1. General Product Description

DermaPlast® Stretch is an elastic gauze bandage.

DermaPlast® Stretch carries the CE mark according to EU directive 93/42/EEC for medical devices. The product is classified as a class l medical device. A conformity assessment has been performed for
DermaPlast® Stretch and it has been shown to be in compliance with all applicable requirements of the above mentioned directive. The safe use of DermaPlast® Stretch, therefore, is ensured if the product is used in line with the intended purpose.

2. Application / Indication

DermaPlast® Stretch is used for light fixations.

3. Presentation

Skin-colored:

Size (width x length dilated)	Bandage/Folded box	Folded boxes/Transport cardboard box
6 cm x 10 m	1	200
4 cm x 4 m	10	125
6 cm x 4 m	1 (in Cellux-foil) 10	240 49
8 cm x 4 m	10	42

Weiss:

Size (width x length dilated)	Bandage/Folded box	Folded boxes/Transport cardboard box
4 cm x 10 m	1	280
6 cm x 10 m	1	200
8 cm x 10 m	1	160
2.5 cm x 4 m	20	1680
4 cm x 4 m	20	560
6 cm x 4 m	20	400
8 cm x 4 m	20	280
10 cm x 4 m	20	240

4. Product Characteristics

Material composition: 40% cot
60% polyamide

5. Product Requirements

Color: skin-colored or white
Extensibility: 100%

6. Labelling

LOT-Number, 7-digit

7. Packaging

DermaPlast® Stretch are packaged in folded boxes. The transport cardboard carton according to DIN is sealed with adhesive tapes: Packaging onto europallet.

Date: 27.06.2011

Marketing

CHANGE CONTROL REPORT

Keywords :	Technical Data Sheet		
Release of	**Substitutes release of**	**Change**	**No. of Pages**
13.07.06		new	3
27.08.08	13.07.06	Kapitel 6: Erklärung der LOT-Nummer gestrichen	3
27.06.2011	27.08.08	hautfarbig, Grösse 4cm x 10m, 8cm x 10m streichen	3

Date/Signature Written by:	Date/Signature Approved by:
E. Knill, R&D	A. Hartmann, Marketing