TECNODRAPE
SURGICAL INCISION ANTIMICROBIAL DRAPE

INSTRUCTIONS FOR USE

Surgical Incise Drape With Iodopovidone

Intended Use

TecnoDrape is designed to prevent the transmission of pathogens from the non-sterile skin towards the sterile surgical field in order to protect the patient from the risk of surgical site infections.

Warnings

Sterile and single-use. Check the integrity of the package before use. Discard any pack that shows signs of damage, since this could compromise the sterility of the product. This product must be used only by qualified staff. Do not re-use.

1. Disinfect the surgical area according to the hospitals Operating Theatre skin prep protocol. Wait for the skin prep to dry completely before applying the product. Do not swab the area to hasten the drying time. TecnoDrape is compatible for use with all skin preps.

2. Open the sterile pouch using aseptic technique and observe the correct method to "PEEL OPEN". Remove the protective paper wrap from the drape. Two operators are preferably required for proper drape placement.

3. Hold the drape (FIG.1) by placing operators hands on the appropriate sides of the drape (The Clear edges)

4. Remove the silicone paper (FIG.2) by firmly holding the drape and maintain a 90-degree angle with the film. Avoid applying on excessive force and do not twist.

5. Apply the film on the incision area by keeping it stretched at both ends (FIG.3). Take care to avoid air bubbles and folds during application.

6. To remove the drape, grab it gently at one end and pull it gently alongside the skin surface. Avoid stretching. It is normal to see Iodine adhesive remaining on the patients skin as this offers post operative skin antisepesis. Simply remove with sterile saline if required.


FIG 1.

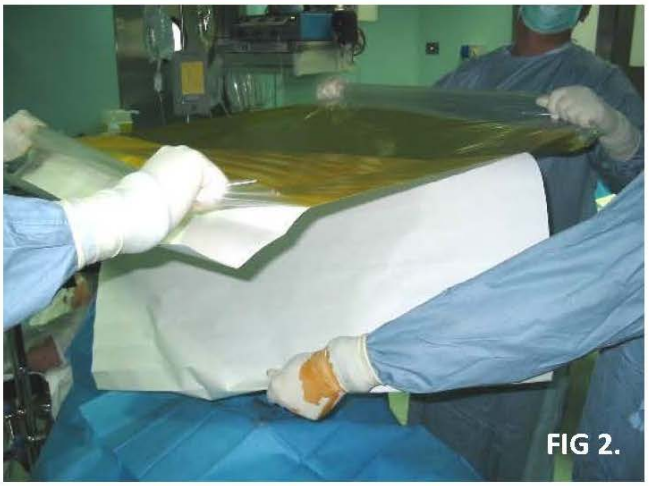
FIG 2.

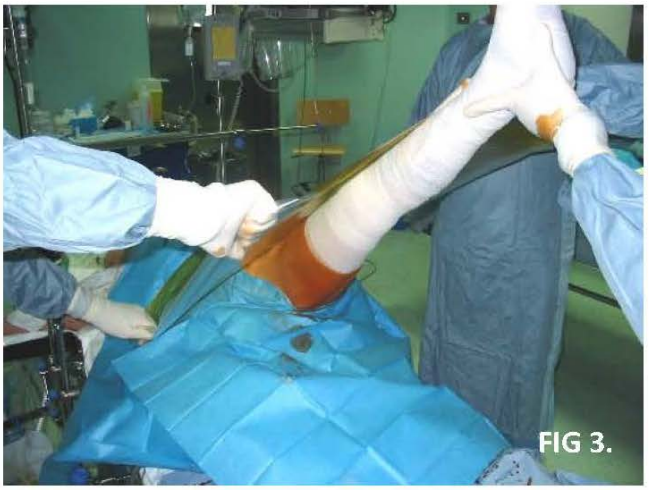
FIG 3.

Contra-indications:
Do not use on patients with known hypersensitivity to the component materials and in particular on patients with a known sensitivity to iodine or iodophor complexes.
Storage conditions: Keep away from sunlight and heating sources.
Special warnings: Do not defibrillate through the drape.

LOT			CE 0373
Batch code	Manufacturing date	Use by date	Class III medical device 93/42/EEC Directive
STERILE EO			
Sterilized by ethylene oxide	Keep dry	Keep away from heat/sunlight	Single use only
PHT			
Not contains PHT	Not to be used in case package is damaged	Manufacturer	

STERILE EO

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