

Gold Probe™ Electrohemostasis Catheters

Bipolar Hemostasis Catheters

GOLD PROBE SINGLE-USE ELECTROHEMOSTASIS CATHETER

Order Number	Description	Working Length(cm)	Type Generator / Connector
M005600707Fr Gold Probe	300.....	Standard Plug*.....
M005600717Fr Gold Probe (Box 5).....	300.....	Standard Plug*.....
M0056010010Fr Gold Probe	300.....	Standard Plug*.....
M0056010110Fr Gold Probe (Box 5)	300.....	Standard Plug*.....
M005602207Fr Gold Probe Enteral	350.....	Standard Plug*.....

* Compatible with Endostat™ Bipolar-Monopolar Electrosurgical Generators. For Banana Plug Connector, use Electrosurgical Bipolar Cable Adaptor.

INSTRUCTIONS FOR USE

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

INDICATIONS

The Gold Probe, Gold Probe Direct and Gold Probe 210cm, 300cm, and 350cm Electrohemostasis Catheters are indicated for use in transendoscopic electrocautery of visible bleeding and non-bleeding sites in the gastrointestinal tract including the esophagus, stomach, duodenum, and colon. The indications include: peptic ulcers, angiomata, Mallory-Weiss tears, watermelon stomach, arteriovenous malformations (AVMs), Barrett's esophagus, dieulafoy lesions, angiodysplasia, bleeding polyp stalks, and esophageal tumors.

CONTRAINDICATIONS

Contraindications known for these devices are those specific to bipolar electrocautery treatment. These contraindications include, but are not limited to non-focal bleeding sites, esophageal/gastric varices, diffuse lesions, and use of these devices with monopolar electrical generators.

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 a patient.

After use, dispose of product and packaging in accordance with hospital, administrative and/or government policy.

POTENTIAL ADVERSE EFFECTS

Potential complications may include, but are not limited to: perforation, bleeding, aspiration pneumonia, and septicemia/infection.

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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