

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): **March 4, 2011**

**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  
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 7 - Regulation FD

### Item 7.01 - Regulation FD Disclosure

On March 4, 2011 BioTime, Inc. issued the press release filed as Exhibit 99.1, which is incorporated by reference.

## 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 March 4, 2011 |

## 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: March 4, 2011

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

Exhibit Number

99.1

Description

Press release dated March 4, 2011

**BioTime CEO Michael West to Present at French-American Biotech Symposium, San Francisco, April 11-12, 2011*****Update on ReCyte™ technology for resetting telomere length  
and its application in cardiovascular disease***

ALAMEDA, Calif.--(BUSINESS WIRE)--March 4, 2011--BioTime, Inc. (NYSE Amex:BTX) and its subsidiary ReCyte Therapeutics, Inc. announced today that BioTime CEO Dr. Michael West will present at the French-American Biotech Symposium on "New Therapeutic Approaches of Aging" in San Francisco, April 11-12, 2011. Dr. West will speak on "Resetting Telomere Length using Transcriptional Reprogramming and its Application in Age-Related Degenerative Disease" on Tuesday, April 12, during Session III: Innovative Approaches to Chronic Diseases Associated with Aging.

On March 16, 2010, BioTime announced in a peer-reviewed scientific publication that iPS cell reprogramming technologies could, under certain conditions, reverse the developmental aging of human cells. In other words, by genetically modifying aged cells from the human body, they could be reverted back to an embryonic state similar to that of embryonic stem cells, which are capable of developing into all other cell and tissue types. In addition, it was previously reported that these reprogramming technologies could reset the telomere clock of cellular aging. On April 12, Dr. West will be presenting new data validating this technology. In particular, evidence will be presented that the technique is capable not only of resetting telomere length, but also of restoring the proliferative lifespan of aged human cells. Dr. West will also describe the plans of BioTime's subsidiary ReCyte Therapeutics, Inc. to commercialize its own proprietary iPS cell technology. The relative advantages of this novel technique over the classic, but potentially genetically unstable, iPS cell methods will be outlined, as will BioTime's progress with reprogrammed young vascular progenitors for the treatment of age-related cardiovascular disease.

This is the fourth French-American Biotech Symposium co-organized by The Office for Science and Technology of the French Embassy in Washington, DC and Eurobiomed, a French biotech cluster dedicated to health sciences, in partnership with the Gladstone Institute for Virology and Immunology.

Dr. West's presentation will be available on BioTime's website at [www.biotimeinc.com](http://www.biotimeinc.com). For more information on ReCyte Therapeutics, please visit our website at [www.recytecorp.com](http://www.recytecorp.com).

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## **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, including a wide array of proprietary ACTCellerate™ cell lines, culture media, and differentiation kits. BioTime's wholly owned subsidiary ES Cell International (ESI) has produced clinical-grade human embryonic stem cell lines that were derived following principles of Good Manufacturing Practice and currently offers them for use in research. 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 an option to clinically develop and commercialize Cell Cure's OpRegen™ 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. ReCyte Therapeutics is developing applications of BioTime's proprietary iPS cell technology to reverse the developmental aging of human cells for cardiovascular and blood cell aging. 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, ReCyte Therapeutics, 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:**

BioTime, Inc.

Judith Segall, 510-521-3390 ext. 301

[jsegall@biotimemail.com](mailto:jsegall@biotimemail.com)