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): 12/07/2010

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 December 7, 2010, Dynavax Technologies Corporation (Dynavax) issued a press release titled "Dynavax Reports Phase 1a Safety and Immunogenicity Results for Universal Flu 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) Exhibit

Exhibit No.	Description
99.1	Press Release, dated December 7, 2010, titled "Dynavax Reports Phase 1a Safety and Immunogenicity Results for Universal Flu 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: December 07, 2010

By: /s/ Michael S. Ostrach

Michael S. Ostrach
Vice President

EXHIBIT INDEX

Exhibit No.	Description
EX-99.1	Press Release, dated December 7, 2010, titled "Dynavax Reports Phase 1a Safety and Immunogenicity Results for Universal Flu Vaccine."

DYNAVAX

DYNAVAX TECHNOLOGIES
2929 Seventh Street, Suite 100
Berkeley, CA 94710

Contact:
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Vice President and Chief Business Officer
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DYNAVAX REPORTS PHASE 1A SAFETY AND IMMUNOGENICITY RESULTS FOR UNIVERSAL FLU VACCINE

Detailed Phase 1a and 1b Results to be Reported at WHO Meeting in February 2011

BERKELEY, CA – December 7, 2010 – Dynavax Technologies Corporation (Nasdaq: DVAX) today reported safety and immunogenicity data from its Phase 1a clinical trial of N8295, one of two key components of its Universal Flu Vaccine candidate. N8295 is a fusion protein comprised of NP and M2e, two highly conserved influenza antigens covalently linked to Dynavax's proprietary second-generation TLR9 agonist. The trial assessed three dose levels of N8295 in a total study population of 39 subjects. The Phase 1a data showed:

- All doses were very safe and generally well tolerated;
- No dose limiting toxicities;
- Positive antibody responses to M2e; and
- Positive T-cell mediated responses to NP.

Based on preliminary safety data, Dynavax initiated a Phase 1b study in September 2010 to evaluate the safety of the combination of N8295, the novel component of Dynavax's Universal Flu vaccine candidate, and an investigational H5N1 avian influenza vaccine. Detailed results of the Phase 1a and 1b studies will be reported at the World Health Organization 7th Meeting on Evaluation of Pandemic Influenza Prototype Vaccines in Clinical Trials in Geneva, Switzerland in February 2011.

Dr. J. Tyler Martin, M.D., Dynavax President and Chief Medical Officer, said, “Now that we have completed the safety assessment of the novel component, N8295, we are eager to assess the combination of N8295 with an avian flu vaccine. Those data are expected to improve our understanding of the immunologic properties of our universal flu vaccine candidate in the absence of pre-existing immunity to the H5N1 flu strain in human subjects. These are key achievements in the continued development of Dynavax's Universal Flu Vaccine as they should allow us to design a proof-of-concept study.“

Dynavax's Universal Flu Vaccine is designed to offer protection against divergent influenza strains as well as to increase the efficacy of a conventional influenza vaccine. Preclinical data have confirmed the expected immunogenicity and mechanistic effects of the vaccine candidate's novel components. The production of cytotoxic T-cells by NP and cytotoxic

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antibodies by M2e have been demonstrated in preclinical studies, as has an increase in neutralizing antibodies provided by a co-administered conventional influenza vaccine. A GLP toxicity study demonstrated that this Universal Flu vaccine candidate is well-tolerated.

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(TM), a Phase 3 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, including statements related to the anticipated timing and nature of data from clinical trials of our universal flu vaccine candidate and the features of the vaccine. 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 results of completed studies can be replicated in human studies with greater numbers of subjects, difficulties or delays in initiation and completion of preclinical or clinical studies, the results of those studies and the impact of those results on the initiation and completion of subsequent studies and issues arising in the regulatory process; achieving the objectives under our collaborative agreement; our ability to obtain additional financing to support our operations; and other risks detailed in the "Risk Factors" section of our current periodic reports filed 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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