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): June 11, 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 8.01. Other Events

On June 11, 2016, Dynavax issued a press release titled “Dynavax Presents New Efficacy Data on Hepatitis B Vaccine, HEPLISAV-B™, in Adults with Type 2 Diabetes at ADA Scientific Sessions.” A copy of the press release is attached hereto as Exhibit 99.1 and incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

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

99.1 Press Release, dated June 11, 2016, titled “Dynavax Presents New Efficacy Data on Hepatitis B Vaccine, HEPLISAV-B™, in Adults with Type 2 Diabetes at ADA Scientific Sessions”

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: June 15, 2016

By: /s/ MICHAEL OSTRACH
Michael Ostrach
Senior Vice President

EXHIBIT INDEX

Exhibit No.	Description
EX-99.1	Press Release, dated June 11, 2016, titled “Dynavax Presents New Efficacy Data on Hepatitis B Vaccine, HEPLISAV-B™, in Adults with Type 2 Diabetes at ADA Scientific Sessions”

DYNAVAX

INNOVATING IMMUNOLOGY

Dynavax Presents New Efficacy Data on Hepatitis B Vaccine, HEPLISAV-B™, in Adults with Type 2 Diabetes at ADA Scientific Sessions

-- Pivotal Phase 3 data show investigational two-dose hepatitis B vaccine provides a significantly higher rate of seroprotection than an approved vaccine in patients with type 2 diabetes --

BERKELEY, Calif. – June 11, 2016 -- Dynavax Technologies Corporation (NASDAQ: DVAX) today announced preliminary results from a pivotal Phase 3 trial demonstrating that HEPLISAV-B [Hepatitis B Vaccine, Recombinant (Adjuvanted)]provided a significantly higher rate of seroprotection than Engerix-B®, an approved hepatitis B vaccine, in adults with type 2 diabetes mellitus. The data are being presented at the 76th Annual Scientific Sessions of the American Diabetes Association (ADA) in New Orleans.

Adults with diabetes are particularly vulnerable to infection with hepatitis B. According to the Centers for Disease Control and Prevention (CDC), people ages 23 to 59 with diabetes are about twice as likely to develop acute hepatitis B as individuals without diabetes. The CDC recommends that all adults age 19 to 59 should be vaccinated against hepatitis B as soon as possible after their diagnosis of diabetes, and those who are age 60 and older may be vaccinated at their physician's discretion after assessing their risk and the likelihood of an adequate immune response. Approximately 20 million U.S. adults have diabetes, and 1.5 million new cases of diabetes are diagnosed each year.

The Phase 3 trial, HBV-23, was a randomized, observer-blinded, active-controlled, multi-center study that compared two doses of HEPLISAV-B with three doses of Engerix-B in adults age 18 to70. Among the over 8,000 randomized participants, there were 1,144 adults with type 2 diabetes of whom two-thirds had diabetes for five years or more. Demographics consisting of age, sex and race were generally similar between the two treatment arms.

Results showed that HEPLISAV-B provided seroprotection in 90.0 percent of participants with diabetes compared with 65.1 percent for Engerix-B – a statistically significant difference of 24.9 percent. Larger differences were observed in participants age 60 to 70, with HEPLISAV-B demonstrating an 85.8 percent rate of seroprotection compared with 58.5 percent for Engerix-B. For participants with a body mass index greater than or equal to 30, HEPLISAV-B demonstrated an 89.5 percent rate of seroprotection compared to 61.4 percent for Engerix-B.

In the total trial population, the rates of adverse events, serious adverse events and deaths were similar between the HEPLISAV-B and Engerix-B groups. All adverse events considered to represent potential immune-mediated disorders were reviewed by an independent, blinded Safety Evaluation and Adjudication Committee, which classified these events as not related to vaccination.

“The hepatitis B virus can spread easily through contact with contaminated medical equipment, such as blood glucose monitors, posing a serious health risk to people living with diabetes,” said Rob Janssen, M.D., chief medical officer for Dynavax. “Results of this study show that, with two doses over one month, HEPLISAV-B provided a significantly higher rate of seroprotection in participants with type 2 diabetes than an existing hepatitis B vaccine.”

The U.S. Food and Drug Administration has established December 15, 2016, as the Prescription Drug User Fee Act (PDUFA) action date for the HEPLISAV-B Biologics License Application.

About Hepatitis B
Hepatitis B is a viral disease of the liver that can become chronic and can lead to cirrhosis of the liver, hepatocellular carcinoma and death. In the U.S., the CDC estimates that approximately 20,000 hepatitis B infections continue to occur annually, with the vast majority occurring in adults. There is no cure for hepatitis B, and disease prevention through effective vaccination is critical to reducing the spread of the disease. Currently marketed hepatitis B vaccines are administered in three doses over a six-month schedule. Results of a published Vaccine Safety Datalink study showed that only 54 percent of adults completed the three-dose hepatitis B vaccine series in one year. Those who do not complete the series may not be adequately protected against hepatitis B.

About HEPLISAV-B
HEPLISAV-B is an investigational adult hepatitis B vaccine that combines hepatitis B surface antigen with a proprietary Toll-like Receptor 9 agonist to enhance the immune response. In Phase 3 trials, HEPLISAV-B demonstrated higher and earlier protection with fewer doses than Engerix-B, a currently licensed hepatitis B vaccine. HEPLISAV-B was administered in two doses over one month.

The investigational vaccine’s safety profile is based on clinical trials that generated safety data from more than 14,000 participants. The most frequently reported local reaction was injection site pain. The most common systemic reactions were fatigue, headache and malaise, all of which were similar to Engerix-B.

Dynavax has worldwide commercial rights to HEPLISAV-B.

About Dynavax
Dynavax, a clinical-stage biopharmaceutical company, discovers and develops novel vaccines and therapeutics in the areas of infectious 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 press release contains forward-looking statements, including statements regarding HEPLISAV-B and FDA review. These statements are subject to a number of risks and uncertainties that could cause actual results to differ materially, including whether there will be changes in the data or interpretation; whether the final study results will be deemed satisfactory by the FDA; whether additional studies or manufacturing process enhancements will be required or other issues will arise that will delay the BLA review or negatively impact the review and approval by the FDA; initiation, enrollment and completion of pre-clinical studies and clinical trials of our other product candidates, including SD-101; 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; 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 press release. We do not undertake any obligation to update publicly any such forward-looking statements, even if new information becomes available. Information on Dynavax's website at www.dynavax.com is not incorporated by reference in our current periodic reports with the SEC.

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