Relmada Therapeutics to Present at the 28th Annual ROTH Conference

Description

Company Presentation scheduled for March 16, 2016 at 11:00 a.m. PT.

NEW YORK, March 8, 2016 /PRNewswire/ — Relmada Therapeutics, Inc. (OTCQB: RLMD), a clinical-stage company developing novel therapies for the treatment of chronic pain, today announced that the Company's management will present at the 28th Annual ROTH Conference, which is being held March 13-16, 2016 in Orange County, CA.

Relmada Therapeutics Corporate Logo

Sergio Traversa, Chief Executive Officer of Relmada Therapeutics, will provide an overview of the Company and recent activities, including topline results of a proof-of-concept pharmacokinetic study in healthy volunteers using BuTab (REL-1028), an investigational, oral formulation of buprenorphine, an opioid that is broadly used to treat both addiction and chronic pain. The presentation is scheduled for Wednesday, March 16, 2016 at 11:00 a.m. PT (2:00 p.m. ET).

Members of the investment community who are interested in meeting with Relmada's management team during the conference event should contact their ROTH representative or Christine Silverstein, director of investor relations at Relmada Therapeutics, at (646) 677-3858 or csilverstein@relmada.com.

A live webcast of Relmada's presentation will be available through the Company's website at www.relmada.com. Please register at least 10 minutes prior to the start of the presentation to ensure timely access. The webcast and presentation will also be archived on the website for 90 days after the conference.

About Relmada Therapeutics, Inc.

Relmada Therapeutics is a clinical-stage, publicly traded specialty pharmaceutical company developing novel versions of proven drug products together with new chemical entities that potentially address areas of high unmet medical need in the treatment of pain. The Company has a diversified portfolio of four lead products at various stages of development including d-Methadone (REL-1017) its N-methyl-D-aspartate (NMDA) receptor antagonist for neuropathic pain; topical mepivacaine (REL-1021), its orphan drug designated topical formulation of the local anesthetic mepivacaine; oral buprenorphine (REL-1028) its oral dosage form of the opioid analgesic buprenorphine; and LevoCap ER (REL-1015), its abuse resistant, sustained release dosage form of the opioid analgesic levorphanol. The Company's product development efforts are guided by the internationally recognized scientific expertise of its research team. The Company's approach is expected to reduce clinical development risks and costs while potentially delivering valuable products in areas of high unmet medical needs. For more information, please visit Relmada's website at: www.relmada.com.

Forward-Looking Statements

The Private Securities Litigation Reform Act of 1995 provides a safe harbor for forward-looking statements made by us or on our behalf. We may from time to time make written or oral statements in this letter, the proxy statements filed with the SEC communications to stockholders and press releases which constitute “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. These forward-looking statements are based upon management’s current expectations, estimates, assumptions and beliefs concerning future events and conditions and may discuss, among other things, anticipated future performance, expected product development, product potential, future business plans and costs. Any statement that is not historical in nature is a forward-looking statement and may be identified by the use of words and phrases such as “expects,” “anticipates,” “believes,” “will,” “will likely result,” “will continue,” “plans to” and similar expressions. No forward-looking statement can be guaranteed and actual results may differ materially from those projected. Relmada undertakes no obligation to publicly update any forward-looking statement, whether as a result of new information, future events, or otherwise. Readers are cautioned that it is not possible to predict or identify all of the risks, uncertainties and other factors that may affect future results and that the risks described herein should not be considered to be a complete list.

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