

3M Steri-Strip™

- Ⓜ Skin Closures Sterile and Non-Sterile
- Ⓜ Sutures cutanées stériles et non stériles
- Ⓜ Sutures cutanées, stériles et non stériles
- Ⓜ Hautverschlüsse steril und nicht-steril
- Ⓜ Suture cutanee sterili e non sterili
- Ⓜ Suturas cutáneas estériles y no estériles
- Ⓜ Suturas cutáneas estériles y no estériles
- Ⓜ Steriele en niet-steriele huidhechtstrips
- Ⓜ Hudförlutningar Steril och icke-steril
- Ⓜ Tapesuturer, sterile og ikke-sterile
- Ⓜ Sårlukningsstrips Sterile og usterile
- Ⓜ Haavansulkuteipit, steriilit ja ei-steriilit
- Ⓜ Fechos de Pele Estéreis e Não Estéreis
- Ⓜ Suturas cutáneas adesivas estéreis e não estéreis
- Ⓜ Ταυλές Συρραφής Δέρματος Αποστειρωμένες και Μη Αποστειρωμένες
- Ⓜ Jafowe i niejafowe paski do zamykania ran
- Ⓜ Steril és nem steril sebzáró csíkok
- Ⓜ Náplastové stehy sterilní a nesterilní
- Ⓜ Kožné náplastové stehy sterilné a nesterilné
- Ⓜ Trakci za zatvaranje ran na koži sterilni in nesterilni
- Ⓜ Haavasulgurid sterillsed ja mittesterillsed
- Ⓜ Ádas plásteri Sterili un nesterili
- Ⓜ Sterilios ir nesterilios odos žaizdų uždarymo juostelės
- Ⓜ Benzii pentru sutura leziunilor sterile și nesterile
- Ⓜ Пластырные кожные швы стерильные и нестерильные
- Ⓜ Стерильні та нестерильні пластирні смужки для зведення країв ран
- Ⓜ Trake za zatvaranje rana Sterilne i nesterilne
- Ⓜ Sterilni i nesterilni lenti za bezševno zatvaranje na rani
- Ⓜ Trake za zatvaranje rana sterilne i nesterilne
- Ⓜ Cilt Kapatıcıları Steril ve Steril Olmayan
- Ⓜ Зарарьсздандырылған және зарарьсздандырылмаған тері жапсырмалары
- Ⓜ 減菌および非減菌皮膚閉鎖材
- Ⓜ Shirita për mbylljen e plagëve, sterile dhe jo sterile
- Ⓜ Ленти за затворање на кожа Стерилни и нестерилни

MD CE 2797

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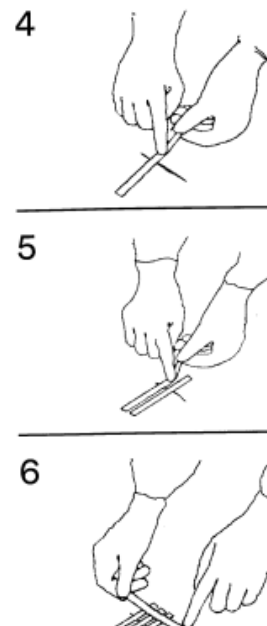
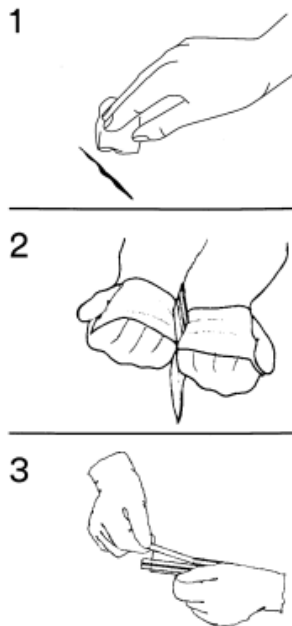
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Explanation of Symbols

