# Delcath Systems to Host First Quarter 2024 Earnings Call

QUEENSBURY, N.Y., May 7, 2024 — Delcath Systems, Inc. (Nasdaq: DCTH), an interventional oncology company focused on the treatment of primary and metastatic cancers of the liver, announced today it will host a conference call on May 14, 2024, at 8:30 AM Eastern Time to discuss results for its first quarter ended March 31, 2024.

# Delcath

### **Conference Call Information**

To participate in this event, dial in approximately 5 to 10 minutes before the beginning of the call.

Event Date: Tuesday, May 14, 2024 Time: 8:30 AM Eastern Time

Participant Numbers: Toll Free: 1-833-630-1960 International: 1-412-317-1841 Webcast: https://app.webinar.net/PKDyZ5PV2aB

### **Conference Replay**

 US Toll Free:
 1-877-344-7529

 International Toll:
 1-412-317-0088

 Replay Access Code:
 9490444

 End Date:
 May 21, 2024

## About Delcath Systems, Inc., HEPZATO KIT and CHEMOSAT

Delcath Systems, Inc. is an interventional oncology company focused on the treatment of primary and metastatic liver cancers. The company's proprietary products, HEPZATO KIT (Hepzato (melphalan) for Injection/Hepatic Delivery System) and CHEMOSAT® Hepatic Delivery System for Melphalan percutaneous hepatic perfusion (PHP) are designed to administer high-dose chemotherapy to the liver while controlling systemic exposure and associated side effects during a PHP procedure.

In the United States, HEPZATO KIT is considered a combination drug and device product and is regulated and approved for sale as a drug by the FDA. HEPZATO KIT is comprised of the

chemotherapeutic drug melphalan and Delcath's proprietary Hepatic Delivery System (HDS). The HDS is used to surgically isolate the liver while simultaneously filtrating hepatic venous blood during melphalan infusion and washout. The use of the HDS results in loco-regional delivery of a relatively high melphalan dose, which can potentially induce a clinically meaningful tumor response with minimal hepatotoxicity and reduce systemic exposure. HEPZATO KIT is approved in the United States as a liver-directed treatment for adult patients with metastatic uveal melanoma (mUM) with unresectable hepatic metastases affecting less than 50% of the liver and no extrahepatic disease, or extrahepatic disease limited to the bone, lymph nodes, subcutaneous tissues, or lung that is amenable to resection or radiation. Please see the full Prescribing Information, including BOXED WARNING for the HEPZATO KIT.

In Europe, the device-only configuration of the HDS is regulated as a Class III medical device and is approved for sale under the trade name CHEMOSAT Hepatic Delivery System for Melphalan, or CHEMOSAT, where it has been used in the conduct of percutaneous hepatic perfusion procedures at major medical centers to treat a wide range of cancers of the liver. Please see the full Prescribing Information, including BOXED WARNING for the HEPZATO KIT.

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