

Laparoscopic Insufflation Tubing Set & Medi-stat Laprovent Vent Tubing Set

For use in laparoscopic procedures.

Medical device class; 2A

Intended use

For connecting an insufflation generator to a laparoscopic insufflation cannula or Verasse needle.

GMDN # ; 38559

Material

Styrene butadiene, PVC, LDPE, ABS, Polyester / PTFE filter media, activate carbon (Medi-Stat Laprovent filter)

Materials are latex free but may contain phthalates.

Instructions for use

Insufflation Tubing Set;

1. Remove unit from its packaging. Ensure there is no obvious damage to the tubing or the filter unit and that the tubing is securely attached to the filter and end connectors.
2. Ensure the generator and cannula / Veress needle connectors are the correct types.

Product Code	Machine manufacturer	Machine end	Patient end (trocar / Veress needle)
MED884170.101	Wolf	15mm Male	Male Swivel Luer lock
MED884710.900	Olympus	Male Swivel Luer lock	Male Swivel Luer lock
MED884170.901	Other	Female Fixed Luer lock	Male Swivel Luer lock
8870-400	N/A	N/A	Male Swivel Luer lock

3. Connect the tubing set short tube to the insufflation generator.
4. Ensure the tubing is not kinked or occluded in any way.
5. Check flow of CO² gas through tubing set.
6. Connect the long tube end connector to the cannula / Veress needle.
7. Follow the instructions provided with the insufflation generator.

Medi-stat Lapro-vent Carbon Filter Laparoscopic Vent Tubing set;

8. Laprovent does not connect to the insufflator it has filter and venting tap.
9. To vent quickly fully turn the tap to open in anti-clockwise direction.
10. Tap can be turned gradually for lower amounts of gas to be vented until correct level attained.

Storage

- ⇒ Store between 10 – 35°C and RH 20 – 80%.
- ⇒ Store in clean and dry place.
- ⇒ Do not use if the package open or damaged
- ⇒ Do not store in direct sunlight

Maintenance

No maintenance or servicing is permitted. This device is for single use only and must not be reprocessed.

Sterilization & reprocessing

The devices are supplied non-sterile and are for single use only. Because of the risk of possible cross contamination these devices must not be reprocessed.

Disposal

Discard the device after use in accordance with local regulation for medical waste.
 The device may be incinerated in a facility capable of burning plastic waste.

Warnings and cautions

- Not suitable for use other than that described above.
- The device contains small amounts of phthalate that may possibly cause harm to neonates, prepubescent males, nursing mothers and pregnant women.

Disposal

- The device should be disposed of after use using the rules applying locally for the disposal of contaminated medical equipment.
- The device may be incinerated in a facility capable of burning PVC waste.
- The devices end of life action should follow local disposal procedure regulations for non-biodegradable plastics

Ordering information:

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Explanation of symbols:

	= Single Use. The definition of single use is for use on a single patient for a single procedure.		= Manufacturer
	= Do not reprocess / re-sterilise		= Temperature range
	= Use by date		= Consult instructions for use
	= Lot or batch number.		= Keep away from sunlight
	= Date of manufacture.		= Keep dry
	= Device catalogue number / product code / stock		= Contains latex
	= Caution		= Do not use of packaging is wet or damaged.
	= Device catalogue number / product code / stock		= Contains or Presence of Phthalates.
	=CE mark with notified body number		

This is an explanation of the symbols that may be found on Meditech Systems Ltd's products.

ADVERSE INCIDENT REPORTING

Adverse incidents may be reported to the Competent Authority in the country / region where the incident occurred.

MIFU-018 (GGV1)

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