UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
under the Securities Exchange Act of 1934

February 14, 2014

Commission File Number 001-35203

THERATECHNOLOGIES INC.

(Translation of registrant's name into English)

2310 Alfred-Nobel Boulevard
Montréal, Québec, Canada
H4S 2B4
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F:

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Yes ☐ No ☒

Note: Regulation S-T Rule 101(b)(1) only permits the submission in paper of a Form 6-K if submitted solely to provide an attached annual report to security holders.

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Yes ☐ No ☒

Note: Regulation S-T Rule 101(b)(7) only permits the submission in paper of a Form 6-K if submitted to furnish a report or other document that the registrant foreign private issuer must furnish and make public under the laws of the jurisdiction in which the registrant is incorporated, domiciled or legally organized (the registrant's "home country"), or under the rules of the home country exchange on which the registrant's securities are traded, as long as the report or other document is not a press release, is not required to be and has not been distributed to the registrant's security holders, and, if discussing a material event, has already been the subject of a Form 6-K submission or other Commission filing on EDGAR.

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes ☐ No ☒

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): 82-_____.

Exhibit	Description
99.1	Press Release dated February 14, 2014

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, thereunto duly authorized.

THERATECHNOLOGIES INC.

By: /s/ Jocelyn Lafond
Name: Jocelyn Lafond
Title: Vice President, Legal Affairs

Date: February 14, 2014

Short Supply of *EGRIFTA*® Announced

Montreal, Canada – February 14, 2014 – Theratechnologies Inc. (TSX: TH) announced today that it expects current inventory of *EGRIFTA*® (tesamorelin for injection) to be depleted in the coming weeks due to a combination of manufacturing delays and issues observed during the production of new batches of *EGRIFTA*®. The depletion of the inventory will result in a shortage of *EGRIFTA*® and an eventual stock-out. EMD Serono, Inc. has notified the United States Food and Drug Administration about this product shortage.

The Company also announced that it has made the decision to temporarily cease the manufacture of *EGRIFTA*®. At this time, the Company is unable to determine a timeline to resume the manufacture and delivery of *EGRIFTA*® and the Company is currently investigating the causes of the issues observed. The Company will update the market when new information is available.

The transaction with EMD Serono, Inc. previously announced on December 13, 2013 relating to the Company regaining all rights to *EGRIFTA*® in the United States is still expected to close during the Company's second quarter of its current fiscal year despite the decision made by the Company to temporarily halt the manufacture of *EGRIFTA*®.

About Theratechnologies

Theratechnologies (TSX: TH) is a biopharmaceutical company which specializes in the development and the commercialization of innovative therapeutic peptide products. Further information about Theratechnologies is available on the Company's website at www.theratech.com, on SEDAR at www.sedar.com and on the Securities and Exchange Commission's website at www.sec.gov.

Forward-Looking Information

This press release contains certain statements that are considered "forward-looking information" within the meaning of applicable securities legislation, which statements may contain such words as "may", "would", "could", "will", "intend", "plan", "anticipate", "believe", "estimate", "expect" and similar expressions. This forward-looking information includes, but is not limited to, information regarding the timing on the depletion of the *EGRIFTA*® inventory and the closing date of the transaction with EMD Serono, Inc. announced on December 13, 2013.

Forward-looking information is based upon a number of assumptions and is subject to a number of risks and uncertainties, many of which are beyond Theratechnologies' control that could cause actual results to differ materially from those that are disclosed in or implied by such forward-looking information. These assumptions include, but are not limited to, the fact that sales of *EGRIFTA*® in the United States will not slow down and that the FDA will not issue any order or decision having the effect of suspending the commercialization of *EGRIFTA*® in the United States.

These risks and uncertainties include, but are not limited to, the risk that the drug shortage occurs earlier than anticipated if demand for *EGRIFTA*® increases, that the Company is unable to rapidly identify the causes of the issues and is unable to rapidly resume the manufacture of *EGRIFTA*®, all of which would result in a long product shortage and a loss of goodwill for *EGRIFTA*®, that the Company is required to develop and implement corrective measures or a new manufacturing process for *EGRIFTA*®, all of which will take time and cost money while no revenue from sales of *EGRIFTA*® will be recorded and, therefore, our financial condition and operating results will be materially adversely affected, that corrective measures, if need be, be subject to the FDA's approval and that we do not obtain such approval, that we may be unable to meet our ongoing and future obligations if the stock-out lasts for a long period of time and we do not control our expenses, or that an order preventing the commercialization of *EGRIFTA*® or the closing of the transaction with EMD Serono, Inc. is issued by the FDA or any other U.S. regulatory authorities.

We refer potential investors to the "Risk Factors" section of our Annual Report on Form 20-F dated February 26, 2013 available at www.sedar.com, www.sec.gov and www.theratech.com. The reader is cautioned to consider these and other risks and uncertainties carefully and not to put undue reliance on forward-looking statements. Forward-looking statements reflect current expectations regarding future events and speak only as of the date of this press release and represent our expectations as of that date.

We undertake no obligation to update or revise the information contained in this press release, whether as a result of new information, future events or circumstances or otherwise, except as may be required by applicable law.

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