CryoLife to Present at 6th Annual Canaccord Genuity Cardiovascular, Aesthetics & Metabolic Disorders Conference

November 29, 2011

ATLANTA, Nov. 29, 2011 /PRNewswire/ -- CryoLife, Inc. (NYSE: CRY), a leading tissue processing and medical device company focused on cardiac and vascular surgery, announced today that it will participate in the upcoming 6th Annual Canaccord Genuity Cardiovascular, Aesthetics & Metabolic Disorders Conference on Tuesday, December 6, 2011 at the St. Regis Hotel in San Francisco.

D. Ashley Lee, executive vice president, chief operating office, and chief financial officer of CryoLife, Inc., is scheduled to present at 1:40 p.m. PT.

CryoLife's live presentation may be accessed through its website, www.cryolife.com, on the Investor Relations page. An archived copy of the presentation will be available for 90 days on the same website.

About CryoLife

Founded in 1984, CryoLife, Inc. is a leader in the processing and distribution of implantable living human tissues for use in cardiac and vascular surgeries throughout the U.S. and Canada. CryoLife's CryoValve® SG pulmonary heart valve, processed using CryoLife's proprietary SynerGraft® technology, has FDA 510(k) clearance for the replacement of diseased, damaged, malformed, or malfunctioning native or prosthetic pulmonary valves. CryoLife's CryoPatch® SG pulmonary cardiac patch has FDA 510(k) clearance for the repair or reconstruction of the right ventricular outflow tract (RVOT), which is a surgery commonly performed in children with congenital heart defects, such as Tetralogy of Fallot, Truncus Arteriosus, and Pulmonary Atresia. CryoPatch SG is distributed in three anatomic configurations: pulmonary hemi-artery, pulmonary trunk, and pulmonary branch. CryoLife's BioGlue® Surgical Adhesive is FDA approved as an adjunct to sutures and staples for use in adult patients in open surgical repair of large vessels. BioGlue is also CE marked in the European Community and approved in Canada and Australia for use in soft tissue repair and was recently approved in Japan for use in the repair of aortic dissections. CryoLife, through its subsidiary Cardiogenesis Corporation, specializes in the treatment of cardiovascular disease and the sale of devices that treat severe angina. Its market leading FDA-approved Holmium: YAG laser system and single use fiber-optic delivery systems are used to perform a surgical procedure known as Transmyocardial Revascularization (TMR). CryoLife distributes PerClot®, an absorbable powder hemostat, in the European Community. CryoLife's BioFoam™ Surgical Matrix is CE marked in the European Community for use as an adjunct in the sealing of abdominal parenchymal tissues (liver and spleen) when cessation of bleeding by ligature or other conventional methods is ineffective or impractical.

For additional information about the company, visit CryoLife's website: http://www.cryolife.com.

Contacts:

CryoLife	**The Ruth Group**
D. Ashley Lee	Nick Laudico / Zack Kubow
Executive Vice President, Chief Financial Officer and Chief Operating Officer	646-536-7030 / 7020
Phone: 770-419-3355	nlaudico@theruthgroup.com
	zkubow@theruthgroup.com

SOURCE CryoLife, Inc.