U.S. SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549

FORM 10-Q

Quarterly Report Under
the Securities Exchange Act of 1934

For Quarter Ended: **March 31, 2017**

Commission File Number: **000-52898**

SUNSHINE BIOPHARMA INC.

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

Colorado	**20-5566275**
(State of other jurisdiction of incorporation)	(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)

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days: Yes ☑ No ☐

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). Yes ☑ No ☐

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, smaller reporting company, or an emerging growth company. See the definitions of "large accelerated filer," "accelerated filer", "smaller reporting company", and "emerging growth company" in Rule 12b-2 of the Exchange Act. (Check one)

Large accelerated filer ☐	Accelerated filer ☐
Non-accelerated filer ☐	Smaller reporting company ☐
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. ☐

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). ☐ Yes ☑ No

The number of shares of the registrant's only class of common stock issued and outstanding as of May 12, 2017, was 858,594,650 shares.

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PART I.
FINANCIAL INFORMATION

Sunshine Biopharma, Inc.
Unaudited Consolidated Balance Sheet

	Unaudited March 31, 2017	Audited December 31, 2016
ASSETS		
Current Assets:		
Cash and cash equivalents	$ 17,143	$ 57,453
Prepaid expenses	846	1,007
Total Current Assets	17,989	58,460
Equipment (net of $2,797 and $2,272 depreciation, resepctively)	5,420	5,944
Patents (net of $58,918 amortization and $556,120 impairment)	-	-
TOTAL ASSETS	$ 23,409	$ 64,404
LIABILITIES AND SHAREHOLDERS' EQUITY		
Current liabilities:		
Current portion of notes payable	71,439	69,939
Current portion of notes payable - related party	183,889	167,032
Accounts payable	27,953	28,122
Interest payable	1,171	9,011
Total current liabilities	284,452	274,104
TOTAL LIABILITIES	284,452	274,104
COMMITMENTS AND CONTINGENCIES		
SHAREHOLDERS' EQUITY (DEFICIT)		
Preferred stock, Series A $0.10 par value per share; Authorized 850,000 Shares; Issued and outstanding -0- shares.	-	-
Preferred stock, Series B $0.10 par value per share; Authorized 500,000 Shares; Issued and outstanding 500,000 shares.	50,000	50,000
Common Stock, $0.001 par value per share; Authorized 3,000,000,000 Shares; Issued and outstanding 817,927,983 and 769,399,858 at March 31, 2017 and December 31, 2016, respectively Reserved for issuance 204,918,033 at March 31, 2017	817,928	769,400
Capital paid in excess of par value	11,642,783	11,548,460
Accumulated comprehensive income	1,903	394
Accumulated (Deficit)	(12,773,657)	(12,577,954)
TOTAL SHAREHOLDERS' EQUITY (DEFICIT)	(261,043)	(209,700)
TOTAL LIABILITIES AND SHAREHOLDERS' EQUITY (DEFICIT)	$ 23,409	$ 64,404

See Accompanying Notes To These Financial Statements.

Sunshine Biopharma, Inc.
Unaudited Consolidated Statement Of Operations and Comprehensive Loss

	Unaudited 3 Months Ended March 31, 2017	Unaudited 3 Months Ended March 31, 2016
Revenue:	$ -	$ -
General & Administrative Expenses		
Accounting	15,600	6,800
Amortization & depreciation	518	15,244
Consulting	25,937	14,184
Legal	14,904	28,099
Office	7,401	3,269
Officer & director remuneration	42,965	-
Stock transfer fee	1,665	2,086
Total G & A	108,990	69,682
(Loss) from operations	(108,990)	(69,682)
Other Income (expense):		
Foreign exchange gain	(639)	-
Interest expense	(9,144)	(6,954)
Litigation settlement proceeds	-	25,000
Loss on debt conversions	(76,929)	(253,658)
Total Other (Expense)	(86,712)	(235,612)
Net (loss)	$ (195,702)	$ (305,294)
Basic (Loss) per common share	$ 0.00	$ 0.00
Weighted Average Common Shares Outstanding	805,604,089	215,596,040
Net Income (Loss)	$ (195,702)	$ (305,294)
Other comprehensive income:		
Unrealized Gain (Loss) from foreign exchange translation	1,509	(9,991)
Comprehensive (Loss)	(194,193)	(315,285)
Basic (Loss) per common share	$ 0.00	$ 0.00
Weighted Average Common Shares Outstanding	805,604,089	215,596,040

See Accompanying Notes To These Financial Statements.

