
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): October 10, 2018

ZIOPHARM Oncology, Inc.
(Exact Name of Registrant as Specified in Charter)

Delaware
(State or Other Jurisdiction
of Incorporation)

001-33038
(Commission File Number)

84-1475642
(IRS Employer
Identification No.)

One First Avenue, Parris Building 34, Navy Yard Plaza
Boston, Massachusetts
(Address of Principal Executive Offices)

02129
(Zip Code)

(617) 259-1970
(Registrant's telephone number, including area code)

Not applicable
(Former Name or Former Address, if Changed Since Last Report)

Check the appropriate box below if the Form 8-K is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- 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)).

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act (17 CFR 230.405) or Rule 12b-2 of the Exchange Act (17 CFR 240.12b-2).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 5.02. Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers.

Election of New Director

Effective October 15, 2018, Laurence Cooper, M.D., Ph.D., the President and Chief Executive Officer of Ziopharm Oncology, Inc., or the Company, was elected as a director to fill an existing vacancy on the Company's Board of Directors, or the Board. Dr. Cooper will serve as a director until the Company's 2019 Annual Meeting of Stockholders and until such time as his successor is duly elected and qualified, or until his earlier death, resignation or removal. Dr. Cooper has not been appointed to any committee of the Board.

The compensation of Dr. Cooper, as the Company's President and Chief Executive Officer, is described in the Company's definitive proxy statement on Schedule 14A relating to its 2018 Annual Meeting of Stockholders, or the 2018 Proxy Statement, which was filed with the Securities and Exchange Commission on August 8, 2018. Dr. Cooper will not receive any additional compensation for his service on the Board.

Dr. Cooper was not selected as a director pursuant to any arrangements or understandings with the Company or with any other person. Except as described in the 2018 Proxy Statement, there are no related party transactions between Dr. Cooper and the Company that would require disclosure under Item 404(a) of Regulation S-K.

Departure of Executive Officer

On October 10, 2018, Francois Lebel, M.D. notified the Company that he was stepping down from his position as the Company's Executive Vice President, Research and Development and Chief Medical Officer, effective October 26, 2018.

On October 16, 2018, the Company issued a press release announcing the election of Dr. Cooper to the Board and Dr. Lebel's departure from the Company. The press release is attached to this Current Report on Form 8-K as Exhibit 99.1.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit

<u>No.</u>	<u>Description</u>
99.1	Press release of Ziopharm Oncology, Inc. dated October 16, 2018.

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.

ZIOPHARM ONCOLOGY, INC.

Date: October 16, 2018

By: /s/ Robert Hadfield

Name: Robert Hadfield

Title: General Counsel and Secretary



Ziopharm Oncology Announces Changes to Board of Directors and Management Team

*- CEO Laurence Cooper Appointed to Board of Directors
- Francois Lebel to Depart Company*

BOSTON, October 16, 2018 (GLOBE NEWSWIRE) — Ziopharm Oncology, Inc. (Nasdaq:ZIOP), a biotechnology company focused on development of next generation immunotherapies utilizing gene- and cell-based therapies to treat patients with cancer, today announced changes to its Board of Directors and management team. Ziopharm’s Chief Executive Officer Laurence Cooper, M.D., Ph.D, is appointed to the Board of Directors effective immediately, and Francois Lebel, M.D., is stepping down from his position as Chief Medical Officer (CMO) and Executive Vice President of Research & Development, effective Oct. 26.

“We are pleased to continue the evolution of the Board of Directors with the addition of Dr. Cooper as the Company’s CEO,” said Ziopharm’s Lead Director Scott Tariff, who is Chief Executive Officer of Eagle Pharmaceuticals. “We support Laurence and his plan for the Company’s management team to advance the Controlled IL-12 and *Sleeping Beauty* platforms.”

Ziopharm recently announced it has full developmental control over its technologies and is now evolving its organizational structure. A recruitment process is underway during this transition period aimed at building upon existing expertise to meet the needs of patients and shareholders.

“We are developing and expanding our R&D and clinical development teams to support the Controlled IL-12 and *Sleeping Beauty* CAR-T and TCR-T programs,” said Dr. Cooper. “On behalf of the Board of Directors and management team, I thank Francois for his service to our clinical programs, and we wish him well in his next endeavor.”

Dr. Lebel said, “I very much enjoyed working with the team at Ziopharm and the many world-class investigators involved with our trials. I am proud that the team has established a strong data set that supports the potential efficacy of our technologies, including Controlled IL-12 to treat patients with recurrent glioblastoma and likely other solid tumors. As I leave the Company, I believe it is on a solid track for success.”

About Ziopharm Oncology, Inc.

Ziopharm Oncology is a Boston-based biotechnology company focused on the development of next-generation immunotherapies utilizing gene- and cell-based therapies to treat patients with cancer. Ziopharm is focused on the development of two platform technologies designed to deliver safe, effective and scalable cell- and viral-based therapies for the treatment of multiple cancer types: Controlled IL-12 and *Sleeping Beauty* for genetically modifying T cells. The Company's lead asset, Ad-RTS-hIL-12 plus veledimex, has demonstrated in clinical trials the potential to control interleukin-12, leading to an infiltration of T cells that fight cancer. Ad-RTS-hIL-12 plus veledimex is being evaluated as a monotherapy and in combination with immune checkpoint inhibitors to treat brain cancer and other tumor types. The Company also is advancing therapies using *Sleeping Beauty*, a non-viral approach to genetically modify chimeric antigen receptor (CAR+) and T-cell receptor (TCR+) T cells, which target specific antigens in blood cancers and neoantigens such as in solid tumors. *Sleeping Beauty* is designed using the Company's "point-of-care" technology, a shortened manufacturing process which potentially can be developed as a decentralized manufacturing process based in hospitals. These programs are being advanced in collaboration with MD Anderson Cancer Center and the National Cancer Institute.

Forward-Looking Statements Disclaimer

This press release contains certain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended. Forward-looking statements are statements that are not historical facts, and in some cases can be identified by terms such as "may," "will," "could," "expects," "plans," "anticipates," and "believes." All such statements are subject to certain risks and uncertainties, many of which are difficult to predict and generally beyond the control of the Company, that could cause actual results to differ materially from those expressed in, or implied by, the forward-looking statements. These risks and uncertainties include, but are not limited to: the Company's ability to advance certain activities; whether chimeric antigen receptor T cell (CAR-T) approaches, Ad-RTS-hIL-12, TCR and T-cell receptor (TCR) cell-based therapies, or any other product candidates will advance further in the preclinical research or clinical trial process and whether and when, if at all, they will receive final approval from the U.S. Food and Drug Administration or equivalent foreign regulatory agencies and for which indications; whether chimeric antigen receptor T cell (CAR-T) approaches, Ad-RTS-hIL-12 and TCR cell-based therapies, and the Company's other therapeutic products it develops will be successfully marketed if approved; the strength and enforceability of the Company's intellectual property rights; competition from other pharmaceutical and biotechnology companies; as well as other risk factors contained in the Company's periodic and interim reports filed from time to time with the Securities and Exchange Commission, including but not limited to, the risks and uncertainties set forth in the "Risk Factors" section of the Company's Annual Report on Form 10-K for the fiscal year ended December 31, 2017 and subsequent reports that the Company may file with the Securities and Exchange Commission. Readers are cautioned not to place undue reliance on these forward-looking statements that speak only as of the date hereof, and the Company does not undertake any obligation to revise and disseminate forward-looking statements to reflect events or circumstances after the date hereof, or to reflect the occurrence of or non-occurrence of any events.

For more information contact:

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