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 7, 2016

Dynavax Technologies Corporation
(Exact name of registrant as specified in its charter)

Commission File Number: 001-34207

Delaware	**33-0728374**
(State or other jurisdiction of incorporation)	**(IRS Employer Identification No.)**

2929 Seventh Street, Suite 100
Berkeley, CA 94710-2753
(Address of principal executive offices, including zip code)

(510) 848-5100
(Registrant's telephone number, including area code)

(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:

☐ **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))**

Item 2.02. Results of Operations and Financial Condition

On November 7, 2016, Dynavax Technologies Corporation ("Dynavax") issued a press release announcing its financial results for the quarter ended September 30, 2016. A copy of the press release is attached as Exhibit 99.1 to this current report and is incorporated herein by reference.

The information with respect to item 2.02 in this current report and its accompanying exhibit shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained in this current report and in the accompanying exhibit shall not be incorporated by reference into any filing with the U.S. Securities and Exchange Commission made by Dynavax, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

Item 9.01. Financial Statements and Exhibits

(d) Exhibits. The following exhibit is furnished herewith:

99.1 Press Release, dated November 7, 2016, titled "Dynavax Reports Third Quarter 2016 Financial Results And Company Update"

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.

Dynavax Technologies Corporation

Date: November 7, 2016

By: /s/ DAVID JOHNSON
David Johnson
Vice President

EXHIBIT INDEX

Exhibit No.	Description
EX-99.1	Press Release, dated November 7, 2016, titled “Dynavax Reports Third Quarter 2016 Financial Results And Company Update”

DYNAVAX

INNOVATING IMMUNOLOGY

2929 Seventh Street, Suite 100
Berkeley, CA 94710

Contact:
Michael Ostrach
Chief Financial Officer
510-665-7257
mostrach@dynavax.com

DYNAVAX REPORTS THIRD QUARTER 2016 FINANCIAL RESULTS AND COMPANY UPDATE

BERKELEY, CA – November 7, 2016 – Dynavax Technologies Corporation (NASDAQ: DVAX) today reported financial results for the third quarter and nine months ended September 30, 2016.

The Company had $109.6 million in cash, cash equivalents and marketable securities as of September 30, 2016, compared to $196.1 million at December 31, 2015. The net loss for the third quarter of 2016 was $34.7 million, compared to $30.1 million for the third quarter of 2015.

Recent Progress

HEPLISAV-B. In late August, the U.S. Food and Drug Administration (FDA) cancelled its previously scheduled Vaccines and Related Biological Products Advisory Committee (VRBPAC) meeting to review the Biologics License Application (BLA) for HEPLISAV-B™ [Hepatitis B Vaccine, Recombinant (Adjuvanted)]. The FDA indicated that remaining questions on the BLA will be addressed between Dynavax and the FDA review team. The Company has since provided responses to information requests by the FDA related to remaining questions. The FDA also confirmed in August that it will not include in its review of the BLA the immunogenicity data submitted by the Company related to sub-populations, including results in individuals with diabetes. The Company plans to submit these data as a supplemental BLA.

The Prescription Drug User Fee Act (PDUFA) date for the HEPLISAV-B BLA is December 15, 2016.

In late October, we reported sub-group results from HBV-23, demonstrating that HEPLISAV-B, when administered as two doses over one month, induced significantly higher seroprotection rates than the approved hepatitis B vaccine Engerix-B®, when administered as three doses over six months. This result was observed in all prespecified groups of study participants, including those with characteristics that are known to have a reduced immune response to currently licensed hepatitis B vaccines, including older age, high body mass index, diabetes mellitus, male gender and persons who smoke. In the total Phase 3 trial population, the rates of adverse events, serious adverse events and deaths were similar between the HEPLISAV-B and Engerix-B groups. The data were presented at the Infectious Diseases Society of America's (IDSA) annual IDWeek 2016 meeting in New Orleans.

Preparations for launch of HEPLISAV-B are continuing, including pre-commercial activities, manufacturing of launch inventory and continued infrastructure spending related to commercial development and information technology capabilities and related increases in headcount.

Immuno-oncology. In October we announced at the European Society of Medical Oncology (ESMO) Annual Congress 2016 the first presentation of findings from an ongoing Phase 1/2 study evaluating SD-101, Dynavax's intratumoral TLR9 agonist, in combination with Keytruda ® (pembrolizumab), Merck's anti-PD-1 treatment. Early results evaluating five patients with metastatic melanoma for efficacy and 16 patients for safety were reported. In patients naïve to anti-PD-1 treatment objective responses were observed in three out of four (75%) including one complete response (CR) and two partial responses (PR's). One patient with progressive disease while receiving anti-PD-1 therapy was observed to have stable disease (SD). The drug combination was well-tolerated. No dose-limiting toxicities of the combination were observed in any dose cohort, and a maximum tolerated dose (MTD) was not identified. No immune-related adverse events were reported. Additional results from this study will be presented at future scientific meetings.