Sunshine Biopharma, Inc.
Unaudited Consolidated Statement Of Cash Flows

	Unaudited 3 Months Ended March 31, 2017	Unaudited 3 Months Ended March 31, 2016
Cash Flows From Operating Activities:		
Net (Loss)	$ (195,702)	$ (305,294)
Adjustments to reconcile net loss to net cash used in operating activities:		
Foreign exchange loss	639	
Depreciation/Amortization	525	15,244
Stock issued for payment interest	3,022	3,120
Loss on debt conversion	76,929	253,658
Stock issued for payment of expenses	14,400	-
(Increase) decrease in prepaid expenses	161	(44)
Increase (decrease) in Accounts Payable & accrued expenses	(171)	(52,116)
Increase (decrease) in interest payable	(7,840)	3,834
Net Cash Flows (used) in operations	(108,037)	(81,598)
Cash Flows From Investing Activities:		
Purchase of equipment	-	-
Net Cash Flows (used) in Investing activities	-	-
Cash Flows From Financing Activities:		
Proceed from note payable	50,256	82,000
Note payable used to pay expenses	13,962	-
Note payable used to pay origionation fees & interest	2,000	3,000
Sale of common stock	-	79,128
Net Cash Flows provided by financing activities	66,218	164,128
Net Increase (Decrease) In Cash and cash equivalents	(41,819)	82,530
Foreign currency translation adjustment	1,509	(9,991)
Cash and cash equivalents at beginning of period	57,453	50,798
Cash and cash equivalents at end of period	$ 17,143	$ 123,337
Supplementary Disclosure Of Cash Flow Information:		
Stock issued for services, licenses and other assets	$ 14,400	$ 100,000
Stock issued for note conversions including interest	$ 128,451	$ 377,814
Cash paid for interest	$ -	$ -
Cash paid for income taxes	$ -	$ -

See Accompanying Notes To These Financial Statements.

Note 1 – Description of Business

Mountain West Business Solutions, Inc. (“MWBS”) was incorporated on August 31, 2006 in the State of Colorado. Sunshine Etopo, Inc. (formerly Sunshine Biopharma, Inc.) was incorporated in the State of Colorado on August 17, 2009. Effective October 15, 2009, MWBS was acquired by Sunshine Etopo, Inc. in a transaction classified as a reverse acquisition. MWBS concurrently changed its name to Sunshine Biopharma, Inc. Sunshine Etopo, Inc. has been inactive and was recently dissolved. In July 2014, the Company formed a wholly owned Canadian subsidiary, Sunshine Biopharma Canada Inc. The financial statements represent the consolidated activity of Sunshine Biopharma, Inc. and Sunshine Biopharma Canada Inc. (hereinafter together referred to as the "Company"). The Company was formed for the purposes of conducting research, development and commercialization of drugs for the treatment of various forms of cancer. The Company may also engage in any other business that is permitted by law, as designated by the Board of Directors of the Company.

The Company’s wholly owned Canadian Subsidiary, Sunshine Biopharma Canada Inc. (“Sunshine Canada”) was formed for the purposes of offering generic pharmaceutical products in Canada and elsewhere around the world. Sunshine Canada has recently signed licensing agreements for four (4) generic prescription drugs for treatment of breast cancer, prostate cancer and BPH (Benign Prostatic Hyperplasia). In addition, Sunshine Canada is currently evaluating several other generic products for in-licensing and is working on securing a Drug Establishment License (“DEL”) and a Drug Identification Number (“DIN”) per product from Health Canada. Once the DEL and the DIN’s are secured, Sunshine Canada will begin its generic pharmaceuticals marketing and sales program in Canada and overseas.

In addition, the Company has been and continues to work on the development of its proprietary anticancer drug, Adva-27a. The next series of steps in the development of Adva-27a include (i) GMP-manufacturing of a 2-kilogram quantity of the drug, (ii) completing requisite IND-enabling studies, and (iii) conducting Phase I clinical trials. In the preclinical studies, Adva-27a was shown to be effective at destroying multidrug resistant cancer cells including Pancreatic Cancer, Breast Cancer, Lung Cancer and Uterine Sarcoma, cells.

During the last three month period the Company has continued to raise money through stock sales and borrowings.

The Company’s activities are subject to significant risks and uncertainties, including failing to secure additional funding to operationalize the Company’s generics business and proprietary drug development program.

Note 2 – Summary of Significant Accounting Policies

See the Notes in the 2016 Form 10-K consolidated financial statements for a complete summary of the Company’s significant accounting policies.

This summary of significant accounting policies is presented to assist the reader in understanding the Company's financial statements. The consolidated financial statements and notes are representations of the Company's management, which is responsible for their integrity and objectivity. These accounting policies conform to generally accepted accounting principles and have been consistently applied in the preparation of the financial statements.

PRINCIPLES OF CONSOLIDATION

The accompanying consolidated financial statements include the accounts of the Company and its wholly owned subsidiaries. All intercompany accounts and transactions have been eliminated in consolidation.

