IMMUNOGEN INC
Form 8-K
July 07, 2010

# 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): **July 6, 2010**

# ImmunoGen, Inc.

(Exact name of registrant as specified in its charter)

| **Massachusetts** | **0-17999** | **04-2726691** |
|---|---|---|
| (State or other jurisdiction of incorporation) | (Commission File Number) | (IRS Employer Identification No.) |

**830 Winter Street, Waltham, MA 02451**

(Address of principal executive offices) (Zip Code)

Registrant s telephone number, including area code: **(781) 895-0600**

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 (*see* General Instruction A.2. below):

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

**ITEM 8.01. OTHER EVENTS**

Genentech, a member of the Roche Group, and Roche simultaneously announced on the evening of July 6/morning of July 7, 2010, respectively, the company s submission of a Biologics License Application, or BLA, to the US Food and Drug Administration to gain marketing approval in the US of trastuzumab-DM1, or T-DM1, to treat patients with advanced HER2-positive breast cancer who had previously received multiple chemotherapies and HER2-targeted medicines. The basis of the submission is the Phase II trial that was reported at the San Antonio Breast Cancer Symposium in December 2009.

T-DM1 consists of ImmunoGen, Inc. s DM1 cancer-cell killing agent attached to Roche s HER2-targeting antibody, trastuzumab, using ImmunoGen s linker and methods of attachment. T-DM1 is in global development by Roche under a collaboration agreement between Genentech and ImmunoGen.

**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.

**ImmunoGen, Inc.**
(Registrant)

Date: July 7, 2010

/s/ Gregory D. Perry

Gregory D. Perry
Senior Vice President and Chief Financial Officer