

Interject[®] Sclerotherapy Needles

Performance you can count on for injection therapy applications.

INTERJECT SINGLE-USE SCLEROTHERAPY NEEDLES

Order Number	Sheath Design	Needle Gauge	Max. Needle Extension Length (mm)	Sheath O.D. (mm)	Sheath Length (cm)
M00518151	Contrast	.23	4	1.8	200
M00518161	Contrast	.25	4	1.8	200
M00518251	Contrast	.23	6	1.8	200
M00518261	Contrast	.25	6	1.8	200
M00518351	Contrast	.23	4	2.3	240
M00518361	Contrast	.25	4	2.3	240

INTERJECT CLEAR SINGLE-USE SCLEROTHERAPY NEEDLES

Order Number	Sheath Design	Needle Gauge	Max. Needle Extension Length (mm)	Sheath O.D. (mm)	Sheath Length (cm)
M00518111	Clear	.25	4	1.8	200
M00518301	Clear	.23	4	2.3	240
M00518311	Clear	.25	4	2.3	240

INSTRUCTIONS FOR USE

Refer to the operator's manual for complete instructions for use.

INDICATIONS

Indications for endoscopy to introduce a sclerosing agent or vasoconstrictor into selected sites to control actual or potential bleeding lesions in the digestive system; and the injection of saline to aid in Endoscopic Mucosal Resection (EMR), polypectomy procedures and to control non-variceal hemorrhage.

CONTRAINDICATIONS

Contraindications for this device are those applicable to injection therapy and include, but may not be limited to, those patients allergic to sclerosing or vasoconstriction agents and patients with lesions inappropriate for injection therapy with sclerosing or vasoconstriction agents.

WARNINGS

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.

POTENTIAL ADVERSE EFFECTS

Potential complications include, but may not be limited to bleeding, post-injection ulceration with delayed bleeding, perforation, aspiration pneumonia, pleural effusion, hepatic failure, septicemia, chest pain, esophageal ulcers, esophageal stricture and dysphasia.

Please be aware that potential adverse effects may arise even with the proper use of medical devices. Accordingly, this device should only be used by persons qualified in the procedures for which it is indicated.

CAUTIONS

Cautions can be found in the product labeling supplied with each device. CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.

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Ordering Information
1.800.225.3226

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