

June 9, 2022



RenovoRx and TIGeR-PaC Leadership to Host RenovoTAMP™ Webinar on June 21, 2022

"Facing Today's Challenges in Pancreatic Cancer Treatment: Understanding RenovoRx's Trans-Arterial Micro-Perfusion Therapy Platform"

LOS ALTOS, Calif.--(BUSINESS WIRE)-- RenovoRx, Inc. (Nasdaq: RNXT), a biopharmaceutical company and innovator in targeted cancer therapy, today announced that they will host a webinar highlighting their RenovoTAMP™ therapy platform. The webinar, "Facing Today's Challenges in Pancreatic Cancer Treatment: Understanding RenovoRx's Trans-Arterial Micro-Perfusion Therapy Platform," will be held on Tuesday, June 21st at 8:00 am (ET). To register for this event, click [here](#).

Speakers include Dr. Michael Pishvaian, director of Gastrointestinal, Developmental Therapeutics and Clinical Research Programs for the Johns Hopkins Kimmel Cancer Center and an associate professor at the School of Medicine, and Shaun R. Bagai, RenovoRx's Chief Executive Officer.

During the presentation, Dr. Pishvaian, who also serves as Study Chair for the Company's Phase 3 TIGeR-PaC clinical trial, will discuss current available treatment options for pancreatic cancer and why they can fall short of providing an effective solution to fighting this deadly disease. He will also discuss how the TIGeR-PaC study is challenging this treatment paradigm by localizing the delivery of chemotherapy directly to the tumor, and how the RenovoRx Trans-Arterial Micro-Perfusion (RenovoTAMP) therapy platform re-envisioned the treatment of pancreatic and other solid tumors with the goal of improving survival and quality of life for patients undergoing chemotherapy. Dr. Pishvaian will also explain the scientific rationale underlying RenovoTAMP's development. Mr. Bagai will conclude by talking about the Company's plan for expanding the utility of its innovative therapy platform and its anticipated milestones for the remainder of the year. The virtual presentation will also include a Q & A session.

A recording of this presentation will be posted to the RenovoRx Website [Events](#) page when it becomes available.

About RenovoRx, Inc.

RenovoRx is a clinical-stage biopharmaceutical company focused on fighting cancer through the localized treatment of difficult to treat tumors via its proprietary RenovoRx Trans-Arterial Micro-Perfusion (RenovoTAMP™) therapy platform. RenovoTAMP utilizes approved chemotherapeutics with validated mechanisms of action and well-established safety and side effect profiles, with the goal of increasing their efficacy, improving their safety, and widening their therapeutic window. RenovoRx's lead product candidate, RenovoGem™, is a

combination of gemcitabine and our patented delivery system, RenovoCath[®], and is regulated by the FDA as a novel oncology drug product to treat unresectable locally advanced pancreatic cancer (LAPC). RenovoGem is currently being studied in the Phase 3 TIGeR-PaC trial for the treatment of LAPC.

RenovoRx's patent portfolio includes seven U.S. patents for its technology. RenovoRx has been granted Orphan Drug Designation for intra-arterial delivery of gemcitabine for the treatment of both pancreatic cancer and bile duct cancer.

RenovoRx won the Drug Delivery Technology category of the Fierce Innovation Awards – Life Sciences Edition 2020 for its RenovoTAMP technology.

Learn more by visiting the RenovoRx [website](#) or following us on [Facebook](#), [LinkedIn](#) and [Twitter](#).

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, and Section 21E of the Securities Exchange Act of 1934, including but not limited to statements regarding our clinical trials and studies, statements regarding the potential of RenovoTAMP[™], RenovoCath[®] or RenovoGem[™], or regarding our ongoing TIGeR-PaC Phase 3 clinical trial in LAPC, and statements regarding the potential for our product candidates to treat or provide clinically meaningful outcomes for certain medical conditions or diseases. Statements that are not purely historical are forward-looking statements. The forward-looking statements contained herein are based upon our current expectations and beliefs regarding future events, many of which, by their nature, are inherently uncertain, outside of our control and involve assumptions that may never materialize or may prove to be incorrect. These may include estimates, projections and statements relating to our research and development plans, clinical trials, therapy platform, business plans, objectives and expected operating results, which are based on current expectations and assumptions that are subject to known and unknown risks and uncertainties that may cause actual results to differ materially from those expressed or implied by these forward-looking statements. These statements may be identified using words such as “may,” “expects,” “plans,” “aims,” “anticipates,” “believes,” “forecasts,” “estimates,” “intends,” and “potential,” or the negative of these terms or other comparable terminology regarding RenovoRx's expectations strategy, plans or intentions, although not all forward-looking statements contain these words. These forward-looking statements are subject to a number of risks, uncertainties and assumptions, that could cause actual events to differ materially from those projected or indicated by such statements, including, among other things: the timing of the initiation, progress and potential results of our preclinical studies, clinical trials and our research programs; our ability to use and expand our therapy platform to build a pipeline of product candidates; our ability to advance product candidates into, and successfully complete, clinical trials; the timing or likelihood of regulatory filings and approvals; our estimates of the number of patients who suffer from the diseases we are targeting and the number of patients that may enroll in our clinical trials; the commercialization potential of our product candidates, if approved; our ability and the potential to successfully manufacture and supply our product candidates for clinical trials and for commercial use, if approved; future strategic arrangements and/or collaborations and the potential benefits of such arrangements; our estimates regarding expenses, future revenue,

capital requirements and needs for additional financing and our ability to obtain additional capital; the sufficiency of our existing cash and cash equivalents to fund our future operating expenses and capital expenditure requirements; our ability to retain the continued service of our key personnel and to identify, hire and retain additional qualified personnel; the implementation of our strategic plans for our business and product candidates; the scope of protection we are able to establish and maintain for intellectual property rights, including our therapy platform, product candidates and research programs; our ability to contract with third-party suppliers and manufacturers and their ability to perform adequately; the pricing, coverage and reimbursement of our product candidates, if approved; developments relating to our competitors and our industry, including competing product candidates and therapies; negative impacts of the COVID-19 pandemic on our operations; and other risks. Information regarding the foregoing and additional risks may be found in the section entitled "Risk Factors" in documents that we file from time to time with the Securities and Exchange Commission.

Forward-looking statements included herein are made as of the date hereof, and RenovoRx does not undertake any obligation to update publicly such forward-looking statements to reflect subsequent events or circumstances, except as required by law.

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