Symbol Title	Symbol	Description and Reference
Manufacturer		Indicates the medical device manufacturer as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC. Source: ISO 15223, 5.1.1
Date of Manufacture		Indicates the date when the medical device was manufactured. Source: ISO 15223, 5.1.3
Use-by date		Indicates the date after which the medical device is not to be used. Source: ISO 15223, 5.1.4
Batch code		Indicates the manufacturer's batch code so that the batch or lot can be identified. ISO 15223, 5.1.5
Do not use if package is damaged or open		Indicates a medical device that should not be used if the package has been damaged or opened. Source: ISO 15223, 5.2.8
Do not re-use		Indicates a medical device that is intended for one use or for use on a single patient during a single procedure. Source: ISO 15223, 5.4.2
Caution		Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself. Source: ISO 15223, 5.4.4
Natural rubber latex is not present		Indicates natural rubber or dry natural rubber latex is not present as a material of construction within the medical device or the packaging of a medical device. Source: ISO 15223, 5.4.5 and Annex B

For more information see, HCBRegulatory.3M.com

3M™ Steri-Strip™



3M™ Steri-Strip™

EN

Skin Closures Sterile and Non-Sterile

Product Description

- 3M™ Steri-Strip™ Skin Closures are made of porous, nonwoven material. They are reinforced with filaments for strength.
- 3M™ Steri-Strip™ Blend Tone Skin Closures are made of porous, nonwoven material and are coated with a hypoallergenic adhesive.
- 3M™ Steri-Strip™ Elastic Skin Closures are made of porous, nonwoven, elastic material. They are designed to allow for tissue expansion and movement and are coated with a hypoallergenic adhesive.

Indications for Use

Steri-Strip skin closures are indicated for use as a skin closure device in the treatment of lacerations and surgical incisions. Steri-Strip skin closures may also be used in conjunction with skin sutures and staples or after their removal for wound support.

Contraindications

1. Steri-Strip skin closures are contraindicated where adhesion cannot be obtained. Potential causes of inadequate adhesion are presence of exudate, skin oils, moisture, or hair.
2. Use of Steri-Strip skin closures on infected wounds is contraindicated.
3. Steri-Strip skin closures are contraindicated for use in high tension wounds which cannot be easily approximated with fingers or forceps.

Warnings

- The development of postoperative edema may cause skin shearing, skin blistering, or loss of tape adhesion to occur at either end of the strip.
- Application of any surgical tape or adhesive skin closure may result in skin stripping upon removal.
- As with all adhesive products applied to the skin, a small percentage of individuals may experience hypopigmentation or hyperpigmentation following removal.
- Occasional cases of mild acne and folliculitis have been observed in testing on healthy volunteers.

Precautions

1. The skin should be clean, dry, and free of skin oils to assure good adhesion.
2. Do not apply skin closures under tension. Skin shearing, skin blistering, or loss of adhesion may result if excessive tension is applied.

Directions for Use

1. Clean and dry skin. If desired, a thin layer of Compound Benzoin Tincture may be applied to enhance strip adhesion.
2. Open package. Remove card and card end tab.
3. Grasp end of strip and lift strip straight upward from card.
4. When using as a skin closure device, without presence of skin suture or staple, appose skin edges using fingers or forceps. Place strips across wound, ensuring tensionless application.
5. Section larger wounds to distribute tension. Space strips approximately 1/4 inch (3 mm) apart.
6. Additional closures may be applied parallel to the wound and approximately 1/2 inch (12 mm) from closure ends. This may reduce shear force beneath the strip ends.

Storage/Shelf Life/Disposal

For best results, store at room temperature. Avoid excessive heat and humidity. For shelf life, refer to the expiration date printed on each package.

How Supplied

1. Supplied in individually packaged cards of sterile skin closures.
2. Store at room temperature. Avoid excessive heat or humidity.
3. Sterile unless envelope is damaged or open.
4. Steri-Strip skin closures may be provided to end users as sterilized via radiation or, when provided in some kits, sterilized via Ethylene Oxide (EO). Refer to label for sterilization type.

Please report a serious incident occurring in relation to the device to 3M and the local competent authority (EU) or local regulatory authority.

For further information, please contact your local 3M representative or contact us at 3M.com and select your country.