

Catchem'

POLYP TRAP



FEDERAL (USA) LAW RESTRICTS THIS DEVICE TO SALE BY OR ON THE ORDER OF A PHYSICIAN

EN - INSTRUCTIONS FOR USE



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Think Within.



REF: ET2211 Catchem® Polyp Trap, 4 Chamber, Removable

REF: ET2213 Catchem® Polyp Trap, 4 Chamber, Fixed

Intended Use

The single use polyp trap is designed for in-line suction retrieval of polyps during endoscopic procedures.

Contraindications

None known within the scope of the intended use.

Warnings

Do not use if package is opened or damaged.

Visually inspect before use. If any abnormality is detected that would prohibit proper working condition, do not use.

This device is designed for single use only. Attempts to reprocess, resterilize, and/or reuse may lead to device failure and/or transmission of disease.

Instructions

1. Connect suction tubing to center port on trap and outside tubing on lid to endoscope.
2. Rotate arrow on lid to position tubing over chamber 1.
3. Retrieve and collect specimen.
4. Rotate lid clockwise to position tubing over chamber 2 after first specimen is collected.
5. Retrieve and collect specimen.
6. Repeat step 4 and 5 by positioning tubing over chamber 3 & 4.
7. To detach, disconnect suction tubing from suction system and trap tubing from endoscope.
8. For ET2211 only: To remove the chambers, position the canister with the desired chamber facing you. Apply firm pressure to the upper portion of the canister, directly on the number corresponding to the desired chamber for removal. At the same time using your other hand, securely grip the desired chamber and gently pull it away from the base of the canister. Repeat these steps until all the chambers have been removed from the canister.
9. After all specimens are transferred to a transport vial, discard remaining portion of trap immediately.

Notes

Use of this device restricted to a trained healthcare professional.

In the event of a malfunction, discontinue use and save for return and contact your local representative.

Serious incidents that have occurred in relation to this medical device should be reported to the manufacturer and competent authority in the country where the incident occurred.

Disposal

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



Storage

Store in a dry location, away from temperature extremes.

Symbols

EN - Symbols

A symbol glossary can be found at endotherapeutics.com/symbol-glossary.

	EN - Quantity of devices in pouch
	EN - Quantity of devices in box

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