January 10, 2004

ENDOLOGIX TO HOST CONFERENCE CALL TO DISCUSS DEVELOPMENT AGREEMENT FOR BIOFOAM

IRVINE, Calif. (January 10, 2004) - Endologix, Inc. (Nasdaq: ELGX) will host a conference call tomorrow, Tuesday, January 11, 2005 at 12:00 p.m. Eastern Time (9:00 a.m. Pacific Time) to discuss the details of its agreement to develop and market CryoLife, Inc.'s (NYSE: CRY) BioFoamT as a self-expanding sealant for endovascular aortic aneurysm grafts.

Individuals interested in participating in the conference call may do so by dialing (888) 463-4487 for domestic callers, or (706) 634-5615 for international callers. Those interested in listening to the conference call live via the Internet may do so by visiting the company's Web site at www.endologix.com.

A replay will be available on the Endologix Web site for 14 days. A telephone replay will be available for 48 hours following the conclusion of the call by dialing (800) 642-1687 for domestic callers, or (706) 645-9291 for international callers, and entering reservation code 3235425.

About Endologix
Endologix, Inc. develops and manufactures minimally invasive treatments for vascular diseases. Endologix's Powerlink System is an endoluminal stent graft for treating abdominal aortic aneurysms (AAA). AAA is a weakening of the wall of the aorta, the largest artery in the body, resulting in a balloon-like enlargement. Once AAA develops, it continues to enlarge and, if left untreated, becomes increasingly susceptible to rupture. The overall patient mortality rate for ruptured AAA is approximately 75%, making it the thirteenth leading cause of death in the United States. In October 2004, Endologix received approval to market the Powerlink in the U.S. Additional information can be found on Endologix’s Web site at www.endologix.com.

The above referenced conference call may contain forward-looking statements, the accuracy of which are necessarily subject to risks and uncertainties, including risks related to the development, clinical success and regulatory approval of a new medical device product, and the risks related to intellectual property rights surrounding new technology, all of which are difficult or impossible to predict accurately and many of which are beyond the control of Endologix, all are more fully described in the risk factors and other matters set forth in the Company's Annual Report on Form 10-K for the year ended December 31, 2003, and the Company's other filings with the SEC.