USE OF ESTIMATES

The preparation of financial statements in conformity with U.S. generally accepted accounting principles requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period. The more significant estimates and assumptions made by management are valuation of equity instruments, depreciation of property and equipment, and deferred tax asset valuation. Actual results could differ from those estimates as the current economic environment has increased the degree of uncertainty inherent in these estimates and assumptions.

CASH AND CASH EQUIVALENTS

For the Balance Sheets and Statement of Cash Flows, all highly liquid investments with maturity of 90 days or less are considered to be cash equivalents. The Company had a cash balance of $17,143 and $57,453 as of March 31, 2017 and December 31, 2016, respectively. At times such cash balances may be in excess of the FDIC limit of $250,000.

EARNINGS PER SHARE

The Company has adopted the FASB ASC Topic 260 regarding earnings / loss per share, which provides for calculation of “basic” and “diluted” earnings / loss per share. Basic earnings / loss per share includes no dilution and is computed by dividing net income / loss available to common shareholders by the weighted average common shares outstanding for the period. Diluted earnings / loss per share reflect the potential dilution of securities that could share in the earnings of an entity similar to fully diluted earnings / loss per share.

Other than the Notes Payable specified under Note 4 and Note 5 below, there were no potentially dilutive instruments outstanding during the interim period ended March 31, 2017 or the year ended December 31, 2016.

INCOME TAXES

The Company follows the asset and liability method of accounting for deferred income taxes. The asset and liability method requires the recognition of deferred tax assets and liabilities for the expected future tax consequences of temporary differences between financial accounting and tax bases of assets and liabilities. The Company accounts for income taxes pursuant to ASC 740. There was no increase in liabilities for unrecognized tax benefits as a result of this implementation. The Company recognizes accrued interest related to unrecognized tax benefits in interest expense and penalties in general and administrative expense.

FUNCTIONAL CURRENCY

The U.S. dollar is the functional currency of the Company which is operating in the United States. The functional currency for the Company's Canadian subsidiary is the Canadian dollar. The Company translates its Canadian subsidiary's financial statements into U.S. dollars as follows:

- Assets and liabilities are translated at the exchange rate in effect as of the financial statement date.
- Income statement accounts are translated using the weighted average exchange rate for the period.

The Company includes translation adjustments from currency exchange and the effect of exchange rate changes on intercompany transactions of a long-term investment nature as a separate component of shareholders' equity. There are currently no transactions of a long-term investment nature, nor any gains or losses from non-U.S. currency transactions.

REVENUE RECOGNITION

Since inception, the Company has been focused on the research, development and commercialization of drugs for the treatment of various forms of cancer. Following its recent entry into the generic pharmaceuticals business, the Company has become a fully integrated pharmaceutical company offering generic and proprietary drugs for the treatment of cancer and other acute and chronic indications. The Company intends to recognize revenues from the sales of generic pharmaceuticals, if or when they occur, at the time the products are sold and collectability is assured. In the event the Company provides consulting services in the future, revenues from such services will be recognized when the services are rendered and invoiced.

GOING CONCERN

The accompanying financial statements have been prepared on a going concern basis, which contemplates the realization of assets and the satisfaction of liabilities in the normal course of business. Since inception, the Company has had recurring operating losses and negative operating cash flows. These factors raise substantial doubt about the Company's ability to continue as a going concern.

The Company's continuation as a going concern is dependent on its ability to obtain additional financing to fund operations, implement its business model, and ultimately, attain profitable operations. The Company will need to secure additional funds through various means, including equity and debt financing or any similar financing. There can be no assurance that the Company will be able to obtain additional equity or debt financing, if and when needed, on terms acceptable to the Company, or at all. Any additional equity or debt financing may involve substantial dilution to the Company's stockholders, restrictive covenants or high interest costs. The Company's long-term liquidity also depends upon its ability to generate revenues and achieve profitability.

The accompanying financial statements do not include any adjustments relating to the recoverability and classification of recorded asset amounts or the amounts and classification of liabilities that might be necessary should the Company be unable to continue as a going concern.

Note 3 – Unaudited Financial Information

The unaudited financial information included for the three month interim period ended March 31, 2017 was taken from the books and records of the Company without audit. However, such information reflects all adjustments, consisting only of normal recurring adjustments, which in the opinion of management are necessary to reflect properly the results of the interim periods presented. The results of operations for the three month interim period ended March 31, 2017 are not necessarily indicative of the results expected for the fiscal year ending December 31, 2017.

Note 4 – Notes Payable

A Note Payable having a face value of $21,439 and a maturity date of December 31, 2017 was entered into on December 31, 2016. This Note accrues interest at a rate of 12% per annum and is convertible after December 31, 2016 into $0.001 par value Common Stock at a price 35% below market value. The Company estimates that the fair value of this convertible debt approximates the face value, so no value has been assigned to the beneficial conversion feature.

