# **UNITED STATES** SECURITIES AND EXCHANGE COMMISSION

**WASHINGTON, DC 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): April 4, 2014

# GTx, Inc.

(Exact Name of Registrant as Specified in Charter)

**Delaware** (State or Other Jurisdiction of Incorporation)

000-50549 (Commission File Number)

62-1715807 (IRS Employer Identification No.)

175 Toyota Plaza 7th Floor Memphis, Tennessee (Address of Principal Executive Offices)

38103 (Zip Code)

Registrant's telephone number, including area code: (901) 523-9700

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

- o Written communication 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.

On April 4, 2014, GTx, Inc. (the "Company") issued a press release announcing that Mitchell S. Steiner, the vice chairman and chief executive officer and a co-founder of the Company, has resigned his employment as chief executive officer of the Company and has resigned as a member of the Board, effective April 3, 2014, in order to pursue other business interests. Marc S. Hanover, a co-founder of the Company and the president and chief operating officer of the Company since the Company's inception in September 1997, has been named interim chief executive officer by the Company's Board of Directors (the "Board") and was elected by the Board to fill Dr. Steiner's remaining term as a Class II director until the annual meeting of stockholders in 2015. The Company intends to file an additional Current Report on Form 8-K within the required time period to provide disclosure under Item 5.02.

A copy of the press release is furnished as Exhibit 99.1 to this Current Report.

#### Item 9.01 Financial Statements and Exhibits.

Exhibits. (d)

**Exhibit No** Description

Press Release issued by GTx, Inc. dated April 4, 2014 99.1

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: April 4, 2014 GTx, Inc.

By: /s/ Henry P. Doggrell

Name: Henry P. Doggrell

Title: Vice President, Chief Legal Officer and Secretary

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### EXHIBIT INDEX

Press Release issued by GTx, Inc. dated April 4, 2014

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#### **GTx Announces Departure of CEO**

- Dr. Mitchell Steiner steps down as CEO and Vice Chairman of the Board -
  - Marc Hanover, GTx's President and COO, named interim CEO -

April 4, 2014—GTx, Inc. (NASDAQ: GTXI) today announced that Mitchell S. Steiner, its vice chairman and chief executive officer (CEO) and a co-founder of the company, is leaving GTx to pursue other business interests. Dr. Steiner has stepped down from his roles as CEO and vice chairman of the Board of Directors at the company effective Thursday April 3, 2014.

Marc S. Hanover, a co-founder of GTx, has been named interim CEO by the Company's Board. He also was elected by the Board to fill Dr. Steiner's remaining term as a Class II director, until the annual meeting of shareholders in 2015. Mr. Hanover has served as president and chief operating officer of GTx since the company's inception in September 1997.

"Since co-founding the company, Mitch has played an instrumental role in helping build GTx to develop novel medicines for cancer patients," said J.R. Hyde, III, chairman of the Board of GTx. "The company remains committed to the implementation of GTx's strategy, which includes pursuing a marketing application in Europe for enobosarm 3mg for the prevention and treatment of muscle wasting in patients with advanced non-small cell lung cancer."

"After more than 15 years as CEO, it's a good time for me to leave my position so that I can spend more time with my family and pursue different opportunities," said Dr. Steiner. "I remain excited by the company's prospects and the potential of GTx's drug candidates to improve the lives of people living with cancer."

#### GTx remains focused on the following:

- · Completing the required Phase I studies and gaining regulatory approval of the company's pediatric investigation plan (PIP) from the European Medicines Agency (EMA) Pediatric Committee in order to submit by the first quarter of 2015 a marketing authorization application (MAA) in the European Union (EU) for enobosarm 3mg for the prevention and treatment of muscle wasting in patients with advanced non-small cell lung cancer (NSCLC)
- · Completing in the 3<sup>rd</sup> quarter of this year, the company's ongoing Phase 2 clinical study of enobosarm 9mg to treat androgen receptor and estrogen receptor positive advanced breast cancer
- · Meeting with the United States Food and Drug Administration (FDA) to determine an appropriate regulatory path forward in the U.S. for enobosarm 3 mg for the prevention and treatment of cancer cachexia in NSCLC, and
- · Completing the company's ongoing Phase 2 clinical study of GTx-758 as a secondary hormonal treatment for men with castration-resistant prostate cancer.

### About GTx

GTx, Inc., headquartered in Memphis, Tenn., is a biopharmaceutical company dedicated to the discovery, development, and commercialization of small molecules for the treatment of cancer, cancer supportive care, including prevention and treatment of cancer-related muscle wasting, and other serious medical conditions.

# Forward-Looking Information is Subject to Risk and Uncertainty

This press release contains forward-looking statements based upon GTx's current expectations. Forward-looking statements involve risks and uncertainties, and include, but are not limited to, statements relating to the anticipated completion of GTx's clinical trials for enobosarm (GTx-024) and GTx-758 (Capesaris®), the potential approval of GTx's PIP from the EMA, the potential submission of a MAA to the EMA for enobosarm 3mg and the timing thereof, and GTx's plans to meet with the FDA and a potential regulatory path forward for enobosarm 3mg in the United States. GTx's actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of these risks and uncertainties, which include, without limitation, the risks (i) that GTx may be unable to successfully complete or develop, in a timely manner or at all, the Phase 1 studies and PIP necessary to enable the submission of the planned MAA; (ii) related to the uncertain and time-consuming regulatory approval process, including the risk that GTx may not be able to obtain required regulatory approvals to commercialize enobosarm 3mg in a timely manner or at all; (iii) that clinical trials being conducted by GTx may not be completed on schedule, or at all, or may otherwise be suspended or terminated or (iv) that GTx could utilize its available cash resources sooner than it currently expects and may be unable to raise additional capital, which would force GTx to delay, reduce or eliminate its product candidate development programs. You should not place undue reliance on these forward-looking statements, which apply only as of the date of this press release. GTx's annual report on Form 10-K filed with the Securities and Exchange Commission on March 12, 2014 contains under the heading, "Risk Factors", a more comprehensive description of these and other risks to which GTx is subject. GTx expressly disclaims any obligation or undertaking to release publicly any updates or revisions to any forward-looking stat

Source: GTx, Inc.

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