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): 02/13/2012

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 February 13, 2012, we issued a press release titled "Phase 3 Data on HEPLISAV in Adults Aged 18-55 Published in Vaccine." 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) Exhibits

Exhibit No.	Description
99.1	Press Release, dated February 13, 2012, titled "Phase 3 Data on HEPLISAV in Adults Aged 18-55 Published in Vaccine."

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: February 14, 2012

By: /s/ Michael S. Ostrach

Michael S. Ostrach
Vice President

EXHIBIT INDEX

Exhibit No.	Description
EX-99.9	PHASE 3 DATA ON HEPLISAV IN ADULTS AGED 18-55 PUBLISHED IN VACCINE

2929 Seventh Street, Suite 100

Berkeley, CA 94710

Contact:
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Vice President and Chief Business Officer
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PHase 3 data on Heplisav in adults aged 18-55 published in vaccine

Berkeley, CA - February 13, 2012 - Dynavax Technologies Corporation (NASDAQ: DVAX) announced today that results of a pivotal Phase 3 trial of HEPLISAV (HBV-10) were published online in the journal VACCINE. Data from this study will be used to support the HEPLISAV Biologics License Application (BLA) submission for an indication in adults 18-70 years of age. The article concludes that a short, two-dose regimen of HEPLISAV over 1 month was well-tolerated and induced superior immunogenicity and earlier onset of protection than a three-dose regimen of a licensed hepatitis B vaccine over 6 months.

The article entitled "Comparison of Safety and Immunogenicity of Two Doses of Investigational Hepatitis B Virus Surface Antigen Co-administered with an Immunostimulatory Phosphorothioate Oligodeoxyribonucleotide and Three Doses of a Licensed Hepatitis B Vaccine in Healthy Adults 18-55 Years of Age" describes the results from one of the two pivotal Phase 3 trials of HEPLISAV. Dr. Scott Halperin of Dalhousie University was the principal investigator and lead author. The trial compared the safety and immunogenicity of HEPLISAV with Engerix-B® in 2,415 adults randomized in a ratio of 3:1, HEPLISAV to Engerix-B. The seroprotection rate at the primary endpoint after 2 doses for HEPLISAV (95%) was significantly higher than after 3 doses for Engerix-B (81%). Superiority of the seroprotection rates for HEPLISAV was demonstrated at all time points measured.

Dynavax plans to submit the BLA for HEPLISAV by the middle of May for an indication in adults 18-70 years of age.

About HEPLISAV

HEPLISAV is an investigational adult hepatitis B vaccine. In earlier Phase 3 trials, HEPLISAV demonstrated increased, rapid protection with fewer doses than current licensed vaccines. Dynavax has worldwide commercial rights to HEPLISAV. HEPLISAV combines hepatitis B surface antigen with a proprietary Toll-like Receptor 9 agonist known as 1018 ISS to enhance the immune response.

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About Dynavax

Dynavax Technologies Corporation, a clinical-stage biopharmaceutical company, discovers and develops novel products to prevent and treat infectious and inflammatory diseases. The Company's lead product candidate is HEPLISAV, a Phase 3 investigational adult hepatitis B vaccine designed to provide rapid and superior protection with fewer doses than current licensed vaccines. For more information visit www.dynavax.com.

Forward-Looking Statements

This press release contains "forward-looking statements," including those relating to the HEPLISAV BLA, planned indication, and timing of the submission, 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 and our process for its manufacture can occur in a timely manner or without significant additional studies or difficulties or delays in development or clinical trial enrollment, whether our 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, including whether the BLA will be accepted for filing; 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 periodic 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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