



Delcath's PHP Therapy Featured in Video Learning Session at the European Conference of Interventional Oncology

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NEW YORK, May 14, 2018 (GLOBE NEWSWIRE) -- Delcath Systems, Inc. (OTCQB:DCTHD), an interventional oncology company focused on the treatment of primary and metastatic liver cancers, announces that the Company's PHP Therapy was featured in a Video Learning Session presented at the Annual Meeting of the European Conference of Interventional Oncology (ECIO). Dr. M.C. Burgmans of Leiden University Medical Center (LUMC) in the Netherlands presented the training in the main auditorium session dedicated to advances in liver cancer therapies. Dr. Burgmans presented an overview of the Percutaneous Hepatic Perfusion (PHP) procedure, discussed the therapy's developmental history, demonstrating how to perform the procedure, as well as outlining its potential in ocular melanoma liver metastases and intrahepatic cholangiocarcinoma, and highlighting ongoing clinical research.

"In his presentation, Dr. Burgmans stated his belief that PHP Therapy should be considered as first line therapy in ocular melanoma liver metastases, an opinion informed by both our commercial experience in Europe and our prior research into this tumor type," said Jennifer K. Simpson, Ph.D., MSN, CRNP, President and CEO of Delcath Systems. "LUMC is one of the our most experienced treatment centers, and with CHEMOSAT recently added to the Dutch National Treatment Guidelines for use in ocular melanoma, we believe Dr. Burgmans comments reflect growing confidence in this therapy's role in treating this disease. We continue to work tirelessly to advance our current clinical trial in this disease in order to further validate this role for our therapy and to deliver on its full potential to patients in need."

The ECIO was held in Vienna, Austria, April 22-25, 2018.

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 have initiated a Registration trial for intrahepatic cholangiocarcinoma (ICC) called The ALIGN Trial. 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: the timing and results of the Company's clinical trials including without limitation the OM and ICC clinical trial programs, timely enrollment and treatment of patients in the global Phase 3 OM clinical trial, IRB or ethics committee clearance of the Phase 3 OM and ICC Registration trial protocols from participating sites and the timing of site activation and subject enrollment in each trial, the impact of the presentations at major medical conferences and future clinical results consistent with the data presented, approval of Individual Funding Requests for reimbursement of the CHEMOSAT procedure, 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 including the key markets of Germany and the UK, the Company's ability to successfully commercialize the Melphalan HDS/CHEMOSAT system and the potential of the Melphalan HDS/CHEMOSAT 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, approval of the current or future Melphalan HDS/CHEMOSAT system for delivery and filtration of melphalan or other chemotherapeutic agents 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, 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.

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