

TherapeuticsMD, Inc.  
Form 8-K  
October 29, 2018

**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): October 29, 2018

**TherapeuticsMD, Inc.**

(Exact Name of Registrant as Specified in its Charter)

**Nevada**  
(State or Other

**001-00100**  
(Commission File Number)

**87-0233535**  
(IRS Employer  
Identification No.)

Jurisdiction of Incorporation)

6800 Broken Sound Parkway NW, Third Floor

Boca Raton, FL 33487

(Address of Principal Executive Office) (Zip Code)

Registrant's telephone number, including area code: (561) 961-1900

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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Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§ 230-405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

**Item 7.01 Regulation FD Disclosure.**

On October 29, 2018, TherapeuticsMD, Inc., a Nevada corporation (the “Company”), issued a press release announcing that the United States Food and Drug Administration (FDA) has approved BIJUVA™ (estradiol and progesterone) capsules, 1 mg/100 mg, the first and only FDA-approved bio-identical hormone therapy combination of estradiol and progesterone in a single, oral capsule for the treatment of moderate to severe vasomotor symptoms (commonly known as hot flashes or flushes) due to menopause in women with a uterus. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated in this Item 7.01 by reference.

The Company intends to host a conference call to discuss the approval of BIJUVA™ on Monday, October 29, 2018, at 8:30 a.m. EDT, and is furnishing as Exhibit 99.2 to this Current Report on Form 8-K a presentation which will be used, in whole or in part, and subject to modification, on October 29, 2018 and at subsequent meetings with investors or analysts.

The information included in this Item 7.01 and in Exhibits 99.1 and 99.2 shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) *Exhibits*

Exhibit

Number Description

- |             |  |
|-------------|--|
| <u>99.1</u> | Press Release from TherapeuticsMD, Inc., dated October 29, 2018. |
| <u>99.2</u> | TherapeuticsMD, Inc. presentation dated October 29, 2018.        |

**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.

Date: October 29, 2018 THERAPEUTICSMD, INC.

By: /s/ Daniel A. Cartwright  
Name: Daniel A. Cartwright  
Title: Chief Financial Officer