A Note payable dated July 1, 2016, having a face value of $55,000 and a principal balance of $48,500 at December 31, 2016, accrued interest at 10%. This note was convertible into $0.001 par value Common Stock at a price 40% below market value. During the three month period ended March 31, 2017, a total of $48,500 in principal and $3,022 of interest was converted into $0.001 par value Common Stock leaving a principal balance of $-0-. In connection with this conversion, 42,528,125 shares of $0.001 par value Common Stock valued at $128,451 were issued generating a loss of $76,929 on conversion.

On February 10, 2017, the Company received net proceeds of $48,000 in exchange for a note payable having a face value of $50,000 and accruing interest at the rate of 8% per annum. The note, due on November 20, 2017, is convertible after 180 days from issuance into $0.001 par value Common Stock at a price 39% below market value. The Company estimates that the fair value of this convertible debt approximates the face value, so no value has been assigned to the beneficial conversion feature. In connection with this note, 204,918,033 shares of the Company's Common Stock have been reserved for issuance.

At March 31, 2017 and December 31, 2016, accrued interest on Notes Payable was $15,124 and $9,011, respectively.

Note 5 – Notes Payable Related Party

In December 2016, the Company received monies from its CEO in exchange for a note payable having a principal amount of $90,000 Canadian ($67,032 US) '–with interest at 12% due March 31, 2017. The note is convertible any time after the date of issuance into $0.001 par value Common Stock at a price 35% below market value. The Company estimated that the fair value of the convertible debt approximates the face value, so no value has been assigned to the beneficial conversion feature. Any gain or loss will be recognized at conversion. This note is collateralized by all of the assets of the Company. In the event of default, the interest rate will increased to 18% per annum and a penalty of $1,000 Canadian ($752 US) per day will accrue. On March 31, 2017, the note, together with accrued interest of $3,021 Canadian ($2,271 US) and an additional principal amount of $3,000 Canadian ($2,247 US) paid to the Company on March 28, 2017, was renewed for a 90-day period under the same terms and conditions as the original note. The new note now having a face value of $96,021 Canadian ($72,198 US) is due on June 30, 2017.

A note payable held by a private individual who became a principal shareholder of the Company having a face value of $100,000 at December 31, 2016 and a maturity date of March 31, 2017, accrues interest at 12%. The Note is convertible any time from the date of issuance into $0.001 par value Common Stock at a 35% discount from market price. The Company estimates that the fair value of this convertible debt approximates the face value, so no value has been assigned to the beneficial conversion feature. On March 31, 2017, the note's principal balance of $100,000 plus accrued interest of $11,715 was renewed for a period of 90-day under the same terms and conditions as the original note. The new note now having a face value of $111,715 matures on June 30, 2017.

Note 6 – Issuance of Common Stock

During the three months ended March 31, 2017, the Company issued a total of 48,428,125 shares of $0.001 par value Common Stock. Of these 42,428,125 shares valued at $128,451 were issued upon conversion of outstanding notes payable, reducing the debt by $48,500 and interest payable by $3,022 and generating a loss on conversion of $76,929.

In addition, the Company issued 6,000,000 shares of $0.001 par value Common Stock valued at $14,400 for $15,000 Canadian ($11,278 US). The $15,000 Canadian was paid directly to a firm which was engaged to conduct a valuation of the Company's patents. The difference of $3,122 was recorded as a consulting expense.

In connection with its convertible notes payable, the Company has 204,918,033 shares of its Common Stock reserved for issuance at March 31,2017.

The Company declared no dividends through March 31, 2017.

Note 7 – Earnings (Loss) Per Share

Earnings (loss) per share is computed using the weighted average number of Common Shares outstanding during the period. The Company has adopted ASC 260 (formerly SFAS128), "Earnings per Share".

Note 8 -- Generic Drugs Licenses

In 2016, the Company entered into License Agreements for the following four Generic Drugs:

- Anastrozole (brand name Arimidex® by AstraZenica) for treatment of Breast Cancer
- Letrozole (brand name Femara® by Novartis) for treatment of Breast Cancer
- Bicalutamide (brand name Casodex® by AstraZenica) for treatment of Prostate Cancer
- Finasteride (brand name Propecia® by Merck) for treatment of Benign Prostatic Hyperplasia

The cost of these Licenses has been fully expensed.

Note 9 – Financial Statements

For a complete set of footnotes, reference is made to the Company's Report on Form 10-K for the year ended December 31, 2016, as filed with the Securities and Exchange Commission and the audited financial statements and notes included therein.

