

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): **November 10, 2010**

**BioTime, Inc.**

(Exact name of registrant as specified in its charter)

**California**

(State or other jurisdiction of  
incorporation)

**1-12830**

(Commission File Number)

**94-3127919**

(IRS Employer Identification  
No.)

**1301 Harbor Bay Parkway, Suite 100  
Alameda, California 94502**

(Address of principal executive offices)

**(510) 521-3390**

(Registrant's telephone number, including area code)

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))
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*Statements made in this Report that are not historical facts may constitute forward-looking statements that are subject to risks and uncertainties that could cause actual results to differ materially from those discussed. Such risks and uncertainties include but are not limited to those discussed in this report and in BioTime's other reports filed with the Securities and Exchange Commission. Words such as "expects," "may," "will," "anticipates," "intends," "plans," "believes," "seeks," "estimates," and similar expressions identify forward-looking statements.*

## **Section 8 – Other Events**

### **Item 8.01 – Other Events**

On December 15, 2010 our subsidiary Embryome Sciences, Inc. will add a group of new products to its product line. The new products will include 31 new human embryonic progenitor cell lines, associated cell culture media, 53 diverse extracellular matrices, and 62 diverse extracts from conditioned media, all of which will be offered for research use only. Additional information about these new products will be found at [www.embryome.com](http://www.embryome.com) beginning with product launch.

Human embryonic progenitor cells are intermediate in the developmental process between embryonic stem cells and fully differentiated cells. The cells may possess the ability to become a wide array of cell types with potential applications in research, drug discovery, and potential novel regenerative stem cell therapies. The cells are relatively easy to manufacture on a large scale and in a purified state, which may make it advantageous to work with these cells compared to the direct use of human embryonic stem cells.

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## Section 9-Financial Statements and Exhibits

### Item 9.01 Financial Statements and Exhibits.

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press release dated November 10, 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.

#### BIOTIME, INC.

Date: November 10, 2010

By: /s/ Robert W. Peabody  
Senior Vice President, Chief Operating  
Officer and Chief Financial Officer

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press release dated November 10, 2010

**BioTime's CEO Michael West to Present Keynote Address at BIT Life Sciences' 3<sup>rd</sup>  
Annual World Congress of Regenerative Medicine & Stem Cells 2010 in Shanghai, China**

ALAMEDA, Calif.--(BUSINESS WIRE)--November 10, 2010--BioTime, Inc. (NYSE Amex: BTX) announced today that BioTime's CEO Dr. Michael West will present the keynote address at the BIT Life Sciences' 3<sup>rd</sup> Annual World Congress of Regenerative Medicine & Stem Cells 2010 in Shanghai, China on December 5, 2010. Dr. West's presentation will be titled "Regenerative Medicine: A Force for Change in Biotechnology." During his talk, Dr. West will present a group of new research products that the Company's subsidiary Embryome Sciences, Inc. will be launching beginning December 15, 2010. The new products will include 31 new human embryonic progenitor cell lines, associated cell culture media, 53 diverse extracellular matrices, and 62 diverse extracts from conditioned media, all of which will be offered for research use only. Additional information about these new products will be found at [www.embryome.com](http://www.embryome.com) beginning with product launch. Dr. West's presentation will be available on BioTime's website at [www.biotimeinc.com](http://www.biotimeinc.com).

***About BioTime, Inc.***

BioTime, headquartered in Alameda, California, is a biotechnology company focused on regenerative medicine and blood plasma volume expanders. Its broad platform of stem cell technologies is developed through subsidiaries focused on specific fields of applications. BioTime develops and markets research products in the field of stem cells and regenerative medicine through its wholly owned subsidiary Embryome Sciences, Inc. BioTime's therapeutic product development strategy is pursued through subsidiaries that focus on specific organ systems and related diseases for which there is a high unmet medical need. BioTime's majority owned subsidiary Cell Cure Neurosciences, Ltd. is developing therapeutic products derived from stem cells for the treatment of retinal and neural degenerative diseases. Cell Cure's minority shareholder Teva Pharmaceutical Industries has optioned its OpRegen<sup>TM</sup> retinal cell product for use in the treatment of Age-related Macular Degeneration (AMD). BioTime's subsidiary OrthoCyte Corporation is developing therapeutic applications of stem cells to treat orthopedic diseases and injuries. Another subsidiary, OncoCyte Corporation, focuses on the therapeutic applications of stem cell technology in cancer. BioTime's Singapore subsidiary, ES Cell International Pte Ltd, has been at the forefront of advances in human embryonic stem ("hES") cell technology, having been one of the earliest distributors of hES cell lines to the research community. ESI has produced clinical-grade human embryonic stem cell lines that were derived following principles of Good Manufacturing Practice and currently offers them for potential use in therapeutic product development. In addition to its stem cell products, BioTime develops blood plasma volume expanders, blood replacement solutions for hypothermic (low temperature) surgery, and technology for use in surgery, emergency trauma treatment and other applications. BioTime's lead product, Hextend®, is a blood plasma volume expander manufactured and distributed in the U.S. by Hospira, Inc. and in South Korea by CJ CheilJedang Corp. under exclusive licensing agreements. Additional information about BioTime, Embryome Sciences, Cell Cure, OrthoCyte, OncoCyte, BioTime Asia, and ESI can be found on the web at [www.biotimeinc.com](http://www.biotimeinc.com).

***Forward-Looking Statements***

Statements pertaining to future financial and/or operating results, future growth in research, technology, clinical development, and potential opportunities for the company and its subsidiaries, along with other statements about the future expectations, beliefs, goals, plans, or prospects expressed by management constitute forward-looking statements. Any statements that are not historical fact (including, but not limited to statements that contain words such as "will," "believes," "plans," "anticipates," "expects," "estimates") should also be considered to be forward-looking statements. Forward-looking statements involve risks and uncertainties, including, without limitation, risks inherent in the development and/or commercialization of potential products, uncertainty in the results of clinical trials or regulatory approvals, need and ability to obtain future capital, and maintenance of intellectual property rights. Actual results may differ materially from the results anticipated in these forward-looking statements and as such should be evaluated together with the many uncertainties that affect the company's business, particularly those mentioned in the cautionary statements found in the company's Securities and Exchange Commission filings. The company disclaims any intent or obligation to update these forward-looking statements.

To receive ongoing BioTime corporate communications, please click on the following link to join our email alert list:  
<http://www.b2i.us/irpass.asp?BzID=1152&to=ea&s=0>

**CONTACT:**

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