U.S. 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): <u>August 4, 2015</u>

SUNSHINE BIOPHARMA, INC.

(Exact name of small business issuer as specified in its charter)

Colorado000-5289820-5566275(State or other jurisdiction of incorporation)(Commission File Number)(IRS Employer ID No.)

469 Jean-Talon West 3rd Floor Montreal, Quebec, Canada H3N 1R4

(Address of principal executive offices)

(514) 764-9698

(Issuer's Telephone Number)

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))

Item 8.01 Other Events

As previously reported on November 14, 2014, we entered into a Manufacturing Services Agreement with Lonza Ltd. and Lonza Sales Ltd. (hereinafter jointly referred to as "Lonza") of Basel, Switzerland whereby we engaged Lonza to be the manufacturer of our Adva-27a anticancer drug.

In June 2015, we received the initial sample of our Adva-27a manufacturing and on August 3, 2015 we completed our laboratory analysis of the same. Our test results show that the sample meets all of the biological specifications, clearing the way for the manufacturing of a 2-kilogram quantity for clinical trials.

Item 7.01 Regulation FD Disclosure

Our Press Release relating to the execution of the Lonza Agreement described above is attached as Exhibit 99.6 and is hereby incorporated.

Item 9.01 Financial Statements and Exhibits

(b) Exhibits. The following exhibits are included in this report:

No. Description

99.6 Press Release Announcing Test Results of Initial Sample of Adva-27a Manufacturing

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.

SUNSHINE BIOPHARMA, INC. Dated: August 4, 2015

(Registrant)

By: /s/ Dr. Steve N. Slilaty
Dr. Steve N. Slilaty, Chief Executive Officer



For Immediate Release August 4, 2015

SUNSHINE BIOPHARMA ANNOUNCES THAT THE INITIAL MANUFACTURING SAMPLE OF ITS ANTICANCER COMPOUND Adva-27a MEETS BIOLOGICAL SPECIFICATIONS PAVING THE WAY FOR CLINICAL TRIALS

Montreal, Quebec, Canada -- (MARKETWIRED) -- Sunshine Biopharma Inc. (OTCQB: "SBFM"), a pharmaceutical company focused on the research, development and commercialization of drugs for the treatment of various forms of cancer, announced today that it has completed analysis of the initial Adva-27a manufacturing sample received in June. The results show that the sample meets all of the biological specifications, clearing the way for the manufacturing of a 2-kilogram quantity for clinical trials.

"This is a major advancement in our Adva-27a drug development program," said Dr. Steve N. Slilaty, President and CEO of Sunshine Biopharma. "We are now well on our way to getting the 2 kilograms we need to conducts our Adva-27a clinical trials," he added.

About Adva-27a

Adva-27a is Sunshine Biopharma's lead anticancer compound, a Topoisomerase II inhibitor, small molecule that has recently been shown to be effective at killing Multidrug Resistant Breast Cancer cells, Small-Cell Lung Cancer cells, Uterine Sarcoma cells and Pancreatic Cancer cells (Published in ANTICANCER RESEARCH, Volume 32, Pages 4423-4432, October 2012). Adva-27a is currently in the IND-Enabling stage of development. The original U.S. patent covering Adva-27a was issued on August 7, 2012 under U.S. patent number 8,236,935. The Company is planning a Phase I clinical trial of Adva-27a for Pancreatic Cancer in parallel to the Phase I clinical trial of Adva-27a for multidrug resistant Breast Cancer to be conducted at McGill University's Jewish General Hospital in Montreal (Canada).

Safe Harbor Forward-Looking Statements

To the extent that statements in this press release are not strictly historical, including statements as to revenue projections, business strategy, outlook, objectives, future milestones, plans, intentions, goals, future financial conditions, future collaboration agreements, the success of the Company's development, events conditioned on stockholder or other approval, or otherwise as to future events, such statements are forward-looking, and are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. The forward-looking statements contained in this release are subject to certain risks and uncertainties that could cause actual results to differ materially from the statements made.

For Additional Information:

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