

# WALLSTENT<sup>®</sup> Colonic & Duodenal Endoprosthesis

With UNISTEP<sup>™</sup> Plus Delivery System

## WALLSTENT SINGLE-USE COLONIC AND DUODENAL ENDOPROSTHESIS WITH UNISTEP PLUS DELIVERY SYSTEM\*

Order Number	Expanded Stent Diameter (mm)	Expanded Stent Length (mm)	Working Length (cm)	Delivery System Length (cm)	Delivery System Diameter (fr)
M00565560.....	20	60	230	255	10
M00565570.....	20	90	230	255	10
M00565580.....	22	60	230	255	10
M00565590.....	22	90	230	255	10
M00565620.....	20	60	135	160	10
M00565630.....	20	90	135	160	10
M00565640.....	22	60	135	160	10
M00565650.....	22	90	135	160	10

\* Recommended Guidewire .035 Wallstent Super Stiff Guidewire

## WALLSTENT SINGLE-USE GI GUIDEWIRE

Order Number	Description	O.D. (in)	Length (cm)	Tip Style
H965180011.....	Wallstent Super Stiff Guidewire.....	.035.....	500.....	Tapered.....

### INSTRUCTIONS FOR USE

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

### INDICATIONS

Indicated for the palliative treatment of colonic, duodenal or gastric outlet obstruction or strictures caused by malignant neoplasms, and to relieve large bowel obstruction prior to colectomy in patients with malignant strictures.

### CONTRAINDICATIONS

Contraindications include:

- Enteral ischemia
- Suspected or impending perforation
- Intra-abdominal abscess/perforation

### WARNINGS

Our devices are designed, manufactured, tested, validated and labeled for single use only. These devices should not be reused, reprocessed or resterilized. 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.

Stents cannot be repositioned after the deployment threshold has been exceeded.

### POTENTIAL ADVERSE EFFECTS

Potential adverse effects may include the usual complications reported for conventional stents and endoscopic procedures such as infection, stent misplacement, stent migration, intestinal perforation and stent obstruction secondary to tumor ingrowth through the stent, tumor overgrowth at the stent ends, or occlusion.

Post stent placement complications may include: bleeding, perforation, pain, stent migration, tumor ingrowth through stent, tumor overgrowth around ends of stent, foreign body sensation, bowel impaction, reflux, ulceration, fever, septicemia, and death (other than that due to normal disease progression).

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.

### TRADEMARKS

WALLSTENT is a registered trademark and UNISTEP is a trademark of Boston Scientific Corporation or its affiliates.

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Ordering Information  
1.800.225.3226

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