**Instructions for use**

**NYLON TAPE**
**STERILE**

**DESCRIPTION**
Nylon Tape is a flat woven or braided Nylon-(Polyamide 6,6) tape. Nylon Tape is available in various widths and lengths, non-needed only.

**PERFORMANCE**
Because of its flat woven structure, Nylon Tape is indicated for temporary ligation or retraction of tissues, organs or other anatomical structures during surgical procedures without injuring the tissue involved.

**INDICATIONS**
Nylon Tape is indicated for temporary ligation or retraction of tissues, organs or other anatomical structures during surgical procedures. The device is not intended to be implanted in the body.

**APPLICATION**
Nylon Tape is used for temporary retraction and/or fixing of organs or parts of organs (e.g. intestine) during an operation. After completion of the operation the tape should be removed and disposed of.

**CONTRAINDICATIONS**
There are no known contraindications. However, Nylon Tape is not indicated for implantation.

**WARNINGS / PRECAUTIONS**
Care should be taken during tissue retraction to avoid injury. Nylon Tape is provided as a sterile product. Do not resterilize. Do not use if packaging is opened or damaged. Discard opened unused products.

Reuse of this device (or portions of this device) may create a risk of product degradation and cross-contamination, which may lead to failure of treatment, infection or transmission of bloodborne pathogens to patients and users.

Safety and effectiveness of Nylon Tape have not been established for use as an implanted device. The risks associated with product implantation have not been established.

**ADVERSE REACTIONS**
None known.

**STERILITY**
Nylon Tape is sterilized by irradiation. Do not resterilize!
Do not use if package is opened or damaged! Discard open, unused product.

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

**SYMBOLS USED ON LABELING**

- **STERILE R**
  - Sterilized using irradiation

- **Do not reuse**
  - Do not resterilize

- **LOT**
  - Batch code

- **CE**
  - CE-mark and Identification Number of Notified Body. The product meets the essential requirements of Medical Device Directive 93/42/EEC

- **Authorized representative in the European Community**

- **Manufacturer**
  - See instructions for use

- **Number of Sachets in Box**

- **Catalogue Number**

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