



REATA ANNOUNCES INITIATION OF INVESTIGATOR-SPONSORED STUDY EVALUATING BARDOXOLONE FOR COMPLICATIONS ASSOCIATED WITH COVID-19

PLANO, Texas—July 28, 2020 (GLOBE NEWSWIRE)—Reata Pharmaceuticals, Inc. (Nasdaq: RETA) (“Reata,” the “Company,” or “we”), a clinical-stage biopharmaceutical company, today announced that researchers at NYU Grossman School of Medicine (“NYU”), led by Sripal Bangalore, MD, interventional cardiologist and professor of Medicine, are initiating an Investigator-Sponsored Trial (“IST”), known as BARCONA, to study the effect of bardoxolone methyl (“bardoxolone”) in patients suffering from COVID-19. At NYU’s request, Reata is providing drug supply for the trial. NYU will be initiating the Phase 2 BARCONA study with a primary endpoint of safety.

The severity of COVID-19 and the development of systemic complications is associated with excessive, systemic inflammation, which can result in dysfunction of the lungs, kidneys, and other organs. Acute kidney injury (“AKI”) has been reported to occur in up to 28% of all hospitalized COVID-19 patients and up to 72% of patients who do not survive COVID-19. Bardoxolone and its analogs have demonstrated anti-inflammatory activity in animal models of acute lung and kidney injury and have shown improvements in kidney function in multiple clinical trials that enrolled over 3,000 patients with various forms of chronic kidney disease.

About Reata Pharmaceuticals, Inc.

Reata is a clinical-stage biopharmaceutical company that develops novel therapeutics for patients with serious or life-threatening diseases by targeting molecular pathways involved in the regulation of cellular metabolism and inflammation. Reata’s two most advanced clinical candidates, bardoxolone and omaveloxolone, target the important transcription factor Nrf2 that promotes the resolution of inflammation by restoring mitochondrial function, reducing oxidative stress, and inhibiting pro-inflammatory signaling. **Bardoxolone and omaveloxolone are investigational drugs, and their safety and efficacy have not been established by any agency.**

Contact:

Reata Pharmaceuticals, Inc.
(972) 865-2219
<http://reatapharma.com>

Investors:

Vinny Jindal
Vice President, Corporate Communications and Strategy
(469) 374-8721
ir@reatapharma.com
<http://reatapharma.com/contact-us/>



Forward-Looking Statements

This press release includes certain disclosures that contain “forward-looking statements,” including, without limitation, statements regarding the success, cost and timing of our product development activities and clinical trials, our plans to research, develop and commercialize our product candidates, our plans to submit regulatory filings, and our ability to obtain and retain regulatory approval of our product candidates. You can identify forward-looking statements because they contain words such as “believes,” “will,” “may,” “aims,” “plans,” “model,” and “expects.” Forward-looking statements are based on Reata’s current expectations and assumptions. Because forward-looking statements relate to the future, they are subject to inherent uncertainties, risks, and changes in circumstances that may differ materially from those contemplated by the forward-looking statements, which are neither statements of historical fact nor guarantees or assurances of future performance. Important factors that could cause actual results to differ materially from those in the forward-looking statements include, but are not limited to, (i) the timing, costs, conduct, and outcome of our clinical trials and future preclinical studies and clinical trials, including the timing of the initiation and availability of data from such trials; (ii) the timing and likelihood of regulatory filings and approvals for our product candidates; (iii) whether regulatory authorities determine that additional trials or data are necessary in order to obtain approval; (iv) the potential market size and the size of the patient populations for our product candidates, if approved for commercial use, and the market opportunities for our product candidates; and (v) other factors set forth in Reata’s filings with the U.S. Securities and Exchange Commission, including the detailed factors discussed under the caption “Risk Factors” in its Annual Report on Form 10-K for the fiscal year ended December 31, 2019. The forward-looking statements speak only as of the date made and, other than as required by law, we undertake no obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events, or otherwise.