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): 09/28/2010

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 September 28, 2010, Dynavax Technologies Corporation (Dynavax) issued a press release titled "Dynavax's Two Phase 3 HEPLISAV Trials Cleared by DSMB to Continue." A copy of the press release is attached as Exhibit 99.1 to this current report and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

(d) Exhibit

Exhibit No.	Description
99.1	Press Release, dated September 28, 2010, titled "Dynavax's Two Phase 3 HEPLISAV Trials Cleared by DSMB to Continue."

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: September 28, 2010

By: /s/ Michael S. Ostrach

Michael S. Ostrach
Vice President

EXHIBIT INDEX

Exhibit No.	Description
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Exhibit 99.1

DYNAVAX

DYNAVAX TECHNOLOGIES

2929 Seventh Street, Suite 100

Berkeley, CA 94710

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Michael Ostrach
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DYNAVAX'S TWO PHASE 3 HEPLISAV TRIALS CLEARED BY DSMB TO CONTINUE

Third Safety Assessment Complete

BERKELEY, CA - September 28, 2010 - Dynavax Technologies Corporation (NASDAQ: DVAX) today announced that the Data Safety Monitoring Board (DSMB) established for Dynavax's two ongoing Phase 3 trials for HEPLISAV™ has completed the third of its planned safety assessments. The DSMB determined that the studies may continue without protocol modification. The Phase 3 lot-to-lot consistency trial was fully enrolled in May 2010; the Phase 3 chronic kidney disease study is expected to complete enrollment within the next few months.

The evaluation included data from 2717 subjects. Of the subjects enrolled in the Phase 3 lot-to-lot consistency study, all are at least two months past their final HEPLISAV injections. Furthermore, the first subjects enrolled in the Phase 3 chronic kidney disease study are now 12 months past their first dose.

Tyler Martin, M.D., President and Chief Medical Officer, commented, "The DSMB's recommendation that both Phase 3 trials continue without protocol modification is consistent with our expectation. Generally, one expects adverse events related to study drugs to occur in temporal proximity to the exposure. With 100% of subjects in the Phase 3 lot-to-lot consistency study now two months past the last HEPLISAV vaccination, our confidence in the safety of this product is reinforced. "

Dino Dina, M.D., Chief Executive Officer, added, "We have clearly advanced past a key milestone in both studies and are working to stay on our projected timeline for making regulatory submissions before the end of 2011. We have demonstrated HEPLISAV's efficacy in multiple prior trials and expect these trials to demonstrate efficacy in line with our previous results. The first of these Phase 3 trials was designed to confirm the consistency of our lots and to provide us a safety database that will be acceptable to the regulatory authorities in the U.S. and in Europe; the second was designed to demonstrate a regimen specifically for CKD patients. "

HEPLISAV is an innovative investigational vaccine designed to protect against hepatitis B infection. The DSMB reviewed safety data from two ongoing multi-center Phase 3 trials evaluating HEPLISAV, the first a lot-to-lot consistency trial in adults 40 years and older, and the second a trial in chronic kidney disease patients. The DSMB is comprised of an independent group of medical experts who are responsible for reviewing and evaluating subject safety data at regular intervals during the ongoing trials.

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Conference Call

Dynavax will webcast a conference call today at 4:10 p.m. EDT (1:10 p.m. PDT). The live and archived webcast can be accessed by visiting the investor relations section of the Company's website at http://investors.dynavax.com/newsevents.cfm.

About HEPLISAV

HEPLISAV is an investigational adult hepatitis B vaccine. The vaccine candidate is being evaluated in two Phase 3 studies that are directed toward fulfilling licensure requirements in U.S., Canada and Europe. In a completed pivotal Phase 3 trial, HEPLISAV demonstrated increased, rapid protection with fewer doses than current licensed vaccines. Dynavax has worldwide commercial rights to HEPLISAV and is developing the vaccine for large, high-value populations that are less responsive to current licensed vaccines, including individuals with chronic kidney disease. HEPLISAV combines hepatitis B surface antigen with a proprietary Toll-like Receptor 9 agonist known as ISS to enhance the immune response.

About Dynavax

Dynavax Technologies Corporation, a clinical-stage biopharmaceutical company, discovers and develops novel products to prevent and treat infectious diseases. The Company's lead product candidate is HEPLISAV, an investigational adult hepatitis B vaccine designed to enhance protection more rapidly and with fewer doses than current licensed vaccines. For more information visit www.dynavax.com.

Forward Looking Statements

This press release contains "forward-looking statements," that are subject to a number of risks and uncertainties. Actual results may differ materially from those set forth in this press release due to the risks and uncertainties inherent in our business, including whether successful clinical and regulatory development and approval of HEPLISAV can occur in a timely manner or without significant additional studies or difficulties or delays in development or clinical trial enrollment, whether the studies can support registration for commercialization of HEPLISAV; the results of clinical trials and the impact of those results on the initiation and completion of subsequent trials and issues arising in the regulatory process; the Company's ability to obtain additional financing to support the development and commercialization of HEPLISAV and its other operations, possible claims against the Company based on the patent rights of others; and other risks detailed in the "Risk Factors" section of our current p eriodic reports with the SEC. We undertake no obligation to revise or update information herein to reflect events or circumstances in the future, even if new information becomes available. Information on Dynavax's website at www.dynavax.com is not incorporated by reference in the Company's current periodic reports with the SEC.

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