CryoLife Announces Release Date and Teleconference Call Details for 2014 Third Quarter Financial Results

October 14, 2014

ATLANTA, Oct. 14, 2014 /PRNewswire/ -- CryoLife, Inc. (NYSE: CRY), a leading medical device and tissue processing company focused on cardiac and vascular surgery, announced today that 2014 third quarter financial results will be released on Tuesday, October 28, 2014. On that day, the Company will hold a teleconference call and live webcast at 10:00 a.m. Eastern Time to discuss the results, followed by a question and answer session hosted by Pat Mackin, president and chief executive officer of CryoLife, Inc.

To listen to the live teleconference, please dial 201-689-8261 a few minutes prior to 10:00 a.m. A replay of the teleconference will be available October 28 through November 4 and can be accessed by calling (toll free) 877-660-6853 or 201-612-7415. The conference number for the replay is 13592923.

The live webcast and replay can be accessed by going to the Investor Relations section of the CryoLife website at www.cryolife.com and selecting the heading Webcasts & Presentations. In addition, a copy of the earnings press release, which will contain financial and statistical information for the completed quarter, can be accessed on the Investor Relations section of the CryoLife website.

About CryoLife, Inc.

CryoLife, Inc. is a leader in medical device manufacturing and distribution and in the processing and distribution of implantable living human tissues for use in cardiac and vascular surgeries. It operates throughout the U.S. and internationally. CryoLife manufactures and distributes BioGlue® Surgical Adhesive, an FDA-approved adjunct to sutures and staples for use in adult patients in open surgical repair of large vessels. BioGlue is also CE marked in Europe for use in soft tissue repair and has received additional marketing approvals in several other countries throughout the world. CryoLife's BioFoam® Surgical Matrix is CE marked in Europe for use as an adjunct to hemostasis in cardiovascular surgery and on abdominal parenchymal tissues (liver and spleen) when control of bleeding by ligature or conventional methods is ineffective or impractical. CryoLife distributes PerClot®, a powdered hemostat, in Europe and other select international countries. CryoLife has received FDA 510(k) clearance for a topical version of PerClot, which is being marketed in the U.S. for use in topical applications, and is conducting a pivotal clinical trial in the U.S. for potential FDA approval of the surgical version of PerClot. CryoLife, through its subsidiary Cardiogenesis Corporation, specializes in the treatment of coronary artery disease for severe angina using a laser console system and single-use, fiber-optic handpieces to perform a surgical procedure known as Transmyocardial Revascularization (TMR). CryoLife and its subsidiary Hemosphere, Inc. market the HeRO® Graft, which is a solution for end-stage renal disease in certain hemodialysis patients. CryoLife distributes ProCol®, a natural biological graft derived from a bovine mesenteric vein that provides vascular access for end-stage renal disease hemodialysis patients, which is approved for sale in the U.S. 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.

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

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