

MOLLELAST® HAFT

Cohesive Bandage

REF 22629, 30063-30072, 30103-30105

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1. Composition of the product

This Mollelast® haft bandage consists of:

- viscose
- polyamide
- coated with natural latex (on both sides)

This product data sheet covers the following items:

REF 30063	Mollelast® haft	4 cm x 4 m stretched	packed individually
REF 30064	Mollelast® haft	6 cm x 4 m stretched	packed individually
REF 30065	Mollelast® haft	8 cm x 4 m stretched	packed individually
REF 30066	Mollelast® haft	10 cm x 4 m stretched	packed individually
REF 30067	Mollelast® haft	12 cm x 4 m stretched	packed individually
REF 30068	Mollelast® haft	4 cm x 20 m stretched	packed individually
REF 30069	Mollelast® haft	6 cm x 20 m stretched	packed individually
REF 30070	Mollelast® haft	8 cm x 20 m stretched	packed individually
REF 30071	Mollelast® haft	10 cm x 20 m stretched	packed individually
REF 30072	Mollelast® haft	12 cm x 20 m stretched	packed individually
REF 30103	Mollelast® haft	6 cm x 20 m	6 pcs./pack
REF 30104	Mollelast® haft	8 cm x 20 m	6 pcs./pack
REF 30105	Mollelast® haft	10 cm x 20 m	6 pcs./pack
REF 22629	Two-way elastic adhesive bandage 10 cm x 20 m stretched Loose, 40 pcs./pack		

2. Packaging, structure and composition

2.1 Unit container

REF 30063 - REF 30072, REF 30103 - REF 30105:

- see 2.2

REF 22629:

- see 2.3

2.2 Shelf container

REF 30063 - REF 30072, REF 30103 - REF 30105:

- a cardboard core (cellulose)

- a folding cardboard box (cellulose)

2.3 Transit container

REF 30063 - REF 30072, REF 30103 - REF 30105:

- corrugated cardboard box (cellulose)

REF 22629:

- a cardboard core (cellulose)

- a folding cardboard box (cellulose)

3. Manufacture

Mollelast[®] haft bandages are produced according to specification under hygienic conditions and packed as described in its relevant packaging specification.

4. Description

Mollelast[®] haft is a ribbon with firm edges produced in a “knitting cum weaving technique”. It is a lengthwise and widthwise elastic conforming bandage. This bandage is cohesive, breathable and skin friendly.

Mollelast[®] haft bandages are available in various sizes.

5. Properties

Mollelast[®] haft are conforming bandages with an extensibility of approx. 80 %. The cohesive finish allows bandage tours to cling to each other, making bandage clips for securing bandage ends obsolete.

Mollelast[®] haft bandages are non-sterile and can be sterilized by steam in compliance with DIN EN ISO 17665-1 if necessary.

Please note:

Steam sterilization will increase the products resistance to unwinding and reduce its extensibility.

6. Intended purpose (see valid instructions for use)

Mollelast[®] haft bandages are intended for the immobilization of body parts, for the retention of dressings and cannulas, and/or for compression purposes.

7. Medical device classification

Mollelast[®] haft bandages are medical devices of Class I in terms of Rule 1. (Council Directive 93/42/EEC concerning medical devices, Annex IX)

- ▶ Exception - REF 22629 is not a medical device, it is not used on humans (for use on animals).

8. Biological evaluation and biocompatibility (DIN EN ISO 10993)

The starting materials used in the manufacture of Mollelast[®] haft bandages are safe provided the product is used appropriately and for the purposes intended.

However, persons allergic to natural latex should avoid direct skin contact with Mollelast[®] haft bandages.

The purpose of this documentation and the statements made therein is to show that there is no risk involved in the use of the medical device Mollelast[®] haft bandage and that it is designed, manufactured and packaged in such a way that it will not compromise the clinical condition or the safety of patients, or the safety and health of users and other persons when used under the conditions and for the purposes intended.

9. Stability

Stored appropriately – in their original packaging and at room temperature and air humidity of not more than 60 % - Mollelast[®] haft bandages have a shelf life of 5 years.

10. Disposal

The user is advised to observe current legislation, norms and guidelines, regulating the disposal of medical refuse.

Packaging materials must also be disposed of in compliance with applicable national requirements.

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