

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: May 20, 2015  
(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))
-

**ITEM 7.01 REGULATION FD DISCLOSURE**

On May 26, 2015, we issued a press release announcing our intent to adjourn our Annual Meeting of Stockholders from May 28, 2015 to June 9, 2015. A copy of the press release is furnished as Exhibit 99.1 and is incorporated by reference herein.

On May 21, 2015, we issued a press release announcing our first quarter 2015 results and providing an update on our clinical programs. The release further announced that management would host a conference call and webcast on May 21, 2015, beginning at 5:00 P.M. EDT. A copy of the press release is furnished as Exhibit 99.2 and is incorporated by reference herein.

On May 20, 2015, we issued a press release announcing our intent to host a conference call to discuss first quarter 2015 results and provide an update on our clinical programs. A copy of the press release is furnished as Exhibit 99.3 and is incorporated by reference herein. Additionally, we are furnishing a copy of our May 11, 2015 press release which announced the cancellation of our previously scheduled first quarter financial and conference call as Exhibit 99.4.

**ITEM 9.01 FINANCIAL STATEMENTS AND EXHIBITS**

(d) Exhibits

<u>Number</u>	<u>Title</u>
99.1	Press release dated May 26, 2015, entitled "Collectar Biosciences Announces Intent to Adjourn Annual Meeting Of Stockholders"
99.2	Press release dated May 21, 2015, entitled "Collectar Biosciences Reports First Quarter 2015 Financial Results and Provides Update on Clinical Programs"
99.3	Press release dated May 20, 2015, entitled "Collectar Biosciences to Host Conference Call on May 21st to Discuss First Quarter 2015 Results and Provide Update on Development Programs"
99.4	Press release dated May 11, 2015, entitled "Collectar Biosciences Reschedules Release of First Quarter Financials and Quarterly Conference Call"

---

**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: May 27, 2015

**CELLECTAR BIOSCIENCES, INC.**

By: /s/ Chad J. Kolean

Name: Chad J. Kolean

Title: Vice President and Chief Financial Officer

---

## EXHIBIT INDEX

Number	Title
99.1	Press release dated May 26, 2015, entitled “Collectar Biosciences Announces Intent to Adjourn Annual Meeting Of Stockholders”
99.2	Press release dated May 21, 2015, entitled “Collectar Biosciences Reports First Quarter 2015 Financial Results and Provides Update on Clinical Programs”
99.3	Press release dated May 20, 2015, entitled “Collectar Biosciences to Host Conference Call on May 21st to Discuss First Quarter 2015 Results and Provide Update on Development Programs”
99.4	Press release dated May 11, 2015, entitled “Collectar Biosciences Reschedules Release of First Quarter Financials and Quarterly Conference Call”

---



**Cellecstar Biosciences Announces Intent to Adjourn Annual Meeting Of Stockholders**

*Meeting to be reconvened on June 9, 2015 at 10:00 AM Central Time*

*Board of Directors Wishes to Provide Stockholders with Sufficient Time to Review Amendments to the Company's Annual Report*

**MADISON, Wis., May 26, 2015**, – Cellecstar Biosciences, Inc. (NASDAQ: CLRB), a clinical stage biopharmaceutical company developing innovative agents for the detection and treatment of cancer, announced today that it intends to call to order and then immediately adjourn its annual meeting of stockholders to be held at its Corporate headquarters on Thursday, May 28, 2015, at 10:00 a.m., local time. The adjournment of the annual meeting of stockholders will be until 10:00 a.m., local time on Tuesday, June 9, 2015 at Cellecstar's headquarters, located at 3301 Agriculture Drive, Madison, Wisconsin. Cellecstar's Board of Directors is taking this action to allow stockholders sufficient time to review its amended annual report on Form 10-K/A, including the Company's restated financial results (available at <http://www.cellecstar.com>).

Stockholders of record may submit their votes for matters to be considered at the annual meeting until the polls are formally closed. Stockholders who have already voted in accordance with the instructions contained in the proxy statement and related materials do not need to submit new proxy cards or give new voting instructions unless they wish to change their votes. Stockholders who have not yet voted can still use the proxy cards and voting instruction forms previously provided to them. Cellecstar's Board of Directors encourages all stockholders to review the proxy statement and Annual Report on Form 10-K/A carefully before voting.

