
**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 6, 2015

LIXTE BIOTECHNOLOGY HOLDINGS, INC.

DELAWARE
(State or other jurisdiction
of incorporation)

000-51436
(Commission
File Number)

20-2903526
(IRS Employer
Identification No.)

248 Route 25A, No. 2
East Setauket, New York 11733
(Address of principal executive offices)

Registrant's telephone number, including area code: 631 942 7959

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 of 1933 (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(e) under the Exchange Act (17 CFR 240.13e-4(c))
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Item 8.01 Other Events

On November 6, 2015, the Company issued a press release regarding certain developments.

Item 9.01 Financial Statements and Exhibits

(d) Exhibits

There is filed as part of this report the exhibit listed on the accompanying Index to Exhibits, which information is incorporated herein by reference.

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.

Date: November 9, 2015

LIXTE BIOTECHNOLOGY HOLDINGS, INC.

By: /s/ JOHN S. KOVACH

John S. Kovach, Chief Executive Officer

Index to Exhibits

Exhibit No. Description

99.1 Press Release.

LIXTE BIOTECHNOLOGY HOLDINGS, INC. ANNOUNCES THAT ITS NOVEL PROTEIN PHOSPHATASE 2A INHIBITOR IS ASSOCIATED WITH STABILIZATION OF SEVERAL TYPES OF CANCER WITHOUT DOSE-LIMITING TOXICITY

East Setauket, New York, November 6, 2015 - Lixte Biotechnology Holdings, Inc. ([OTCQB: LIXT](#)) announced today that in an ongoing Phase I trial its lead anti-cancer compound, LB-100, was associated with stabilization of a variety of advanced cancers that had been progressing despite extensive prior treatment. The results were presented at the AACR-NCI-EORTC Molecular Targets and Cancer Therapeutics Conference, Boston, on November 6. The authors were Vincent Chung, City of Hope, Duarte, CA; Donald Richards, Texas Oncology, Tyler, TX; Fadi Braiteh, Comprehensive Cancer Centers of Nevada, Las Vegas, NV; John S. Kovach, Lixte Biotechnology Holdings, Inc., East Setauket, NY; Aaron Scott Mansfield, Mayo Clinic, Rochester, MN.

A total of 21 patients received LB-100 for 3-consecutive days in 3-week cycles. Nine of these patients had stabilization of their disease without significant toxicity. One patient with pancreatic cancer received 14 cycles of LB-100 (42 weeks); one with thymoma, 9 cycles; one with testicular and one with carcinoid of the lung, 5 cycles; two with ovarian, 3 and 6 cycles; and one with NSCLC and one with duodenal cancer, 3 cycles. The average number of cycles received at the last four evaluable dose levels of LB-100 was 4.8 (14.4 weeks). In a study by Mansfield et al (2015 JCO 33[15] suppl:2567), the average number of cycles of a new drug in 51 NCI-sponsored Phase I trials conducted from 1994-2014 involving 1841 patients was 2.0 (6-8 weeks).

John S. Kovach M.D., founder and president of Lixte, said “Lixte interprets the results as showing that LB-100 has single agent activity in suppressing the growth of several types of cancer. What is highly encouraging is that stabilization of disease occurred in the absence of dose-limiting toxicity. In fact, most patients tolerated repeated doses without any ill-effects. As pre-clinical studies have shown that LB-100 potentiates the effectiveness of cytotoxic agents, we believe that LB-100 alone and in combination with standard anti-cancer drugs and/or radiation may offer new therapeutic options for a spectrum of neoplastic diseases.”

About Lixte Biotechnology Holdings, Inc.

Lixte is a drug discovery company that uses biomarker technology to identify enzyme targets associated with serious common diseases and then design novel compounds to attack those targets. Lixte’s product pipeline encompasses two major categories of compounds at various stages of pre-clinical and clinical development which the Company believes have broad therapeutic potential not only for cancer but for other debilitating and life-threatening diseases. Lixte’s unique phosphatase inhibitor, LB-100, is in a Phase I clinical trial at two NCI designated Comprehensive Cancer Centers and three US Oncology Research sites (see [ClinicalTrials.gov: Identifier NCT01837667](#)).

Forward-Looking Statements .

This announcement contains certain 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. For example, statements regarding the Company’s financial position, business strategy and other plans and objectives for future operations, and assumptions and predictions about future product demand, supply, manufacturing, costs, marketing and pricing factors are all forward-looking statements. These statements are generally accompanied by words such as “intend,” “anticipate,” “believe,” “estimate,” “potential(ly),” “continue,” “forecast,” “predict,” “plan,” “may,” “will,” “could,” “would,” “should,” “expect” or the negative of such terms or other comparable terminology. The Company believes that the assumptions and expectations reflected in such forward-looking statements are reasonable, based on information available to it on the date hereof, but the Company cannot provide assurances that these assumptions and expectations will prove to have been correct or that the Company will take any action that the Company may presently be planning. However, these forward-looking statements are inherently subject to known and unknown risks and uncertainties. Actual results or experience may differ materially from those expected or anticipated in the forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to, regulatory policies, available cash, research results, competition from other similar businesses, and market and general economic factors. This discussion should be read in conjunction with the Company’s filings with the United States Securities and Exchange Commission at <http://www.sec.gov/edgar.shtml>.

Additional information on the Company is available at www.liخته.com.

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