

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: April 25, 2017  
(Date of earliest event reported)

**CELLECTAR BIOSCIENCES, INC.**  
(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**1-36598**  
(Commission  
File Number)

**04-3321804**  
(IRS Employer  
Identification Number)

**3301 Agriculture Drive**  
**Madison, WI 53716**  
(Address of principal executive offices)

**(608) 441-8120**  
(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 (see General Instruction A.2. below):

- 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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**ITEM 7.01 REGULATION FD DISCLOSURE**

On April 25, 2017, we issued a press release announcing the United States Patent and Trademark Office granted us a method of use patent for CLR 1501, CLR 1502 and an additional CLR 1401-boron-dipyrromethene analog for the detection of multiple cancer types. All of these compounds utilize our proprietary phospholipid drug conjugate (PDC) delivery platform. A copy of the press release is furnished as Exhibit 99.1 and is incorporated by reference herein.

**ITEM 9.01 FINANCIAL STATEMENTS AND EXHIBITS**

(d) Exhibits

<u>Number</u>	<u>Title</u>
99.1	Press release dated April 25, 2017, entitled "Collectar Biosciences Receives Additional U.S. Patents for PDC Optical Agents in the Detection of Multiple Cancers"

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**SIGNATURE**

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.

Dated: April 25, 2017

**CELLECTAR BIOSCIENCES, INC.**

By: /s/ Chad J. Kolean

Name: Chad J. Kolean

Title: Vice President and Chief Financial Officer

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**Collectar Biosciences Receives Additional U.S. Patents for PDC Optical Agents in the Detection of Multiple Cancers**

Madison, Wisc. (April 25, 2017) -- Collectar Biosciences, Inc. (Nasdaq: CLRB) (the “company”), an oncology-focused, clinical stage biotechnology company, today announces the United States Patent and Trademark Office has granted a method of use patent for CLR 1501, CLR 1502 and an additional CLR 1401-boron-dipyrrromethene analog for the detection of multiple cancer types. All of these compounds utilize Collectar’s proprietary phospholipid drug conjugate (PDC) delivery platform.

The recently issued patent, 9,616,140, outlines the method of use of these fluorophore compounds to detect a variety of solid tumors in patients, including melanomas, colorectal adenocarcinoma, uterine carcinoma, pancreatic carcinoma, ovarian adenocarcinoma, glioblastoma, clear cell carcinoma, and prostate adenocarcinoma. The current patent provides intellectual property protection through May 11, 2029.

“We continue to successfully execute our plan to expand the company’s intellectual property portfolio to protect and enhance the value of our PDC pipeline assets, both in diagnostic and therapeutic applications,” said Jim Caruso, president and CEO of Collectar. “While our focus continues to be the development of our therapeutic assets, specifically CLR 131, for the treatment of multiple myeloma and other hematologic malignancies, our platform assets offer significant additional opportunity in a variety of clinical applications.”

**About Phospholipid Drug Conjugates (PDCs)**

Collectar’s product candidates are built upon its patented cancer cell-targeting delivery and retention platform of optimized phospholipid ether-drug conjugates (PDCs). The company deliberately designed its phospholipid ether (PLE) carrier platform to be coupled with a variety of payloads to facilitate both therapeutic and diagnostic applications. The basis for selective tumor targeting of our PDC compounds lies in the differences between the plasma membranes of cancer cells compared to those of normal cells. Cancer cell membranes are highly enriched in lipid rafts, which are glycolipoprotein microdomains of the plasma membrane of cells that contain high concentrations of cholesterol and sphingolipids, and serve to organize cell surface and intracellular signaling molecules. PDCs have been tested in more than 80 different xenograft models of cancer.

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## **About Collectar Biosciences, Inc.**

Collectar Biosciences is developing phospholipid drug conjugates (PDCs) designed to provide cancer targeted delivery of diverse oncologic payloads to a broad range of cancers and cancer stem cells. Collectar's PDC platform is based on the company's proprietary phospholipid ether analogs. These novel small-molecules have demonstrated highly selective uptake and retention in a broad range of cancers.

Collectar's PDC pipeline includes product candidates for cancer therapy and cancer diagnostic imaging. The company's lead therapeutic PDC, CLR 131, utilizes iodine-131, a cytotoxic radioisotope, as its payload. CLR 131 is currently being evaluated under an orphan drug designated Phase I clinical study in patients with relapsed or refractory multiple myeloma. In addition, the company has initiated a Phase II clinical study to assess efficacy in a range of B-cell malignancies. The company is also developing PDCs for targeted delivery of chemotherapeutics such as paclitaxel (CLR 1602-PTX), a preclinical stage product candidate, and plans to expand its PDC chemotherapeutic pipeline through both in-house and collaborative R&D efforts. For more information please visit [www.collectar.com](http://www.collectar.com).

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This news release contains forward-looking statements. You can identify these statements by our use of words such as "may," "expect," "believe," "anticipate," "intend," "could," "estimate," "continue," "plans," or their negatives or cognates. These statements are only estimates and predictions and are subject to known and unknown risks and uncertainties that may cause actual future experience and results to differ materially from the statements made. These statements are based on our current beliefs and expectations as to such future outcomes. Drug discovery and development involve a high degree of risk. Factors that might cause such a material difference include, among others, uncertainties related to the ability to raise additional capital, uncertainties related to the ability to attract and retain partners for our technologies, the identification of lead compounds, the successful preclinical development thereof, the completion of clinical trials, the FDA review process and other government regulation, our pharmaceutical collaborators' ability to successfully develop and commercialize drug candidates, competition from other pharmaceutical companies, product pricing and third-party reimbursement. A complete description of risks and uncertainties related to our business is contained in our periodic reports filed with the Securities and Exchange Commission including our Form 10-K for the year ended December 31, 2016. These forward-looking statements are made only as of the date hereof, and we disclaim any obligation to update any such forward-looking statements.

**CONTACT:**

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