Note 10 – Income Taxes

Deferred income taxes arise from the temporary differences between financial statement and income tax recognition of net operating losses and other items. Loss carryovers are limited under the Internal Revenue Code should a significant change in ownership occur.

A deferred tax asset at each date has been offset by a 100% valuation allowance.

Note 11 – Royalties Payable

As part of a subscription agreement entered into in 2016, the Company has an obligation to pay a royalty of 5% of net sales on one of its generic products (Anastrozole) for a period of three (3) years from the date of the first sale of that product.

Note 12 – Related Party Transactions

In addition to the related party transactions detailed in Note 5 above, during the three month period ended March 31, 2017, the Company paid its Officers and Directors cash compensation totaling $42,965. $1,519 was paid to the Company's COO and $41,446 was paid to the Company's CEO.

During the fiscal year ended December 31, 2016, Advanomics Corporation paid on behalf of the Company $13,725 Canadian in patenting fees. Advanomics was fully reimbursed by the Company in January 2017.

Note 13 – Subsequent Events

In April 2017, the Company received monies in exchange for a convertible note payable having a face value of $100,000 Canadian (approximately $75,190 US).

On April 3, 2017, the Company sold 34,000,000 shares of its $0.001 par value Common Stock valued at $98,600 for $85,000 Canadian (approximately $63,912 US).

On April 24, 2017, the Company issued 6,666,667 shares of its $0.001 par value Common Stock for the purchase of laboratory equipment valued at $22,000.

On April 26, 2017, the Company received monies in exchange for a convertible note payable having a face value of $65,000. In connection with this note, 100,000,000 shares of the Company's Common Stock have been reserved for issuance.

PART I.

ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

The following discussion should be read in conjunction with our consolidated financial statements and notes thereto included herein. In connection with, and because we desire to take advantage of, the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995, we caution readers regarding certain forward looking statements in the following discussion and elsewhere in this report and in any other statement made by, or on our behalf, whether or not in future filings with the Securities and Exchange Commission. Forward looking statements are statements not based on historical information and which relate to future operations, strategies, financial results or other developments. Forward looking statements are necessarily based upon estimates and assumptions that are inherently subject to significant business, economic and competitive uncertainties and contingencies, many of which are beyond our control and many of which, with respect to future business decisions, are subject to change. These uncertainties and contingencies can affect actual results and could cause actual results to differ materially from those expressed in any forward looking statements made by, or on our behalf. We disclaim any obligation to update forward looking statements.

Overview and History

We were incorporated in the State of Colorado on August 31, 2006 under the name "Mountain West Business Solutions, Inc." Until October 2009, our business was to provide management consulting with regard to accounting, computer and general business issues for small and home-office based companies. Effective October 15, 2009, we executed an agreement to acquire Sunshine Biopharma, Inc., a Colorado corporation, in exchange for the issuance of 21,962,000 shares of our Common Stock and 850,000 shares of Convertible Preferred Stock, each convertible into twenty (20) shares of our Common Stock. As a result of this transaction we changed our name to "Sunshine Biopharma, Inc. and our officers and directors resigned their positions with us and were replaced by our current management. The majority of the Common Shares and all of the Convertible Preferred Shares we issued for this transaction were issued to Advanomics Corporation, a privately held Canadian company ("Advanomics"). On December 21, 2011, Advanomics exercised its right to convert the 850,000 shares of Series "A" Preferred Stock it held in our Company into 17,000,000 shares of Common Stock.

Following the above detailed transactions, we began to operate as a pharmaceutical company focusing on development of the Adva-27a anticancer compound. We operated under a the exclusive technology license agreement with Advanomics until December 2015, at which time we acquired all of the worldwide right to the technology and became direct owner of all issued and pending patents pertaining to the Adva-27a technology. Following acquisition of the Adva-27a patents, the exclusive license agreement with Advanomics was terminated and Sunshine Etopo, Inc., Sunshine Biopharma Inc.'s subsidiary holding the exclusive license with Advanomics, was dissolved.

In July 2014, we formed a wholly owned Canadian subsidiary, Sunshine Biopharma Canada Inc. ("Sunshine Canada"), for the purposes of offering generic pharmaceutical products in Canada and elsewhere around the globe. Sunshine Canada has recently signed licensing agreements for four (4) generic prescription drugs for the treatment of cancer and BPH (Benign Prostatic Hyperplasia). With our entry into the generic pharmaceuticals business, we have become a fully integrated pharmaceutical company offering generic and proprietary drugs for the treatment of cancer and other acute and chronic indications.

Our principal place of business is located at 469 Jean-Talon West, 3rd Floor, Montreal, Quebec, Canada H3N 1R4. Our phone number is (514) 764-9698 and our website address is www.sunshinebiopharma.com.

