Overview

Product Rapid Refill™ Continuous Injection System – IFU 51999304-01 Rev. A

Rx Statement

CAUTION: Federal law (USA) restricts this device to sale by or on the order of a licensed practitioner.

This device should only be used by or under the supervision of physicians trained in Endoscopic procedures. A thorough understanding of the technical principles, clinical applications and risks associated with endoscopic procedures is necessary before using this device.

REUSE WARNING

For single use only. Do not reuse, reprocess or resterilize. 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. Reuse, reprocessing or resterilization may also create a risk of contamination of the device and/or cause patient infection or cross-infection, including, but not limited to, the transmission of infectious disease(s) from one patient to another. Contamination of the device may lead to injury, illness or death of the patient.

Prior to use, please refer to all applicable “Instructions for Use” for more information on Intended Use/Indications for Use, Contraindications, Warnings, Precautions, Potential Adverse Events, and Operator’s Instructions.

INTENDED USE/INDICATIONS FOR USE

This device is indicated for use in fluid administration procedures.

CONTRAINDICATIONS
None known.

PRECAUTIONS
- Spike cover must be removed before being inserted.
- Verify spike vent is in open position to allow fluid flow.

POTENTIAL ADVERSE EVENTS
None known.

CAUTION: The law restricts these devices to sale by or on the order of a physician. Indications, contraindications, warnings, and instructions for use can be found in the product labelling supplied with each device or at www.IFU-BSCI.com. Products shown for INFORMATION purposes only and may not be approved or for sale in certain countries. This material not intended for use in France.