---

For stockholders that made arrangements to attend the annual meeting on May 28<sup>th</sup>, Dr. Stephen Hill, chairman of the board of directors, and Dr. Simon Pedder, president and chief executive officer, plan to make themselves available for an informal gathering following the adjournment of the formal meeting on May 28th.

**Restatement of Prior Financial Statements:**

As disclosed in a Form 8-K filed by the Company on May 18, 2015, the Audit Committee of the Board of Directors of Collectar, in connection with an internal review conducted by Collectar's management, concluded that, because of a misapplication of the accounting guidance related to certain of the Company's warrants, Collectar's previously issued unaudited condensed consolidated financial statements for the three and nine months ended September 30, 2014 and audited consolidated financial statements for the twelve months ended December 31, 2014 should no longer be relied upon. On May 20, 2015, Collectar filed an amended Form 10-Q/A for the third quarter of 2014 and an amended 2014 Form 10-K/A for the year ended December 31, 2014 reflecting the restatements which result in non-cash, non-operating financial statement corrections and have no impact on the Collectar's current or previously reported cash position, operating expenses or total operating, investing or financing cash flows.

On August 20, 2014, in addition to other securities, Collectar issued 3,833,333 warrants to purchase shares of our common stock at an exercise price of \$4.68 per share as part of an underwritten offering. In connection with the election to participate in this offering by the holders of debentures representing \$4,000,000 principal amount and related accrued interest of \$172,435, Collectar issued an additional 1,109,690 warrants. These warrants contain a cash settlement feature in the event there is no current prospectus to support the issuance of stock and warrant holder wishing to exercise the warrant, requests gross settlement rather than net settlement via cashless exercise.

The Audit Committee, together with management, determined that the financial statements subsequent to this issuance should be restated to reflect the warrants issued in August 2014 as liabilities, with subsequent changes in their estimated fair value recorded as non-cash income or expense in each affected period.

---

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

Collectar Biosciences is developing agents to detect, treat and monitor a broad spectrum of cancers. Using a novel phospholipid ether analog (PLE) platform technology as a targeted delivery and retention vehicle, Collectar's compounds are designed to be selectively taken up and retained in cancer cells including cancer stem cells. With the ability to attach both imaging and therapeutic agents to its proprietary delivery platform, Collectar has developed a portfolio of Phase I and Phase II product candidates engineered to leverage the unique characteristics of cancer cells to "find, treat and follow" malignancies in a highly selective way. For additional information please visit [www.collectar.com](http://www.collectar.com).

## **INVESTOR CONTACT**

Kate McNeil, Vice President of IR, PR & Corporate Communications

Collectar Biosciences, Inc.

Phone: (347) 204-4226

Email: [kmcneil@collectar.com](mailto:kmcneil@collectar.com)

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/A for the year ended December 31, 2014. These forward-looking statements are made only as of the date hereof, and we disclaim any obligation to update any such forward-looking statements.

---



## Cellecstar Biosciences Reports First Quarter 2015 Financial Results and Provides Update on Clinical Programs

*Management to Host Conference Call and Webcast Today at 5:00 PM EDT*

**MADISON, Wis., May 21, 2015**, – Cellecstar Biosciences, Inc. (NASDAQ: CLRB), a clinical stage biopharmaceutical company developing innovative agents for the detection and treatment of cancer, is providing an update on its development programs and financial results for the first quarter 2015.

### **Clinical Development Program Updates:**

#### ***I-124-CLR1404***

- After activating five large new centers this year, Cellecstar has seen an increase in patient screening and anticipates a significant increase in enrollment in its Phase II trial of I-124-CLR1404 in glioblastoma. However, enrollment remains slower than expected and the company is evaluating potential strategies to leverage existing data from investigator-sponsored studies to modify its ongoing Phase II trial of I-124-CLR1404 in glioblastoma and reduce the time to successful trial completion.
- Progress in investigator-sponsored clinical trials of I-124-CLR1404, including imaging of 30 patients with various brain cancers, should provide meaningful evidence for the optimal dose and imaging time-point for the use of I-124-CLR1404 in glioblastoma, the primary objective of the on-going company-sponsored Phase II trial in glioblastoma.

