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

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**FORM 8-K**

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**CURRENT REPORT**  
Pursuant to Section 13 or 15(d)  
of the Securities Exchange Act of 1934

Date of report (Date of earliest reported) **October 30, 2019**

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**BRICKELL BIOTECH, INC.**

(Exact name of Registrant as specified in its charter)

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

**000-21088**  
(Commission File  
Number)

**93-0948554**  
(IRS Employer  
Identification No.)

**5777 Central Avenue**  
**Suite 102**  
**Boulder, CO 80301**  
(Address of Principal Executive Offices)

Registrant's telephone number, including area code: **(720) 505-4755**

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):

- Written communication 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))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common stock, par value \$0.01 per share	BBI	The Nasdaq Capital Market

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.

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## Item 8.01. Other Events

On October 25, 2019, NovaQuest Co-Investment Fund X, L.P. (“NovaQuest”) provided written notice to Brickell Biotech, Inc. (the “Company”) of its determination that a “Material Adverse Event” (as defined in Section 1.1 of the Funding Agreement previously entered into between NovaQuest and the Company (the “Funding Agreement”)) occurred as a result of the matters described in the Form 8-K filed by the Company on October 24, 2019 (the “October 24 Form 8-K”). As a result, NovaQuest exercised its right under Section 3.3(e) of the Funding Agreement to suspend further Development Payments (as defined in Section 3.1(a) of the Funding Agreement). Pursuant to the Funding Agreement, NovaQuest is obligated to resume Development Payments if the Material Adverse Event is resolved or cured by the Company to NovaQuest’s reasonable satisfaction by October 25, 2020. If the Material Adverse Event is not resolved or cured to NovaQuest’s reasonable satisfaction by such date, then NovaQuest may, in its sole discretion, terminate any future payment obligation under the Funding Agreement and Brickell may be obligated to make certain payments to NovaQuest.

Additionally, as a result of the matters described in the October 24 Form 8-K, the timeline for the Company’s Phase 3 clinical trials in subjects with primary axillary hyperhidrosis in the United States may be impacted. We intend to provide an update on the timeline when we have further clarity.

Today the Company initiated an arbitration proceeding pursuant to Article 9 of the License Agreement previously entered into between Bodor Laboratories, Inc. (“Bodor”), Nicholas S. Bodor and the Company (the “License Agreement”) with the American Arbitration Association (“AAA”) in Florida against Bodor and Nicholas S. Bodor. This arbitration seeks a declaratory judgment that the purported termination of the License Agreement by Bodor and Nicholas S. Bodor was invalid and unenforceable and asserts (i) a claim for breach of the License Agreement against Bodor and Nicholas S. Bodor, in his individual capacity, and (ii) a claim against Bodor and Nicholas S. Bodor for tortious interference with the Company’s business relations. We have requested expedited treatment of the arbitration proceeding and concurrent mandatory mediation under the AAA rules. The Company concurrently filed today with the United States District Court for the Southern District of Florida a motion to dismiss the complaint brought against the Company by Bodor and Nicholas S. Bodor described in the October 24 Form 8-K.

A copy of the Company’s press release announcing the aforementioned events is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

<b>Exhibit No.</b>	<b>Description of Exhibit</b>
<a href="#">99.1</a>	Press Release dated October 30, 2019

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## SIGNATURES

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: October 30, 2019

Brickell Biotech, Inc.

By: /s/ Robert B. Brown  
Name: Robert B. Brown  
Title: Chief Executive Officer

October 30, 2019

## **Brickell Biotech Provides Update on Bodor Labs Complaint and Funding Agreement with NovaQuest**

*NovaQuest temporarily suspended R&D funding payments*

*Brickell filed a motion to dismiss the complaint filed by Bodor Labs and Nicholas S. Bodor in federal court*

*Brickell initiated arbitration proceedings against Bodor Labs and Nicholas S. Bodor including a claim for tortious interference*

