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): 06/03/2009

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 3, 2009, Dynavax Technologies Corporation (Dynavax) issued a press release titled "Dynavax Presents Additional Phase 3 Data for HEPLISAV(TM) Hepatitis B Vaccine at DDW Medical Conference." 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 June 3, 2009, titled "Dynavax Presents Additional Phase 3 Data for HEPLISAV(TM) Hepatitis B Vaccine at DDW Medical Conference."

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 03, 2009

By: /s/ Michael S. Ostrach

Michael S. Ostrach
Vice President

Exhibit Index

Exhibit No.	Description
EX-99.1	Press Release, dated June 3, 2009, titled "Dynavax Presents Additional Phase 3 Data for HEPLISAV(TM) Hepatitis B Vaccine at DDW Medical Conference."

DYNAVAX

DYNAVAX TECHNOLOGIES
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DYNAVAX PRESENTS ADDITIONAL PHASE 3 DATA FOR HEPLISAV(TM) HEPATITIS B VACCINE AT DDW MEDICAL CONFERENCE

-- HEPLISAV Demonstrates Rapid, Increased Protection for Adults --

BERKELEY, CA – June 3, 2009 – Dynavax Technologies Corporation (Nasdaq: DVAX) today presented additional Phase 3 clinical data for HEPLISAV(TM) hepatitis B vaccine in a poster session at the Digestive Disease Week (DDW) medical conference in Chicago. In addition to meeting its primary endpoint in this Phase 3 trial as previously reported, HEPLISAV provided more rapid and increased seroprotection against hepatitis B viral infection and with fewer doses than the licensed vaccine. The data show the differences to be particularly significant in the subset of subjects over 40 years of age who are usually less likely to respond to immunization.

“Vaccination is critical for the prevention of hepatitis B viral infection and its spread, but the lengthy dosing regimen of current vaccines leaves many unprotected,” commented Scott A. Halperin, M.D., Director Clinical Trials Research Center, Dalhousie University, IWK Heath Centre of Halifax, Nova Scotia and Lead Investigator in the Phase 3 trial. “The Phase 3 data demonstrate HEPLISAV’s superior immunogenicity and similar safety profile versus the licensed vaccine, particularly for adults over 40 who are more difficult to protect against this preventable viral infection.”

This Phase 3 trial referred to as PHAST (Phase 3 HeplisAv Short-regimen Trial) evaluated more than 2,400 adults. The seroprotection rate at the primary endpoint was 95% in subjects receiving 2 doses of HEPLISAV at 0 and 1 month, compared to 81% in subjects receiving 3 doses of licensed vaccine Engerix-B(R) at 0, 1, and 6 months.

In a subanalysis of subjects over 40 years of age, at each time point during the trial there was a statistically significant ($p < 0.0001$) difference in the seroprotection rate for subjects receiving HEPLISAV or Engerix-B.

Ages 40-55 Treatment Group	Dosing Regimen				Seroprotection Rate (1) at Month				
					1	2	3	6	7
HEPLISAV	2	doses	(0,	1 month)	18%	84%	92% (2)	97%	97%
Engerix-B	3	doses	(0,	1, 6 months)	3%	21%	17%	27%	75% (2)

(1) Seroprotection rate – percentage of subjects with anti-HBsAg antibodies 10 mIU/mL

(2) Primary endpoint

-- More --

A copy of the poster is available at http://investors.dynavax.com/newsevents.cfm. Dynavax’s abstract #587659 is titled “A Phase 3 Safety and Efficacy Study to Compare Immune Responses following Either Two Doses of Hepatitis B Surface Antigen Combined with Immunostimulatory Phosphorothioate Oligonucleotide (HBsAg-ISS) or Three Doses of Conventional Hepatitis B Vaccine.”

As previously reported, safety results from this trial demonstrated the safety profile of HEPLISAV and Engerix-B appeared similar. Subjects were randomized 3 to 1 to receive HEPLISAV or Engerix-B and one case of vasculitis was reported in each of the treatment groups. Following the report of the severe adverse event of Wegener's granulomatosis, an uncommon form of vasculitis, HEPLISAV was placed and remains on clinical hold by the U.S. Food and Drug Administration (FDA). Dynavax is in active discussions with regulatory agencies to resolve the FDA’s clinical hold on HEPLISAV and identify an appropriate path for its further development and approval in the United States, Europe, and the rest of the world.

About HEPLISAV

HEPLISAV is a Phase 3 hepatitis B vaccine aimed at unmet medical needs in the vaccination of adults and end-stage renal disease patients by providing rapid and increased protection with fewer doses. HEPLISAV combines a proprietary immunostimulatory sequence (ISS), which targets Toll-like Receptor 9, with hepatitis B surface antigen (HBsAg).

About Hepatitis B

Hepatitis B is a chronic disease which can lead to cirrhosis of the liver and hepatocellular carcinoma. There is no cure for hepatitis B and disease prevention through effective vaccines is critical to reducing the spread of the disease. Current hepatitis B vaccines for adults usually require 3 doses given over 6 months to provide seroprotection of approximately 30%, 75%, and 90% after the first, second, and third doses respectively. The effectiveness of current vaccines is further compromised because only 30% of people receive all 3 doses.

About Dynavax

Dynavax Technologies Corporation, a clinical-stage biopharmaceutical company, discovers and develops a diversified pipeline of novel Toll-like Receptor (TLR) based product candidates. Based on Dynavax's proprietary technologies, these products specifically modify the innate immune response to infectious, respiratory, autoimmune, and inflammatory diseases. Dynavax has partnerships with leading pharmaceutical companies such as GlaxoSmithKline, AstraZeneca, and Novartis as well as funding from Symphony Dynamo, Inc. and the National Institutes of Health. For more information visit www.dynavax.com.

About DDW

DDW is the largest international gathering of physicians, researchers and academics in the fields of gastroenterology, hepatology, endoscopy and gastrointestinal surgery. Jointly sponsored by the American Association for the Study of Liver Diseases, the American Gastroenterological Association (AGA) Institute, the American Society for Gastrointestinal Endoscopy and the Society for Surgery of the Alimentary Tract, DDW takes place May 30 – June 4, 2009, at the McCormick Place, Chicago, IL. The meeting showcases approximately 5,000 abstracts and hundreds of lectures on the latest advances in GI research, medicine and technology. For more information, visit www.ddw.org.

Forward Looking Statements

This press release contains "forward-looking statements," that are subject to a number of risks and uncertainties, including statements related to the nature of communications with regulatory agencies regarding HEPLISAV. 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 the FDA will remove the clinical hold for HEPLISAV, whether HEPLISAV can be further developed, financed or commercialized, or even if further development is permitted, that successful clinical development and regulatory approval can occur in a timely manner or without significant additional studies and difficulties or delays in development, the Company's ability to obtain

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additional financing to support its operations; 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.

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