We have not been subject to any bankruptcy, receivership or similar proceeding.

Results Of Operations

Comparison of Results of Operations for the three months ended March 31, 2017 and 2016

For the three months ended March 31, 2017 and 2016, we did not generate any revenues.

General and administrative expenses during the three month period ended March 31, 2017 were $108,990, compared to general and administrative expense of $69,682 incurred during the three month period ended March 31, 2016, an increase of $39,308. This increase is primarily attributable to an increase of approximately $43,000 in executive compensation. We also incurred increased accounting and consulting fees, but lower legal fees during the three months ended March 31, 2017, compared to the similar period in 2016. Most of our other expenses remained relatively constant during the three month period ended March 31, 2017 compared to the similar period in 2016. We also incurred $9,144 in interest expense during the three months ended March 31, 2017, compared to $6,954 in interest expense during the similar period in 2016 as a result of decreased borrowings. However, we incurred $76,929 in losses arising from debt conversion during the three months ended March 31, 2017, compared to $253,658 in losses from debt conversion during the similar period in 2016, a difference of $176,729 as a result of a smaller amount of convertible notes outstanding. We also received $25,000 from the settlement of litigation in 2016 that we did not receive during the three months ended March 31, 2017.

As a result, we incurred a net loss of $195,702 ($0.00 per share) for the three month period ended March 31, 2017, compared to a net loss of $305,294 ($0.00 per share) during the three month period ended March 31, 2016.

Because we did not generate any revenues since our inception, following is our Plan of Operation.

PLAN OF OPERATION

Since inception, we have been operating as a pharmaceutical company focused on the research, development and commercialization of proprietary drugs for the treatment of various forms of cancer. In July 2014, we formed Sunshine Biopharma Canada Inc., a Canadian wholly owned subsidiary, for the purposes of conducting generic pharmaceuticals business in Canada and elsewhere around the world. During 2016, we intensified our activities in the generic pharmaceuticals area as we continued to pursue our proprietary anticancer drug development efforts. Accordingly, we have become a fully integrated pharmaceutical company offering generic and proprietary drugs for the treatment of cancer and other acute and chronic indications. Below we describe our Generic Pharmaceuticals Operations followed by our Proprietary Drug Development Program.

GENERIC PHARMACEUTICALS OPERATIONS

In 2016, our Canadian wholly owned subsidiary, Sunshine Biopharma Canada Inc. ("Sunshine Canada"), signed Cross Referencing Agreements with a major pharmaceutical company for four prescription generic drugs for the treatment of Breast Cancer, Prostate Cancer and Enlarged Prostate. We will market and sell these new pharmaceutical products under our own Sunshine Biopharma label. These four generic products are as follows:

- Anastrozole (brand name Arimidex® by AstraZeneca) for treatment of Breast Cancer;
- Letrozole (brand name Femara® by Novartis) for treatment of Breast Cancer;
- Bicalutamide (brand name Casodex® by AstraZenica) for treatment of Prostate Cancer;
- Finasteride (brand name Propecia® by Merck) for treatment of BPH (Benign Prostatic Hyperplasia)

Worldwide sales of the brand name version of these products as reported in the SEC filing of the respective owner of the registered trademark are as follows:

- Arimidex® $232M in 2016
- Femara® $380M in 2014
- Casodex® $247M in 2016
- Propecia® $183M in 2015

Sunshine Canada is currently in the process of securing a Drug Identification Number ("DIN") for each of these products from Health Canada. We are also working on finding an appropriate facility and obtaining a Drug Establishment License ("DEL") from Health Canada. Upon receipt of the DEL and DIN's, we will be able to accept orders for our own label SBI-Anastrozole, SBI-Letrozole, SBI-Bicalutamide and SBI-Finasteride. We cannot estimate the timing in our obtaining either the DIN or DEL due to variables involved that are out of our control. The Figure below shows our 30-Pill blister pack of Anastrozole.

We currently have twenty three (23) additional Generic Pharmaceuticals under review for in-licensing. While no assurances can be provided, when completed, this will bring our Generic Products portfolio to a total of twenty seven (27). We believe that a larger product portfolio provides us with more opportunities and a greater reach into the marketplace. We hope to further build our generics portfolio of "SBI" label Generic Pharmaceuticals over time.

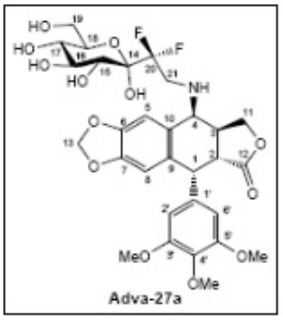
Various publicly available sources indicate that the worldwide sales of generic pharmaceuticals are approximately $200 billion per year. In the United States and Canada, the sales of generic pharmaceuticals are approximately $50 billion and $5 billion, respectively. The generic pharmaceuticals business is fairly competitive and there are several multinational players in the field including Teva (Israel), Novartis - Sandoz (Switzerland), Hospira (USA), Mylan (Netherlands), Sanofi (France), Fresenius Kabi (Germany) and Apotex (Canada). While no assurances can be provided, with our offering of Canadian approved products we believe that we will be able to access at least a small percentage of the generic pharmaceuticals marketplace.

