Instructions for Use Soft Silicone Polyurethane Foam Wound Dressing

SilFoam V

1. PRODUCT DESCRIPTION

SilFoam V is a sterile, absorbent, self adherent soft silicone wound dressing. It comprises of a soft silicone skin and wound contact layer, a polyurethane foam layer with high absorption capacity and a vapour permeable, water and bacteria resistant polyurethane film outer layer. It is available in a border and non border version.

In the presence of exudate, SilFoam V helps maintain a moist wound environment conducive to natural healing conditions.

Product Composition:

Polyurethane, copolyamide, polyacrylate, silicone, polyethylene

2. INTENDED PURPOSE

Long term, non-invasive wound dressings intended principally for the management of light to moderately exuding, partial to full thickness wounds which have breached the dermis on injured skin and can only heal by secondary intent.

2.1 INTENDED POPULATION

Individuals of all ages who are at risk of developing

- pressure ulcers
- venous and arterial leg ulcers
- diabetic foot ulcers
- first and second degree burns

2.2 INTENDED USER

Intended for use by health professionals and caregivers and may be used in a hospital, community and home setting.

3. INDICATIONS

SilFoam V is indicated for the management of light to moderately exuding, partial to full thickness wounds, such as

- pressure ulcers
- venous and arterial leg ulcers
- diabetic foot ulcers
- first and second degree burns

SilFoam V may also be used as an aid for the prevention of skin breakdown.

4. CONTRAINDICATION/SAFETY INFORMATION

SilFoam V is contraindicated for

- ulcers resulting from infections, such as tuberculosis, syphilis, deep fungal infections;
- bites or third degree burns.

In case of infection with inflammatory signs (temperature, oedema, redness, pain) contact proper medical authority. Resume use of SilFoam V when normal healing conditions are present again.

Skin reactions may occur in very rare cases.

WARNINGS

Do not use if pouch is damaged or opened.

Do not re-use. Re-use of single-use devices creates a potential risk to the patient or user. It may lead to contamination and/ or impairment of functional capability. Contamination and/ or limited functionality of the device may lead to injury, illness or death of the patient.

6. INFORMATION FOR USE

SilFoam V is very simple to apply, requiring no special skills or equipment. The interval between dressing changes will depend entirely upon the state of the wound. On moderately exuding wounds, daily changes may be required at the beginning of treatment but this may be reduced to every 2 to 3 days for low exuding or epithelialising wounds.

6.1 Skin Preparation

- a) Cleanse the wound with sterile saline or Ringer solution and sterile swabs.
- b) Dry the skin surrounding the wound.

6.2 Dressing Application

- a) Select the appropriate SilFoam V size that will completely cover the wound surface, ensuring a 2 to 3 cm margin beyond the edges of the wound. If necessary, several dressings can be overlapped to cover very large wound areas.
- b) Grasp the tabs with both hands. Position the dressing over the wound site with the tab sides facing downward. Slowly peel away the tab from one side of the dressing.
- c) Apply with a rolling motion to the wound site.
- d) Remove the second tab. To secure, gently apply pressure to the dressing as it attaches to the wound.
- e) When dressing a sacral ulcer, slightly flex dressing and place into the gluteal fold.
 - Smooth outward to ensure adhesion. Examine the dressing on a daily basis for leakage or other problems. If no problems arise, the dressing may be left in place for
 - up to 7 days before another dressing is required.

f) In case of venous leg ulcers, compression therapy may be used in conjunction with Silfoam V treatment, when so directed by a physician.

6.3 Dressing Changes

SilFoam V should be changed when the dressing is saturated with exudate (2 to 3 days on average). Dressing may be left in place up to 7 days when there is little exudate or changed every 24 hours when the amount of exudate is significant. Where leakage occurs the dressing should be changed immediately.

a) Gently remove SilFoam V.

b) Follow procedure 5.1 a) to 5.2 c) to apply a new dressing.

7. CLINICAL BENEFIT / SPECIAL NOTES

 Continue the causal treatment such as compression therapy for venous leg ulcers or pressure relief for decubital ulcers.

The product may not be used in case of a known intolerance or an allergy to one

or several of its components.

 Because of its good adhesive properties the product should be removed with caution if the skin is fragile (e.g. parchment skin).

Do not use the product simultaneously with oxidizing solutions, e.g. hydrogen

peroxide or hypochlorite solution.

 The wound may initially appear to increase in size in the early stages of SilFoam V treatment. This is normal and occurs as any wound debris is removed from the edges of the wound. This clears the way for healing.

The frequency with which the dressing is changed should be adapted to reflect

the wound's healing progress, depending on the amount of exudate.

 Particularly in cases of an advanced degree of epithelialization over the wound and drying exudate, it may prove prudent to change to other modern wound treatment products.

• In the management of light to moderately exuding wounds, SilFoam V can only make the overlying environment more conducive to healing. There are cases where healing is impaired as a result of underlying conditions; in these instances, SilFoam V alone may make little or no progress, and suitable treatment of the underlying conditions will be necessary as well. Therefore, if after 4-6 weeks of SilFoam V treatment, there has been no improvement then, in line with accepted wound management practice, the original diagnosis and overall therapy should be reassessed.

SilFoam V should be left in place as long as possible in order to prevent trauma
to the fragile newly formed tissue and to reduce cross contamination through
frequent dressing changes. Thick necroses should be removed before applying

SilFoam V.

8. STORAGE

Store dressings away from direct sunlight at ambient temperature and humidity.

9. DISPOSAL

The foam dressings should be disposed based on the hospital or healthcare professional advice.

10. SYMBOLS OON THE LABELLING

WARNINGS		
	Do not use if the pouch is damaged or opened.	
STERILISATION		
STERILE EO (2)		
STORAGE		
淡	K	

11. PRESENTATION

SilFoam V is available in a bordered and non-bordered version in a range of sizes.

Size	
5 x 5 cm	
7.5 x 7.5cm	
10 x 10 cm	
15 x 15 cm	
15 x 20 cm	

12. COMPLAINTS

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

For complaints, questions or comments, contact XXXX Customer Support at XXXXXX.



Finesse Medical Ltd., IDA Business Park, Ballinalee Road, Longford Ireland C E 0120

Made in Ireland

Date of Issue: 12, June 2019

Version: Rev4