



Cara Therapeutics to Present at the 2018 Cantor Global Healthcare Conference

September 26, 2018

STAMFORD, Conn., Sept. 26, 2018 (GLOBE NEWSWIRE) -- Cara Therapeutics, Inc. (Nasdaq:CARA), a biopharmaceutical company focused on developing and commercializing new chemical entities designed to alleviate pruritus and pain by selectively targeting peripheral kappa opioid receptors, today announced a Company presentation at the 2018 Cantor Global Healthcare Conference on Wednesday, October 3, 2018, at 12:15 p.m. ET in New York, NY.

A live webcast of the presentation can be accessed under "Events and Presentations" in the News & Investors section of the Company's website at www.CaraTherapeutics.com. An archived webcast recording will be available on the Cara website for approximately 30 days.

About Cara Therapeutics

Cara Therapeutics is a clinical-stage biopharmaceutical company focused on developing and commercializing new chemical entities designed to alleviate pruritus and pain by selectively targeting peripheral kappa opioid receptors. Cara is developing a novel and proprietary class of product candidates, led by KORSUVA™ (CR845/difelikefalin), a first-in-class kappa opioid receptor agonist that targets the body's peripheral nervous system, as well as certain immune cells. In Phase 2 trials, KORSUVA injection has demonstrated statistically significant reductions in itch intensity and concomitant improvement in quality of life measures in hemodialysis patients suffering from moderate-to-severe chronic kidney disease-associated pruritus (CKD-aP). KORSUVA is currently being investigated in global Phase 3 trials in hemodialysis patients with CKD-aP. Additionally, in a recently completed Phase 2/3 trial in post-operative patients, I.V. CR845/difelikefalin has demonstrated reduction in moderate-to-severe pain, while also reducing the incidence and intensity of nausea and vomiting throughout the post-operative period.

The FDA has conditionally accepted KORSUVA™ as the trade name for difelikefalin injection. CR845/difelikefalin is an investigational drug product, and its safety and efficacy have not been fully evaluated by any regulatory authority.

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Source: Cara Therapeutics, Inc.