



eSuction™

 ET2000, ET2001



INSTRUCTIONS FOR USE

Read this document in its entirety prior to use

Rx Only
SINGLE USE, DISPOSABLE

INDICATIONS

The eSuction™ is intended to be used in the endoscopic retrieval of food bolus, foreign bodies and excised tissue such as polyps.

Product Description	Product Number	Outer Diameter	Fits scopes
eSuction™	ET2000	19 mm	8.6 mm – 10.0 mm
eSuction™	ET2001	19 mm	10.8 mm – 13.2 mm

CONTRAINDICATIONS

None known within the scope of the indicated use.

WARNINGS AND CAUTIONS:



Caution:

Federal (USA) Law restricts this device to sale by or on the order of a physician.



Warning:

For single use only. Do not reuse, reprocess or sterilize. Reuse, reprocessing or resterilization may compromise the structural integrity of the device and/ or lead to device failure which, in turn, may result in patient injury, illness or death.



Warning:

Inspect the package containing the product. If any damage is noticed, do not use the instrument as sterility may be compromised.



Warning:

Do not use if any abnormalities are found or if the instrument is not operating properly.



Warning:

After use, this product may be a potential biohazard. Handle or dispose of it in accordance with accepted medical practice and applicable local, state and federal laws and regulations.



Caution:

Verify device is fully retracted into outer sheath before pulling device through scope channel.



Caution:

During retrieval, allow for proper spacing between the end of the eSuction™ cap and the object being retrieved before deploying the snare. Deploying the snare too closely to an object may result in the snare detaching from the anchor side of the eSuction™ cap.

DIRECTIONS FOR USE

1. Verify that the outer diameter of the endoscope is compatible with the inner diameter of the eSuction™ cap. (Refer to chart.)
2. Peel open the pouch and remove the device. Visually inspect the device for kinks, loose or broken parts.
3. Deploy and retract the snare to ensure the device is functioning properly.
4. Verify that the endoscope has a 2.8 mm accessory channel or larger.
5. Dry the tip of the endoscope with a 4x4 gauze or dry wash cloth. It is important that the eSuction™ cap and the tip of the endoscope are dry to ensure that the device stays attached. Do not use any lubricant; doing so could result in the device falling off the endoscope inside the patient.
6. Load the retrieval snare into the accessory channel of the endoscope.
7. Position the distal tip of the snare into the catheter channel on the eSuction™ cap so that the snare distal tip is flush with the cap's distal end.
8. Place the eSuction™ cap onto the endoscope and align the catheter channel with the endoscope's accessory channel.
9. Gently pull the eSuction™ cap to ensure there is a proper fit on the endoscope.
10. Deploy the snare, fold over and attach the tip onto the anchor side of the eSuction™ cap.
11. Retract and deploy the snare to ensure attachment to anchor side of the eSuction™ cap.
12. The snare should be retracted before inserting into patient
13. Deploy snare loop around food bolus, foreign body or excised tissue by moving finger loops forward.
14. Retract snare loop with food bolus, foreign body or excised tissue by moving finger loops backward.
15. Endoscopic suction may be used as necessary to aid in the removal of food bolus impactions, foreign bodies and excised tissue.
16. After the procedure is completed, remove the endoscope from the patient.
17. Deploy and detach the snare loop from the anchor side of the eSuction™ cap and retract the snare.
18. Remove the eSuction™ cap from the endoscope and discard.

DISPOSAL

After use, this product may be a potential biohazard. Handle or dispose of it in accordance with accepted medical practice and applicable local, state and federal laws and regulations.

STORAGE
















Store sterile instruments at room temperature. Do not expose to solvents, radiation, UV light or any other any other substance or environment that could compromise the sterility of the unit.

WARRANTY

Endo-Therapeutics, Inc. warrants that the instrument will be free from defects in materials and workmanship for use in a single procedure.

This warranty does not apply to use beyond the expiration date. This device is not intended to be reprocessed. The user assumes responsibility for any consequences related to the use of reprocessed instruments. Endo-Therapeutics, Inc. obligations under this warranty are limited to the replacement of the instrument.

LABELING SYMBOLS

	Consult Instruction for Use
	Warning - Indicates a hazardous situation which, if not avoided, could result in death or serious injury.
	Caution - Indicates a hazardous situation which, if not avoided, could result in minor or moderate injury.
	Single use - Do not reuse
	Do not resterilize
	Keep dry
	Ethylene Oxide Sterilization
	Catalog Number / Product Code
	Lot Number
	Use by Date
	Date of manufacture
RxOnly	Federal (USA) Law restricts this device to sale by or on the order of a physician
	Manufacturer
	Quantity of devices in pouch
	Quantity of devices in box
	Do not use if package is opened or damaged


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