

## **CO<sub>2</sub> SOURCE TUBING ADAPTER**

Instructions for Use.
Read carefully prior to use.

Manufactured for Ruhof Corporation 393 Sagamore Avenue Mineola, NY 11501 1-800-537-8463 www.ruhof.com

Not Made with natural rubber latex.

Does not contain DEHP

<u>Part Number</u> <u>Description</u>

GAR093 CO<sub>2</sub> Source Tubing Adapter (Non-Sterile)

#### **INTENDED USE**

The Vacutore® CO<sub>2</sub> Source Tubing Adapter connects to the gas outlet port on the front of the Olympus® UCR CO<sub>2</sub> insufflator. The CO<sub>2</sub> Source Tubing Adapter provides a connection for the CO<sub>2</sub> Source Tubing and Extend Tubing with Luer Connector.

#### WARNINGS AND PRECAUTIONS

- ALWAYS prime the air/water channel prior to inserting the GI endoscope into the patient.
- NEVER attach or reattach the CO<sub>2</sub> Source Tubing Adapter to a GI endoscope that is/was inside a patient and has not been reprocessed.
- NEVER turn off the CO<sub>2</sub> supply while the CO<sub>2</sub> Source Tubing Adapter is attached to the GI endoscope.
- If any of the above prevention guidelines were not followed, or there is any reason to believe that the CO<sub>2</sub> Source Tubing Adapter was contaminated, immediately discard the entire tubing, accessories and water bottle. Replace with new tubing and an unopened sterile water bottle.
- ALWAYS inspect the GI endoscope's o-ring and gaskets (air/water button, GI endoscope connection, etc.) for damage. Damage components can result in less than optimal performance of the CO<sub>2</sub> Source Tubing Adapter.
- Endoscopic procedures should only be performed by persons having adequate training and familiarity with endoscopic techniques.
- Consult the medical literature relative to complications, hazards, and techniques prior to the performance of any endoscopic procedure.
- Use proper aseptic techniques to avoid contamination of the CO<sub>2</sub> Source Tubing Adapter during device setup, replacement of water bottles and between patient uses.

#### **DIRECTIONS FOR USE**

- Prior to clinical use, familiarize yourself with the device and read all the Instructions for Use.
- Inspect the package for shipping or handling damage. If damage is evident, DO NOT USE THESE DEVICES, SAVE THEM OR RETURN, AND CONTACT YOUR LOCAL PRODUCT SPECIALIST.
- Open the peel pack and remove the Vacutore® CO<sub>2</sub> Source Tubing Adapter.
- Place the CO<sub>2</sub> Source Tubing Adapter on the "gas outlet port" on the front of the Olympus® UCR CO<sub>2</sub> Insufflator.
- Turn the CO<sub>2</sub> Source Tubing Adapter clockwise until it is held securely in place.1
- Use the luer connector of the CO<sub>2</sub> Source Tubing to connect to the CO<sub>2</sub> Source Tubing Adapter. Turn luer until firmly connected.
- 7. For directions related to the CO<sub>2</sub> Source Tubing, please see the product specific instructions for use.

#### CLEANING, DISINFECTING OR STERILIZING

Always follow national and local guidelines and your hospital's infection control department to determine appropriate high-level disinfection or sterilization procedures.

### WARNING

An issued or revision date for these instructions is included for the user's information. In the event that two years have elapsed between this date and product use, the user should contact GA Health Company Limited to determine if additional information is available.

Report any serious incident that has occurred in relation to the device to the distributor or manufacturer or competent authority of the member state.

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 $\textbf{Vacutore}^{\textcircled{e}}$  is a registered trademark of GA Health Company Limited.

Olympus® is a registered trademark of Olympus corporation.

# EXPLANATION OF SYMBOLS USED ON LABELS AND INSTRUCTIONS FOR USE



Use-by date



Batch or lot code



Manufacturer



Date of Manufacture



**Medical Device** 



Catalogue number Product Code



Notified body TÜV SÜD



Product not made with natural rubber latex



Do not contain DEHP



**Consult Instructions for Use** 



Authorized representative in the European Community



Keep dry



Keep away from sunlight



Federal law (U.S.A.) restricts this device to sale, distribution and use by or on the order of a physician.