As part of a subscription agreement, we have an obligation to pay a royalty of 5% of net sales on one of our generic products (Anastrozole) for a period of three (3) years from the date of the first sale of that product.

While no assurances can be provided and subject to the availability of adequate financing, of which there is no assurance, we anticipate that profits from the sales of Generic Products will be used to finance our proprietary drug development program, including Adva-27a, our flagship anticancer compound. In addition to near-term revenue generation, building the generics business infrastructure and securing the proper permits will render us appropriately positioned for the marketing and distribution of our proprietary Adva-27a drug candidate, provided that Adva-27a is approved for such marketing and distribution, of which there can be no assurance.

PROPRIETARY DRUG DEVELOPMENT OPERATIONS

Our proprietary drug development activities have been focused on the development of a small molecule called Adva-27a for the treatment of aggressive cancers. A Topoisomerase II inhibitor, Adva-27a has been shown to be effective at destroying Multidrug Resistant cancer cells including Pancreatic Cancer cells, Breast Cancer cells, Small-Cell Lung Cancer cells and Uterine Sarcoma cells (Published in ANTICANCER RESEARCH, Volume 32, Pages 4423-4432, October 2012). Sunshine Biopharma is direct owner of all issued and pending worldwide patents pertaining to Adva-27a including U.S. Patent Number 8,236,935. See "Part I, Item 1 – Business - Intellectual Property."



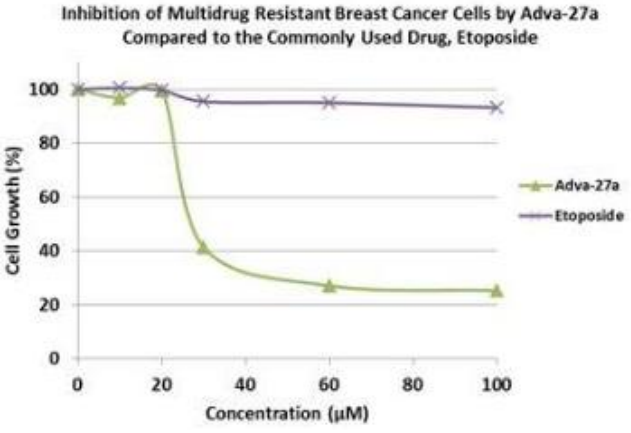
Adva-27a

Summary of Adva-27a Preclinical Studies

Adva-27a is a GEM-difluorinated C-glycoside derivative of Podophyllotoxin. Another derivative of Podophyllotoxin called Etoposide is currently on the market and is used to treat various types of cancer including leukemia, lymphoma, testicular cancer, lung cancer, brain cancer, prostate cancer, bladder cancer, colon cancer, ovarian cancer, liver cancer and several other forms of cancer. Etoposide is one of the most widely used anticancer drugs. Adva-27a and Etoposide are similar in that they both attack the same target in cancer cells, namely the DNA unwinding enzyme, Topoisomerase II. Unlike Etoposide, and other anti-tumor drugs currently in use, Adva-27a is able to destroy Multidrug Resistant Cancer cells. Adva-27a is the only compound known today that is capable of destroying Multidrug Resistant Cancer. In addition, Adva-27a has been shown to have distinct and more desirable biological and pharmacological properties compared to Etoposide. In side-by-side studies using Multidrug Resistant Breast Cancer cells and Etoposide as a reference, Adva-27a showed markedly improved cell killing activity (see Figure below). Our preclinical studies to date have shown that:

- Adva-27a is effective at killing different types of Multidrug Resistant cancer cells, including Pancreatic Cancer Cells (Panc-1), Breast Cancer Cells (MCF-7/MDR), Small-Cell Lung Cancer Cells (H69AR), and Uterine Sarcoma Cells (MES-SA/Dx5).
- Adva-27a is unaffected by P-Glycoprotein, the enzyme responsible for making cancer cells resistant to anti-tumor drugs.
- Adva-27a has excellent clearance time (half-life = 54 minutes) as indicated by human microsomes stability studies and pharmacokinetics data in rats.
- Adva-27a clearance is independent of Cytochrome P450, a mechanism that is less likely to produce toxic intermediates.
- Adva-27a is an excellent inhibitor of Topoisomerase II with an IC50 of only 13.7 micromolar (this number has recently been reduce to 1.44 micromolar as a result of resolving the two isomeric forms of Adva-27a).
- Adva-27a has shown excellent pharmacokinetics profile as indicated by studies done in rats.
- Adva-27a does not inhibit tubulin assembly.

