

Instruction for Use

Product: Veress Needle

The Following information should be read before using this device.

Description:

The Veress Needle is intended for percutaneous insertion into the peritoneal cavity for the purpose of insufflation with carbon dioxide to establish pneumoperitoneum prior to the placement of trocars during laparoscopic procedures.

Contraindications:

This device is not intended for use when endoscopic techniques are contraindicated.

Preparation:

1. Remove the product from the package tray.
2. To hold straight and insert into abdominal, the inner shaft will engage shield and protect to damage internal organ
3. To connect Insufflation generator with luer lock adaptor into needle or Trocar's gas port for insufflation gas into patient's abdominal.
4. Please follow instructions provided by Insufflation pump manufacturer.

Cautions/Warnings:

- This device was designed, tested and manufactured for single patient use only. Reuse or reprocessing of this device may lead to its failure and subsequent patient injury. Reprocessing and/or re-sterilization of this device may create the risk of contamination and patient infection. Do not reuse, reprocess or re-sterilize this device.
- Contents sterile unless enclosed package has been opened or damaged. Store at 15°C~25°C.
- Dispose used product following local state or federal guidelines.
- Federal (U.S.A) law restricts this device to sale, distribution, and use by, or on the order of a physician.



Caution



Sterilized using Ethylene Oxide



Consult instructions for use



Do not use if package is damaged



Batch code



Do not reuse



Keep away from sunlight



Date of manufacture



Keep dry



Catalogue number



Manufacturer



Authorised representative in the European community



Do not re-sterilize



Use by



Temperature limitation: 15°C~25°C



Medical device



Single sterile barrier system



Single sterile barrier system with protective packaging outside



Notified body: DNV Product Assurance AS, with CE2460

If further information is required, please contact:



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