BOULDER, Colo., Oct. 30, 2019 (GLOBE NEWSWIRE) -- Brickell Biotech, Inc. ("Brickell") (NASDAQ: BBI), a clinical-stage pharmaceutical company, today announced that it has initiated an arbitration proceeding pursuant to Article 9 of the License Agreement previously entered into between Bodor Laboratories, Inc. ("Bodor"), Nicholas S. Bodor and Brickell with the American Arbitration Association ("AAA") in Florida against Bodor and Nicholas S. Bodor. This arbitration seeks a declaratory judgment that the purported termination of the License Agreement by Bodor and Nicholas S. Bodor was invalid and unenforceable and asserts (i) a claim for breach of the License Agreement against Bodor and Nicholas S. Bodor, in his individual capacity, and (ii) a claim against Bodor and Nicholas S. Bodor for tortious interference with Brickell's business relations. Brickell has requested expedited treatment of the arbitration proceeding and concurrent mandatory mediation under the AAA rules. Brickell concurrently filed today with the United States District Court for the Southern District of Florida a motion to dismiss the complaint brought against Brickell by Bodor and Nicholas S. Bodor on October 24, 2019.

On October 25, 2019, NovaQuest Co-Investment Fund X, L.P. ("NovaQuest") provided written notice to Brickell of its determination that a material adverse event occurred as a result of the matter described above. As a result, NovaQuest exercised its right to suspend further development payments under the Funding Agreement. NovaQuest is obligated to resume development payments if the material adverse event is resolved or cured by Brickell to NovaQuest's reasonable satisfaction by October 25, 2020. If the material adverse event is not resolved or cured to NovaQuest's reasonable satisfaction by such date, then NovaQuest may, in its sole discretion, terminate any future payment obligation under the Funding Agreement and Brickell may be obligated to make certain payments to NovaQuest.

Additionally, as a result of the matters described above, the timeline for Brickell's Phase 3 clinical trials in subjects with primary axillary hyperhidrosis in the United States may be impacted. Brickell intends to provide an update on the timeline when there is further clarity.

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## **About Brickell**

Brickell is a clinical-stage pharmaceutical company focused on developing innovative and differentiated prescription therapeutics for the treatment of debilitating skin diseases. Brickell's pipeline consists of potential novel therapeutics for hyperhidrosis, cutaneous T-cell lymphoma, psoriasis, and other prevalent dermatological conditions. Brickell's executive management team and board of directors bring extensive experience in product development and global commercialization, having served in leadership roles at large global pharmaceutical companies and biotechs that have developed and/or launched successful products, including several that were first-in-class and/or achieved iconic status, such as Cialis<sup>®</sup>, Taltz<sup>®</sup>, Gemzar<sup>®</sup>, Prozac<sup>®</sup>, Cymbalta<sup>®</sup> and Juvederm<sup>®</sup>. Brickell's strategy is to leverage this experience to in-license, acquire, develop and commercialize innovative products that Brickell believes can be successful in the currently underserved dermatology global marketplace. For more information, visit [www.brickellbio.com](http://www.brickellbio.com).

## **Cautionary Note Regarding Forward-Looking Statements**

Any statements made in this press release relating to future financial, business and/or research and clinical performance, conditions, plans, prospects, trends, or strategies and other such matters, including without limitation, the anticipated timing, scope, design and/or results of future clinical trials and prospects for commercializing any of Brickell's product candidates are forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. In addition, when or if used in this press release, the words "may," "could," "should," "anticipate," "believe," "estimate," "expect," "intend," "plan," "predict" and similar expressions and their variants, as they relate to Brickell, may identify forward-looking statements. Brickell cautions that these forward-looking statements are subject to numerous assumptions, risks, and uncertainties, which change over time, often quickly and in unanticipated ways.

Further information on the factors and risks that could cause actual results to differ from any forward-looking statements are contained in Brickell's filings with the United States Securities and Exchange Commission (SEC), which are available at [www.sec.gov](http://www.sec.gov) (or at [www.brickellbio.com](http://www.brickellbio.com)). The forward-looking statements represent the estimates of Brickell as of the date hereof only, and Brickell specifically disclaims any duty or obligation to update forward-looking statements.

## **Brickell Investor / Media Contact:**

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