#### ***I-131-CLR1404***

- In April 2015, Cellecstar initiated patient dosing in a proof-of-concept trial of I-131-CLR1404 in patients with relapsed or refractory multiple myeloma, an indication for which I-131-CLR1404 previously received orphan drug designation from the U.S. Food and Drug Administration. Based on data from the company's Phase Ib trial, Cellecstar anticipates that evidence of clinical activity will be assessable relatively early in the dose escalation process.
-



- Collectar continues to expect data from this program to be available by year-end 2015.

#### ***CLR1502***

- In February 2015, a publication featured on the cover of *Neurosurgery*, Official Journal of the Congress of Neurological Surgeons - the largest neurosurgical society in the world, demonstrated that Collectar's fluorescent, cancer-selective agents successfully provide visualization of glioma cells with high fidelity, and suggest their practical and promising potential to optimize tumor surgery.
- During the first quarter of 2015, Collectar submitted an investigational new drug (IND) application to the FDA to allow for initiation of a Phase I proof-of-concept trial of CLR1502 in breast cancer patients undergoing lumpectomy. The trial is intended to establish the safety and tolerability of CLR1502 while demonstrating its utility in the real-time identification of malignant tissue.
- Collectar is currently working with the FDA to determine if CLR1502 should be evaluated as an imaging agent through the Center for Drug Evaluation and Research (CDER) or as a combination product along with an imaging system (light source) through the Center for Devices and Radiological Health (CDRH).
- Collectar is working closely with the FDA to resolve this matter and expects to initiate its planned proof-of-concept study in the second half of 2015.

“Having recently initiated our proof-of-concept therapeutic trial in multiple myeloma, Collectar now has two promising product candidates in registration-enabling clinical trials, both of which leverage the company's proprietary broad-spectrum cancer targeting and retention technology and seek to address significant unmet medical needs in cancer care,” commented Dr. Simon Pedder, Collectar's president and chief executive officer. “We look forward to advancing both programs, reporting preliminary data from each, and initiating our fluorescence-guided surgery program with our tumor illuminating agent, CLR1502.”

---

**Financial Results for the Quarter Ended March 31, 2015:**

Cellectar reported a net loss for the quarter ended March 31, 2014 of \$2.3 million or (\$0.30) per share versus a net loss of \$2.9 million or (\$1.03) per share for the comparable period in 2014.

Research and development (R&D) expenses for the quarter ended March 31, 2015 were \$1.6 million, compared to \$1.7 million for the first quarter of 2014. The decrease in first quarter 2015 R&D expense reflects a decrease in costs associated with supporting investigator-sponsored clinical studies partially offset by increases in personnel related costs.

Cellectar's general and administrative (G&A) expenses for first quarter 2015 totaled \$0.9 million reflecting a 13% decrease from the \$1.1 reported for the comparable prior year period. The decrease reflects a reduction in consulting charges and legal fees, partially offset by a slight increase in travel-related activities.

Cellectar ended the quarter with \$7.0 million in cash and cash equivalents compared to \$9.4 million in cash and cash equivalents at December 31, 2014. Cellectar anticipates that available cash and cash equivalents should fund the company's planned operations into the fourth quarter 2015 and that additional capital will be required to complete all ongoing and planned clinical and preclinical trials of its product candidates.

**Restatement of Prior Financial Statements:**

As disclosed in a Form 8-K filed by the Company on May 18, 2015, the Audit Committee of the Board of Directors of Cellectar, in connection with an internal review conducted by Cellectar's management, concluded that, because of a misapplication of the accounting guidance related to certain of the Company's warrants, Cellectar's previously issued unaudited condensed consolidated financial statements for the three and nine months ended September 30, 2014 and audited consolidated financial statements for the twelve months ended December 31, 2014 should no longer be relied upon. Cellectar has filed an amended Form 10-Q/A for the third quarter of 2014 and an amended 2014 Form 10-K/A for the year ended December 31, 2014 reflecting the restatements which result in non-cash, non-operating financial statement corrections and have no impact on the Cellectar's current or previously reported cash position, operating expenses or total operating, investing or financing cash flows.

