



September 5, 2017

Evoke Pharma to Present at the 19th Annual Rodman & Renshaw Global Investment Conference

SOLANA BEACH, Calif., Sept. 05, 2017 (GLOBE NEWSWIRE) -- Evoke Pharma, Inc. (NASDAQ:EVOK), a specialty pharmaceutical company focused on treatments for gastrointestinal (GI) diseases, today announced that it will be featured as a presenting company at the 19th Annual Rodman & Renshaw Global Investment Conference, being held on September 10-12, 2017 at The Lotte New York Palace Hotel in New York City.

David Gonyer, Evoke's President and Chief Executive Officer, will provide a business overview and update on the Company's development of Gimoti™ during the live presentation and will be available to participate in one-on-one meetings with investors who are registered to attend the conference.

If you are an institutional investor, and would like to attend the Company's presentation, please click on the link (www.rodmanevents.com) to register for the conference. Once your registration is confirmed, you will be prompted to log into the conference website to request a one-on-one meeting with the Company.

Details for the presentation are as follows:

Date: Tuesday, September 12, 2017

Time: 10:00 — 10:25 a.m. EDT

Room: Louis

About Evoke Pharma, Inc.

Evoke is a specialty pharmaceutical company focused primarily on the development of drugs to treat GI disorders and diseases. The Company is developing Gimoti, a metoclopramide nasal spray for the relief of symptoms associated with acute and recurrent gastroparesis in women with diabetes mellitus. Diabetic gastroparesis is a disorder afflicting millions of sufferers worldwide, in which the stomach takes too long to empty its contents resulting in serious digestive system symptoms. Metoclopramide is the only product currently approved in the United States to treat gastroparesis, and is currently available only in oral and intravenous forms. Gimoti is a novel formulation of this drug, designed to provide systemic delivery of metoclopramide through nasal administration. Visit www.EvokePharma.com for more information.

Safe Harbor Statement

Evoke cautions you that statements included in this press release that are not a description of historical facts are forward-looking statements. In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "expect," "plan," "anticipate," "could," "intend," "target," "project," "contemplates," "believes," "estimates," "predicts," "potential" or "continue" or the negatives of these terms or other similar expressions. These statements are based on the company's current beliefs and expectations. These forward-looking statements include statements regarding: Evoke's plans to include the PK data in the 505(b)(2) NDA for Gimoti; the timing of announcement of the results of the PK trial and the timing of the submission of the NDA to the FDA; Evoke's expectation that Spaulding Clinical Research will complete the study; Evoke's expectation that the PK trial will be the final clinical trial for Gimoti; and Evoke's belief that there is a large unmet need for an effective treatment for diabetic gastroparesis. Actual results may differ from those set forth in this press release due to the risks and uncertainties inherent in Evoke's business, including, without limitation: risks associated with successfully initiating, conducting and receiving favorable results from the PK trial; later developments with the FDA that may be inconsistent with the already completed pre- NDA meetings, including inconsistent conclusions reflected in the official meeting minutes from the FDA; the inherent risks of clinical development of Gimoti; Evoke is entirely dependent on the success of Gimoti, and Evoke cannot be certain that it will be able to submit an NDA for Gimoti or obtain regulatory approval for or successfully commercialize Gimoti; risks associated with manufacturing new formulations of Gimoti for use in the PK trial; Evoke's dependence on third parties for the manufacture of Gimoti as well as the submission of the NDA; Evoke's dependence on Spaulding Clinical Research to conduct the PK trial; Evoke may require additional funding to complete the PK trial and submit the NDA, and will require substantial additional funding to commercialize Gimoti, and may

be unable to raise capital when needed, including to fund ongoing operations; and other risks detailed in Evoke's prior press releases and in the periodic reports it files with the Securities and Exchange Commission. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof, and Evoke undertakes no obligation to revise or update this press release to reflect events or circumstances after the date hereof. All forward-looking statements are qualified in their entirety by this cautionary statement. This caution is made under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995.

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