Glubran®^2
APPLICATION DEVICES

Sterile 1 ml Luer Lock Syringe
Ref. G-LLS

Dispensing Tip
Ref. G-DT

Drop control device
Ref. G2-DCD-210-8T

Catheter for laparoscopy
Ref. G2-LPC

Ref. G2-LPC-RIG
**DESIGNATION**
Dispensing Tip

**DESCRIPTION**
Device consisting of polypropylene dispenser with a special connector to allow connection with the 2 different types of GLUBRAN glue vials.

**PROPERTIES**
The device allows topical application of GLUBRAN glues both during surgical procedures and during treatment of traumatic and atraumatic cutaneous wounds.

**METHOD OF USE**
The device can be used exclusively with GLUBRAN glues. Twist the cap to open the mono-dose vial and screw the dispensing tip onto the vial neck. Squeeze the vial to allow glue to flow through the dispensing tip and to be applied on the area to be treated. The device cannot be re-used. Discharge after the procedure.

**MATERIAL**
Polypropylene

**CODE**
G-DT

**STERILIZATION**
Gamma rays

**SHELF LIFE**
5 years

**PACKAGING**
Single sterile peel pouch

Any reuse of the disposable product involves risk of infection to the patient for a potential contamination of the product itself and the harm due to a possible impairment in its effectiveness.

**STORAGE**
Standard – Keep product in its original packaging in a clean, cool, dry place at temperatures below 30°C.

**COMPLIANCE DIRECTIVE**
MMD/93/42/EEC and subsequent updating

**QUALITY CONTROL**
During production and on the biocompatibility of the materials

**CLASS**
IIa

---

**DESIGNATION**
Catheter for laparoscopy

**DESCRIPTION**
Device consisting of:
- Hollow catheter with luer tip, for the exclusive application of GLUBRAN 2 surgical glue;
- (Flexible or rigid) Sheath

**INSTRUCTIONS**
Single use, exclusively for application of GLUBRAN 2 surgical glue during laparoscopic surgery. The device can be used only with 5 mm diameter trocar. Insert the entire device into the trocar, controlling compatibility and seal. Then connect the syringe of GLUBRAN 2 surgical glue to the catheter luer lock. Apply the glue drop by drop on the area to be treated, and to avoid the tip to be clogged up, avoid the catheter tip gets in contact with the area. When the procedure is completed, remove first the catheter and then the guide.

**MATERIALS**
Polyethylene, ethacrylate, polyester, PEBAX7233SA01, TERLUX 2802 TR, Stainless steel

**CODE**
G2-LPC
G2-LPC-RIG

**STERILIZATION**
Gamma rays

**SHELF LIFE**
5 years

**PACKAGING**
Double sterile peel pouch

Any reuse of the disposable product involves risk of infection to the patient for a potential contamination of the product itself and the harm due to a possible impairment in its effectiveness.

**STORAGE**
Standard – Keep product in its original packaging in a clean, cool, dry place at temperatures below 30°C.

**COMPLIANCE DIRECTIVE**
MMD/93/42/EEC and subsequent updating

**QUALITY CONTROL**
During production and on the biocompatibility of the materials

**CLASS**
IIa

---

**DESIGNATION**
Drop control device

**DESCRIPTION**
Device consisting of:
- catheter with luer connection complete with bevelled needle exclusively for application of GLUBRAN 2 surgical glue;
- 2.5 ml luer syringe complete with bevelled needle

**INSTRUCTIONS**
Single use, exclusively for the application of GLUBRAN 2 surgical glue during surgical procedures. In sterile conditions, remove the syringe with needle from the packaging and aspirate GLUBRAN 2 surgical glue from the vial as prescribed. Remove the needle used to aspirate the glue. Connect the drop control device to the syringe and press the piston several times until the glue begins to appear at the tip of the needle.

**MATERIALS**
Medical Grade PVC, polypropylene and polystyrene

**CODE**
G2-DCD-210-8T

**STERILIZATION**
Gamma rays

**SHELF LIFE**
5 years

**PACKAGING**
Single sterile peel pouch

Any reuse of the disposable product involves risk of infection to the patient for a potential contamination of the product itself and the harm due to a possible impairment in its effectiveness.

**STORAGE**
Standard – Keep product in its original packaging in a clean, cool, dry place at temperatures below 30°C.

**COMPLIANCE DIRECTIVE**
MMD/93/42/EEC and subsequent updating

**QUALITY CONTROL**
During production and on the biocompatibility of the materials

**CLASS**
IIa

Rev. 04 Ed. 07/05/2014