi cea europeană		
Dimensiunea USP	Desemnarea dimensiunii USP (mm)	Surplusul maxim (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

ET

Resorbeeruv monofilamentne õmblusmaterjal

ENNE TOOTE KASUTAMIST LUGEGE TÄHELEPANELIKULT LÄBI JÄRGNEV TEAVE.

Õmbluse maksimaalne liigne diameeter (mm) (USP & E.P).		
USP suurus	USP suuruse märgistus (mm)	Maksimaalne ülesuurus (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

LV

Viendzīslas absorbējama šuve

PIRMS PRODUKTA LIETOŠANAS RŪPĪGI IZLASIET TĀLĀK SNIEGTO INFORMĀCIJU.

Maksimālais USP un EP šuves virsizmēra diametrs (mm)		
USP izmērs	USP izmēra apzīmējums (mm)	Maksimālais "overage" (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

HR

Monofilamentni resorptivni konac

PRIJE KORIŠTENJA PROIZVODA PAŽLJIVO PROČITAJTE SLJEDEĆE INFORMACIJE.

Maksimalno prekoračenje promjera konca (u mm) – USP i EP		
Oznaka vličina prema USP	Veličina (u mm) prema USP	Maksimalno prekoračenje (u mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

SR

Monofilamentni resorptivni konac

PRE UPOTREBE PROIZVODA PAŽLJIVO PROČITAJTE SLEDEĆE INFORMACIJE.

Maksimalne pozitivne varijacije prečnika konca (mm) u odnosu na USP i EP		
USP veličina	USP oznaka veličine (mm)	Maksimalna pozitivna varijacija (mm)
7-0	0,050-0,069	0,015
6-0	0,070-0,099	0,025
5-0	0,10-0,149	0,035
4-0	0,15-0,199	0,040
3-0	0,20-0,249	0,055
2-0	0,30-0,339	0,055
0	0,35-0,399	0,070
1	0,40-0,499	0,070

LT

Vienagijis absorbuojamasis siūlas

PRIEŠ NAUDODAMI PRODUKTĄ, ATIDŽIAI PERSKAITYKITE PATEIKTĄ INFORMACIJĄ

Maksimalus siūlo skersmens viršijimas (mm) pagal USP ir EP		
USP dydis	USP dydžio žymėjimas (mm)	Maksimalus viršijimas (mm)
7-0	0,050–0,069	0,015
6-0	0,070–0,099	0,025
5-0	0,10–0,149	0,035
4-0	0,15–0,199	0,040
3-0	0,20–0,249	0,055
2-0	0,30–0,339	0,055
0	0,35–0,399	0,070
1	0,40–0,499	0,070

Do not use if package is opened or damaged.
Ne pas utiliser en cas d'endommagement ou d'ouverture de l'emballage.
Bei geöffneter oder beschädigter Verpackung nicht verwenden.
Non utilizzare se la confezione è aperta o danneggiata.
No usar el dispositivo si la envoltura está abierta o dañada.
Não utilizar se a embalagem estiver aberta ou danificada.
Niet gebruiken als de verpakking beschadigd of geopend is.
Får ej användas om förpackningen är öppnad eller skadad.
Må ikke anvendes, hvis emballagen er åbnet eller beskadiget.
Ei saa käyttää, jos pakkaus on avattu tai vaurioitunut.
Μη χρησιμοποιείτε αν η συσκευασία έχει ανοιχτεί ή υποστεί ζημιά.
Nie stosować, jeżeli opakowanie zostało otwarte lub uszkodzone.
Ambalaj açılmış ya da zarar görmüşse kullanmayın.
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Pokud je balení otevřené nebo poškozené, produkt nepoužívejte.
Tilos felhasználni, ha a csomagolás kinyílt vagy megsérült.
Nepoužívajte, ak je obal otvorený alebo poškodený.
Skal ikke brukes hvis emballasjen er åpnet eller skadet.
Ne uporabljajte, če je embalaža odprta ali poškodovana.
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Не използвайте, ако опаковката е отворена или повредена.
A nu se utiliza în cazul în care ambalajul este deschis sau deteriorat.
Ärge kasutage, kui pakend on avatud või kahjustatud.
Nelietot, ja iepakojums ir atvērts vai bojāts.
Ne koristite ako je pakiranje otvoreno ili oštećeno.
Ne koristiti ako je pakovanje otvoreno ili oštećeno.
Nenaudokite, jei pakuotė atidaryta arba pažeista.

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Rx ONLY

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