Delcath Systems Announces Completion of Underwritten Public Offering

NEW YORK, Dec. 11, 2020 — Delcath Systems, Inc. (Nasdaq: DCTH), an interventional oncology company focused on the treatment of rare primary and metastatic cancers of the liver, today announced the completion of an underwritten public offering of 1,679,031 shares of its common stock, including 219,004 shares sold pursuant to the full exercise of an option previously granted to the underwriters to purchase additional shares of common stock. All of the shares were offered by Delcath at a price to the public of $13.25 per share. The Company now estimates that the gross proceeds of the offering to the Company are expected to be approximately $22.2 million, before deducting the underwriting discounts and commissions and other estimated offering expenses.

Delcath intends to use the net proceeds from this offering for (i) the completion of its FOCUS Clinical Trial for Patients with Hepatic Dominant Ocular Melanoma (the "Focus Trial"), a global registration clinical trial that is investigating the primary endpoint of objective response rate, as well as other secondary and exploratory endpoints, in metastatic ocular melanoma, or mOM; (ii) preparation of the federal regulatory application for the HEPZATO™ KIT (melphalan hydrochloride for injection/hepatic delivery system), or HEPZATO™, a drug/device combination product regulated as a drug, designed to administer high-dose chemotherapy to the liver while controlling systemic exposure and associated side effects; (iii) preparation for the commercial launch of HEPZATO; (iv) continued clinical development, including additional indications and expanded access trials in metastatic ocular melanoma; and (v) general corporate purposes, which may include capital expenditures and other operating expenses.

Canaccord Genuity and Roth Capital Partners acted as joint book-running managers for the offering.

A shelf registration statement relating to these securities has been filed with the U.S. Securities and Exchange Commission (SEC) and became effective on December 21, 2018. A final prospectus supplement relating to and describing the terms of the offering was filed with the SEC on December 9, 2020 and is available on the SEC's website at http://www.sec.gov. Copies of the final prospectus supplement and the accompanying prospectus relating to this offering may be obtained by contacting Canaccord Genuity LLC, Attention: Syndicate Department, 99 High Street, Suite 1200, Boston, MA 02110, by telephone at (617) 371-3900 or by email at prospectus@cgf.com or Roth Capital Partners, LLC, 888 San Clemente, Newport Beach, CA 92660, Attention: Prospectus Department, or by telephone at (800) 678-9147.

This press release shall not constitute an offer to sell or a solicitation of an offer to buy, nor shall there be any sale of these securities in any state or jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction.

About Delcath Systems, Inc.

Delcath Systems, Inc. is an interventional oncology company focused on the treatment of primary and metastatic liver cancers. Our investigational product, HEPZATO KIT (melphalan hydrochloride for injection/hepatic delivery system), is designed to administer high-dose chemotherapy to the liver while controlling systemic exposure and associated side effects. HEPZATO KIT has not been approved by the U.S. Food & Drug Administration (FDA) for sale in the U.S. In Europe, our system is marketed under the trade name Delcath CHEMOSAT® Hepatic Delivery System for Melphalan (CHEMOSAT) and has been CE Marked and used at major medical centers to treat a wide range of cancers of the liver. CHEMOSAT is being marketed under an exclusive licensing agreement with medac GmbH, a privately held multi-national pharmaceutical company headquartered in Germany that specializes in the treatment and diagnosis of oncological, urological and autoimmune diseases.

Safe Harbor / Forward-Looking Statements

The 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 mOM and ICC clinical trial programs, our determination whether to continue the ICC clinical trial program or to focus on other alternative indications, and timely monitoring and treatment of patients in the global Phase 3 mOM clinical trial and the impact of the COVID-19 pandemic on the completion of our clinical trials; 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 product; 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 HEPZATO KIT/CHEMOSAT system and the potential of the HEPZATO KIT/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 HEPZATO KIT/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 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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