---

On August 20, 2014, in addition to other securities, Collectar issued 3,833,333 warrants to purchase shares of our common stock at an exercise price of \$4.68 per share as part of an underwritten offering. In connection with the election to participate in this offering by the holders of debentures representing \$4,000,000 principal amount and related accrued interest of \$172,435, Collectar issued an additional 1,109,690 warrants. These warrants contain a cash settlement feature in the event there is no current prospectus to support the issuance of stock and warrant holder wishing to exercise the warrant, requests gross settlement rather than net settlement via cashless exercise.

The Audit Committee, together with management, determined that the financial statements subsequent to this issuance should be restated to reflect the warrants issued in August 2014 as liabilities, with subsequent changes in their estimated fair value recorded as non-cash income or expense in each affected period.

**Conference Call and Webcast Today at 5:00 PM EDT:**

Interested investors may participate in the conference call by dialing 888-646-8293 (domestic) or 973-453-3065 (international). Participants may also access both the live and archived webcast of the conference call on the investor relations section of Collectar's web site, [www.collectar.com](http://www.collectar.com).

**About Collectar Biosciences, Inc.**

Collectar Biosciences is developing agents to detect, treat and monitor a broad spectrum of cancers. Using a novel phospholipid ether analog (PLE) platform technology as a targeted delivery and retention vehicle, Collectar's compounds are designed to be selectively taken up and retained in cancer cells including cancer stem cells. With the ability to attach both imaging and therapeutic agents to its proprietary delivery platform, Collectar has developed a portfolio of Phase I and Phase II product candidates engineered to leverage the unique characteristics of cancer cells to "find, treat and follow" malignancies in a highly selective way. For additional information please visit [www.collectar.com](http://www.collectar.com).

---

## **INVESTOR CONTACT**

Kate McNeil, Vice President of IR, PR & Corporate Communications  
Cellecstar Biosciences, Inc.  
Phone: (347) 204-4226  
Email: [kmcneil@cellectar.com](mailto:kmcneil@cellectar.com)

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/A for the year ended December 31, 2014. These forward-looking statements are made only as of the date hereof, and we disclaim any obligation to update any such forward-looking statements.

---



**Cellecstar Biosciences to Host Conference Call on May 21st to Discuss First Quarter 2015 Results and Provide Update on Development Programs**

**MADISON, Wis., May 20, 2015**, – Cellecstar Biosciences, Inc. (NASDAQ: CLRB), a clinical stage biopharmaceutical company developing innovative agents for the detection and treatment of cancer, announced that management will host a conference call and live webcast to discuss first quarter 2015 financial results and provide an update on each of its development programs on Tuesday, May 21st at 5:00 PM EDT.

**Event Details:**

Interested investors may participate in the conference call by dialing 888-646-8293 (domestic) or 973-453-3065 (international). Participants may also access both the live and archived webcast of the conference call on the investor relations section of Cellecstar's web site, [www.cellecstar.com](http://www.cellecstar.com).

**About Cellecstar Biosciences, Inc.**

Cellecstar Biosciences is developing agents to detect, treat and monitor a broad spectrum of cancers. Using a novel phospholipid ether analog (PLE) platform technology as a targeted delivery and retention vehicle, Cellecstar's compounds are designed to be selectively taken up and retained in cancer cells including cancer stem cells. With the ability to attach both imaging and therapeutic agents to its proprietary delivery platform, Cellecstar has developed a portfolio of product candidates engineered to leverage the unique characteristics of cancer cells to "find, treat and follow" malignancies in a highly selective way. I-124-CLR1404 is a small-molecule, broad-spectrum, cancer-targeted PET imaging agent currently being evaluated in a Phase II glioblastoma imaging trial. Additionally, multiple investigator-sponsored Phase I/II clinical trials are ongoing across 11 solid tumor indications. I-131-CLR1404 is a small-molecule, broad-spectrum, cancer-targeted molecular radiotherapeutic that delivers cytotoxic radiation directly and selectively to cancer cells including cancer stem cells. A Phase Ib dose-escalation trial of I-131-CLR1404 in patients with advanced solid tumors was completed in the first quarter of 2014 and results presented at the American Society of Clinical Oncology (ASCO) 2014 Annual Meeting. CLR1502 is a preclinical, cancer-targeted, non-radioactive optical imaging agent for intraoperative tumor margin illumination and non-invasive tumor imaging. For additional information please visit [www.cellecstar.com](http://www.cellecstar.com).

