
UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d) of the
Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): November 5, 2015

RELMADA THERAPEUTICS, INC.
(Exact name of registrant as specified in its charter)

Nevada

(State or other jurisdiction of
incorporation)

333-184881

(Commission File Number)

45-5401931

(IRS Employer Identification No.)

**757 Third Avenue Avenue, Suite 2018
New York, NY**

(Address of principal executive offices)

10017

(Zip Code)

Registrant's telephone number, including area code **(212) 376-5776**

N/A

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
 - Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
 - Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
 - Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))
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Item 8.01. Other Events.

On November 5, 2015, Relmada Therapeutics, Inc. (the “Company”) issued a press release in response to the Schedule 13D filing by Laidlaw & Company (UK) Ltd. on October 22, 2015. A copy of the Company’s press release is attached as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated into this Item 8.01 by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description
99.1	Relmada Therapeutics, Inc. Press Release, dated November 5, 2015.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Dated: November 5, 2015

RELMADA THERAPEUTICS, INC.

By: /s/ Sergio Traversa
Name: Sergio Traversa
Title: Chief Executive Officer



Relmada Therapeutics Responds to 13D Filing By Laidlaw

NEW YORK, November 5, 2015 - Relmada Therapeutics, Inc. (OTCQB: RLMD), a clinical-stage company developing novel therapies for the treatment of chronic pain, today issued the following statement in response to the Schedule 13D filing by Laidlaw & Company (UK) Ltd. ("**Laidlaw**") on October 22, 2015:

While Relmada welcomes the views and input of all of its shareholders, the Board of Directors and management of the Company strongly disagree with Laidlaw's mischaracterizations and "proposals" set forth in their 13D filing. Such filing, which appears not to have been timely made, contains numerous inaccuracies and appears to disclose information about Relmada that could only have been obtained in violation of Laidlaw's responsibilities to the Company.

The Board and management are committed to maximizing value for all shareholders of the Company and firmly believe that can best be achieved by continuing to focus on the successful execution of the Company's strategy. That strategy, which includes the continual evaluation of in-licensing and out-licensing opportunities, is to develop the Company's four unique product candidates to treat chronic pain.

In addition, the Company has implemented a number of initiatives, including the previously disclosed filing of an application to list the Company's common stock on the NASDAQ Capital Market, intended to make the Company's shares more available and attractive to a larger audience of potential investors.

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.

Relmada Therapeutics
W: www.relmada.com | E: info@relmada.com



Forward-Looking Statements

This news release contains “forward-looking statements.” These statements are based on management’s current expectations and involve risks and uncertainties, which may cause actual results to differ materially from those set forth in the statements. The forward-looking statements may include statements regarding product development, product potential, or financial performance. 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.

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