

AFFYMAX INC  
Form 8-K  
June 21, 2010

**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

Washington, DC 20549

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**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 21, 2010**

**AFFYMAX, INC.**

(Exact name of registrant as specified in charter)

**Delaware**  
(State or other jurisdiction of  
incorporation)

**001-33213**  
(Commission File Number)

**77-0579396**  
(I.R.S. Employer Identification No.)

**4001 Miranda Avenue  
Palo Alto, California 94304**  
(Address of principal executive offices and zip code)

Registrant's telephone number, including area code: **(650) 812 - 8700**

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## Edgar Filing: AFFYMAX INC - Form 8-K

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(b) under the Exchange Act (17 CFR 240.14a-12(b))
  
  - 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))
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**Item 8.01. Other Events.**

On June 21, 2010, Affymax, Inc. (the Company), Takeda Global Research & Development, Inc. and Takeda Pharmaceutical Company, Ltd ( Takeda ) announced preliminary top line results from the Hematide /peginesatide Phase 3 clinical program for the treatment of anemia associated with chronic renal failure anemia. A press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

In connection with database locks for Phase 3 clinical trials, the Company has received \$30 million of milestone payments from Takeda.

**Item 9.01. Financial Statement and Exhibits.**

(d) Exhibits:

<b>Exhibit No.</b>	<b>Description</b>
99.1	Press Release entitled Affymax and Takeda announce Phase 3 Trials Meet Primary Endpoints for Investigational Drug, Hematide /peginesatide, to Treat Anemia in Chronic Renal Failure with Some Differences Noted in Secondary Analyses, dated June 21, 2010.

**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.

**AFFYMAX, INC .**

Dated: June 21, 2010

By:

*/s/ Paul B. Cleveland*

Paul B. Cleveland  
Executive Vice President, Corporate Development and  
Chief Financial Officer

**EXHIBIT INDEX**

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