---

## **INVESTOR CONTACT**

Kate McNeil, Vice President of IR, PR & Corporate Communications  
Cellecstar Biosciences, Inc.  
Phone: (347) 204-4226  
Email: [kmcneil@cellectar.com](mailto:kmcneil@cellectar.com)

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, 2014. These forward-looking statements are made only as of the date hereof, and we disclaim any obligation to update any such forward-looking statements.

---



### **Collectar Biosciences Reschedules Release of First Quarter Financials and Quarterly Conference Call**

**MADISON, Wis., May 11, 2015**, – Collectar Biosciences, Inc. (NASDAQ: CLRB), a clinical stage biopharmaceutical company developing innovative agents for the detection and treatment of cancer, announced that it is rescheduling the release of its first quarter financial results. The Company currently expects that it will file a Form 12b-25 to obtain additional time in which to complete its quarterly report on Form 10-Q for the first quarter of 2015, file its quarterly report with the SEC on or prior to May 20th and host a conference call with investors during the week of May 18<sup>th</sup>. Collectar will issue a press release with the revised date and time for the call prior to the event.

Collectar is postponing the release of its first quarter financial results to allow the audit committee, management and external auditors additional time to further evaluate the impact of a change in the accounting treatment of previously issued warrants. The potential change in accounting treatment may result in a non-cash, non-operating liability related to warrants previously reported as equity. The audit committee does not anticipate any impact on the Company's current or previously reported cash position, operating expenses or total operating, investing or financing cash flows.

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

Collectar Biosciences is developing agents to detect, treat and monitor a broad spectrum of cancers. Using a novel phospholipid ether analog (PLE) platform technology as a targeted delivery and retention vehicle, Collectar's compounds are designed to be selectively taken up and retained in cancer cells including cancer stem cells. With the ability to attach both imaging and therapeutic agents to its proprietary delivery platform, Collectar has developed a portfolio of product candidates engineered to leverage the unique characteristics of cancer cells to "find, treat and follow" malignancies in a highly selective way. I-124-CLR1404 is a small-molecule, broad-spectrum, cancer-targeted PET imaging agent currently being evaluated in a Phase II glioblastoma imaging trial. Additionally, multiple investigator-sponsored Phase I/II clinical trials are ongoing across 11 solid tumor indications. I-131-CLR1404 is a small-molecule, broad-spectrum, cancer-targeted molecular radiotherapeutic that delivers cytotoxic radiation directly and selectively to cancer cells including cancer stem cells. A Phase Ib dose-escalation trial of I-131-CLR1404 in patients with advanced solid tumors was completed in the first quarter of 2014 and results presented at the American Society of Clinical Oncology (ASCO) 2014 Annual Meeting. CLR1502 is a preclinical, cancer-targeted, non-radioactive optical imaging agent for intraoperative tumor margin illumination and non-invasive tumor imaging. For additional information please visit [www.collectar.com](http://www.collectar.com).

---

## **INVESTOR CONTACT**

Kate McNeil, Vice President of IR, PR & Corporate Communications  
Cellecstar Biosciences, Inc.  
Phone: (347) 204-4226  
Email: [kmcneil@cellectar.com](mailto:kmcneil@cellectar.com)

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. Factors that might cause such a material difference include, among others, potential delays in the completion of our analysis of the impact of the potential change in the accounting treatment of the previously issued warrants on our financial statements and the risk that additional information will come to light during the course of our analysis or the review thereof by our registered independent accounting firm that alters the scope or magnitude of the impact of that change. 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, 2014. These forward-looking statements are made only as of the date hereof, and we disclaim any obligation to update any such forward-looking statements.

---