Financial. In late October we secured a $100 million loan commitment from Deerfield upon FDA approval of HEPLISAV-B and satisfaction of other conditions. We intend to use the net proceeds for general corporate purposes, including the commercialization of HEPLISAV-B.

Financials

Total revenues for the third quarter of 2016 were $0.2 million compared to $1.2 million for the same period in 2015. The $1.0 million decrease was due to the conclusion of our work under the research collaboration and license agreement with AstraZeneca for the clinical development of AZD 1419.

Research and development expenses for the third quarter of 2016 were $23.2 million compared to $24.1 million for the same period in 2015. This $0.9 million decrease was primarily due to reduction in outside services expense associated with the completion of HBV-23 in the fourth quarter of 2015, partially offset by an increase in employee headcount and regulatory and manufacturing activities in preparation for the anticipated commercial launch of HEPLISAV-B.

General and administrative expenses for the third quarter of 2016 were $11.8 million compared to $5.5 million for the same period in 2015. This $6.3 million increase reflects expenses related to preparation for the commercial launch of HEPLISAV-B including additional headcount, information technology systems and infrastructure to support commercial development.

The net loss for the third quarter of 2016 was $34.7 million, or $0.90 per basic and diluted share, compared to $30.1 million, or $0.82 per basic and diluted share, for the same period in 2015.

About Dynavax

Dynavax, a clinical-stage biopharmaceutical company, uses TLR biology to discover and develop novel vaccines and therapeutics in the areas of infectious and inflammatory diseases and oncology. Dynavax's lead product candidates are HEPLISAV-B, a Phase 3 investigational adult hepatitis B vaccine, and SD-101, an investigational cancer immunotherapeutic currently in several Phase 1/2 studies. For more information visit www.dynavax.com.

Forward Looking Statements

This release contains forward-looking statements, including statements regarding anticipated approval and launch of HEPLISAV-B. These statements are subject to a number of risks and uncertainties that could cause actual results to differ materially, including whether there will be the need for additional studies, further manufacturing enhancements or other activities, a VRBPAC will be scheduled, whether the FDA will find our responses to their questions to be sufficient, or other issues will arise that will negatively impact the review, duration of review and approval of the BLA by the FDA; whether we will successfully launch the product, possible claims against us, including enjoining sales of HEPLISAV-B based on the patent rights of others, and the potential size and value of approved indications addressable with HEPLISAV-B; whether we will satisfy the conditions to the Deerfield loan; initiation and completion of pre-clinical studies and clinical trials of our other product candidates, including SD-101, in a timely manner; the results of clinical trials and the impact of those results on the initiation or continuation of subsequent trials and issues arising in the regulatory process; our ability to execute on our commercial strategies; whether our financial resources will be adequate without the need to obtain additional financing and other risks detailed in the "Risk Factors" section of our most recent current periodic report filed with the SEC. These statements represent our estimates and assumptions only as of the date of this release. We do not undertake any obligation to update publicly any such forward-looking statements, even if new information becomes available.

DYNAVAX TECHNOLOGIES CORPORATION
CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS
(In thousands, except per share amounts)
(Unaudited)

	Three Months Ended September 30,		Nine Months Ended September 30,	
	2016	**2015**	**2016**	**2015**
Revenues:				
Collaboration revenue	$ -	$ 829	$ 2,578	$ 2,230
Grant revenue	162	359	289	608
Service and license revenue	-	-	884	527
Total revenues	162	1,188	3,751	3,365
Operating expenses:				
Research and development	23,234	24,105	66,051	66,011
General and administrative	11,766	5,524	29,086	15,481
Total operating expenses	35,000	29,629	95,137	81,492
Loss from operations	(34,838)	(28,441)	(91,386)	(78,127)
Interest income	170	33	615	78
Interest expense	-	(62)	-	(572)
Other (expense) income, net	(26)	17	68	360
Loss on extinguishment of debt	-	(1,671)	-	(1,671)
Net loss	$ (34,694)	$ (30,124)	$ (90,703)	$ (79,932)
Basic and diluted net loss per share	$ (0.90)	$ (0.82)	$ (2.36)	$ (2.43)
Weighted average shares used to compute basic and diluted net loss per share	38,512	36,532	38,493	32,880

DYNAVAX TECHNOLOGIES CORPORATION
SELECTED BALANCE SHEET DATA
(In thousands)
(Unaudited)

	September 30, 2016	December 31, 2015
Assets		
Cash, cash equivalents and marketable securities	$ 109,551	$ 196,125
Property and equipment, net	18,739	13,804
Goodwill	2,100	2,043
Other assets	8,908	4,661
Total assets	$ 139,298	$ 216,633
Liabilities and stockholders' equity		
Deferred revenues	$ -	$ 2,654
Other liabilities	31,026	26,900
Total liabilities	31,026	29,554
Stockholders' equity	108,272	187,079
Total liabilities and stockholders' equity	$ 139,298	$ 216,633