New Outcomes Data Using Delcath's Melphalan/HDS To Treat Unresectable Metastatic Ocular Melanoma To The Liver Accepted For Oral Presentation At Regional Cancer Therapies Symposium

NEW YORK, Jan. 18, 2017 — Delcath Systems, Inc. (NASDAQ: DCTH), an interventional oncology Company focused on the treatment of primary and metastatic liver cancers, today announced that new data from a retrospective study of Melphalan/HDS, entitled "*Percutaneous Hepatic Perfusion For Unresectable Metastatic Ocular Melanoma To The Liver: A Multi-Institutional Report Of Outcomes,"* has been accepted for oral presentation at the Regional Cancer Therapies 12th International Symposium, taking place February 18 – 20, 2017 at the Snowbird Ski and Summer Resort in Snowbird, UT.

The retrospective study was conducted by teams from Moffitt Cancer Center in Tampa, FL and the University of Southampton in the United Kingdom. The study explores patient treatment outcomes with Percutaneous Hepatic Perfusion (PHP® Therapy) with Melphalan/HDS in patients with primary metastatic ocular melanoma (OM) with liver metastasis treated between 2008 and 2016. The study was led by Dr. Alexandra Gangi of the Moffitt Cancer Center.

"We are very pleased that Dr. Gangi and her team's research has been accepted for oral presentation at this prestigious event and believe that their research results will further inform on the use of Melphalan/HDS as a potentially viable additional option for treatment of hepatic metastases in patients with metastatic ocular melanoma," said Jennifer K. Simpson, Ph.D., MSN, CRNP, President and Chief Executive Officer of Delcath. "We look forward to announcing the details of these new data after they are presented."

About Delcath Systems

Delcath Systems, Inc. is an interventional oncology Company focused on the treatment of primary and metastatic liver cancers. Our investigational product-Melphalan Hydrochloride for Injection for use with the Delcath Hepatic Delivery System (Melphalan/HDS) -is designed to administer high-dose chemotherapy to the liver while controlling systemic exposure and associated side effects. We have commenced a global Phase 3 FOCUS clinical trial for Patients with Hepatic Dominant Ocular Melanoma (OM) and a global Phase 2 clinical trial in Europe and the U.S. to investigate the Melphalan/HDS system for the treatment of primary liver cancer (HCC) and intrahepatic cholangiocarcinoma (ICC). Melphalan/HDS has not been approved by the U.S. Food & Drug Administration (FDA) for sale in the U.S. In Europe, our system has been commercially available since 2012 under the trade name Delcath Hepatic CHEMOSAT® Delivery System for Melphalan (CHEMOSAT), where it has been used at major medical centers to treat a wide range of cancers of the liver.

Forward Looking Statements:

Private Securities Litigation Reform Act of 1995 provides a safe harbor for forward-looking statements made by the Company or on its behalf. This news release contains forward-looking statements, which are subject to certain risks and uncertainties that can cause actual results to differ materially from those described. Factors that may cause such differences include, but are not limited to, uncertainties relating to: our ability to repay and comply with the obligations under our senior secured convertible notes, the timing and results of the Company's clinical trials including without limitation the OM, HCC ,and ICC clinical trial programs, timely enrollment and treatment of patients in the global Phase 3 FOCUS Clinical Trial for Patients with Hepatic Dominant Ocular Melanoma and the global Phase 2 HCC and ICC clinical trials, IRB or ethics committee clearance of the Phase 2 HCC/ICC and/or Phase 3 OM protocols from participating sites and the timing of site activation and subject enrollment in each trial, the impact, if any, of publication of the Phase 3 trial manuscript to support the Company's efforts, the impact of the presentations at major medical conferences and future clinical results consistent with the data presented, the impact, if any of ZE reimbursement on potential CHEMOSAT product use and sales in Germany, clinical adoption, use and resulting sales, if any, for the CHEMOSAT system to deliver and filter melphalan in Europe, the Company's ability to successfully commercialize the CHEMOSAT/Melphalan HDS system and the potential of the CHEMOSAT/Melphalan HDS system as a treatment for patients with primary and metastatic disease in the liver, our ability to obtain reimbursement for the CHEMOSAT system in various markets, the Company's ability to satisfy the remaining requirements of the FDA's Complete Response Letter and provide the same in a timely manner, approval of the current or future Melphalan HDS/CHEMOSAT system for delivery and filtration of melphalan for various indications in the U.S. and/or in foreign markets, actions by the FDA or other foreign regulatory agencies, the Company's ability to successfully enter into strategic partnership and distribution arrangements in foreign markets and the timing and revenue, if any, of the same, uncertainties relating to the timing and results of research and development projects, our ability to maintain NASDAQ listing, and uncertainties regarding the Company's ability to obtain financial and other resources for any research, development, clinical trials and commercialization activities. These factors, and others, are discussed from time to time in our filings with the Securities and Exchange Commission. You should not place undue reliance on these forward-looking statements, which speak only as of the date they are made. We undertake no obligation to publicly update or revise these forward-looking statements to reflect events or circumstances after the date they are made.

Contact Information:

Investor Contact:
Anne Marie Fields
LHA
212-838-3777
afields@lhai.com

To view the original version on PR Newswire, visit:http://www.prnewswire.com/news-releases/new-outcomes-data-using-delcaths-melphalanhds-to-treat-unresectable-metastatic-ocular-melanoma-to-the-liver-accepted-for-oral-presentation-at-regional-cancer-therapies-symposium-300392530.html

SOURCE Delcath Systems, Inc.