SMTL Dressings Datacard

Product Name:	**TELFA**
Classification Name:	**Dressing Perforated Film Absorbent BP (Type 2)**
Manufacturer:	**The Kendall Company Ltd**

Description

Telfa consists of a thin layer of absorbent cotton fibres, enclosed in a sleeve of poly(ethylene terephthalate) that is perforated in a regular pattern and sealed along two edges. The plastic film is present to prevent the dressing adhering to the surface of the wound, and is perforated to allow the passage of exudate from the wound into the body of the pad.

Indications

Telfa may be used on its own to dress dry sutured wounds, superficial cuts and abrasions[*** linked me-ab-00 ***], and other lightly exuding wounds. It may also be used as the primary wound contact layer for more heavily exuding wounds, if backed by a second absorbent dressing.

Contra-Indications

In common with other perforated plastic film dressings, Telfa should be used with caution in the treatment of leg ulcers that produce copious quantities of very viscous exudate. Under these circumstances, the exudate may become trapped under the dressing, leading to maceration and inflammation of the surrounding skin.

Method of use

Telfa has a low absorbent capacity and is therefore intended for use on lightly exuding wounds. Under these circumstances, a single dressing may simply be secured in position with surgical tape. On more heavily exuding wounds, Telfa may be used as a wound contact layer, beneath a secondary absorbent pad held in position with tape or a bandage.

Frequency of Change

The frequency with which the dressing should be changed depends entirely upon the nature and condition of the wound.

Presentation

Telfa is available in two forms: individually wrapped in paper peel pouches, sterilised by ethylene oxide; or as a dressing with the perforated film on one side only, supplied in bulk, non-sterile.

Sizes

Individually wrapped

5 cm × 7.5 cm

15 cm × 7.5 cm

10 cm × 7.5 cm

20 cm × 7.5 cm

Bulk

20 cm × 7.5 cm

25 cm × 20 cm

Revision Details

Revision Author	Dr S. Thomas
Revision No	1.3
Revision date	1997/12/16

Please inform SMTL of any inaccuracies or changes required in this datacard.

Disclaimer

This datacard has been prepared as an aid to contracting for purchasers in the NHS in Wales, and should not be used as a substitute for literature, instructions or training from the manufacturers of the devices.