These and other preclinical data have been published in ANTICANCER RESEARCH, a peer-reviewed International Journal of Cancer Research and Treatment. The publication which is entitled "Adva-27a, a Novel Podophyllotoxin Derivative Found to Be Effective Against Multidrug Resistant Human Cancer Cells" [ANTICANCER RESEARCH 32: 4423-4432 (2012)] is available on our website at www.sunshinebiopharma.com.



Clinical Development Path

The early stage preclinical studies for our lead compound, Adva-27a, were successfully completed and the results have been published in ANTICANCER RESEARCH 32: 4423-4432 (2012). We have been delayed in our implementation of our clinical development program due to lack of funding. Our fund raising efforts are continuing and as soon as adequate financing is in place we will continue our clinical development program of Adva-27a by conducting the next sequence of steps comprised of the following:

- GMP Manufacturing of 2 kilogram for use in IND-Enabling Studies and Phase I Clinical Trials
- IND-Enabling Studies
- Regulatory Filing (Fast-Track Status Anticipated)
- Phase I Clinical Trials (Pancreatic Cancer and in parallel Multidrug Resistant Breast Cancer)

GMP Manufacturing

In November 2014, we entered into a Manufacturing Services Agreement with Lonza Ltd. and Lonza Sales Ltd. (hereinafter jointly referred to as "Lonza"), whereby we engaged Lonza to be the manufacturer of our Adva-27a anticancer drug. In June 2015 we received a sample of the pilot manufacturing run for evaluation. Our laboratory analyses showed that, while the sample meets the required biological specifications, the amount of material generated (the "Yield") by the pilot run was found to be significantly lower than planned. During the course of our discussions concerning the problem of the low Yield, Lonza informed us that they required us to pay them $687,818 prior to moving forward with any activity pertaining to the manufacturing agreement we have with them. We have repeatedly indicated to Lonza that a clear path defining exactly how the extremely low Yield issue would be addressed is imperative prior to us making any payments. As of the date of this report, neither party has changed its position. See "Part I, Item 3 – Legal Proceedings."

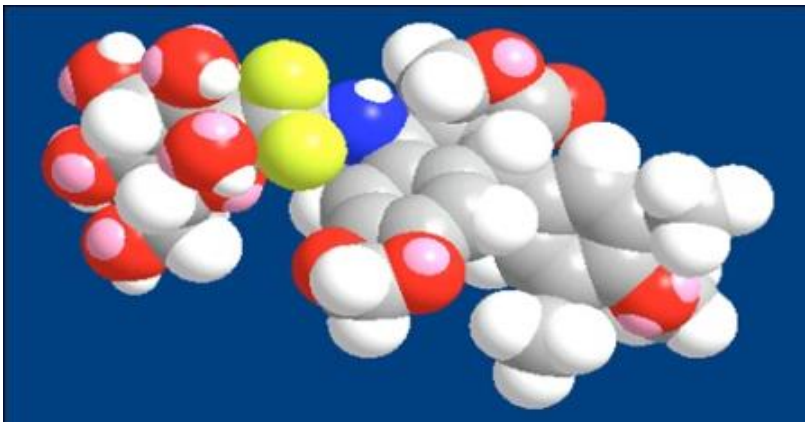
Clinical Trials

Adva-27a's initial indication will be pancreatic cancer for which there are currently little or no treatment options available. We have concluded an agreement with McGill University's Jewish General Hospital in Montreal, Canada to conduct Phase I clinical trials for this indication. All aspects of the planned clinical trials in Canada will employ FDA standards at all levels. Subject to obtaining the necessary financing, we now anticipate that Phase I clinical trials will commence in mid-2018 and we estimate that it will take 18 months to complete, at which time we expect to receive limited marketing approval for "compassionate-use" under the FDA and similar guidelines in Canada. See "Potential Near-Term Opportunities" below.

Potential Near-Term Opportunities

According to the American Cancer Society, nearly 1.5 million new cases of cancer are diagnosed in the U.S. each year. Given the terminal and limited treatment options available for the pancreatic cancer indication we are planning to study, we anticipate being granted limited marketing approval ("compassionate-use") for our Adva-27a following receipt of funding and a successful Phase I clinical trial. There are no assurances that either will occur. Such limited approval will allow us to make the drug available to various hospitals and health care centers for experimental therapy and/or "compassionate-use", thereby generating revenues in the near-term.

In addition, we believe that upon successful completion of Phase I Clinical Trials we may receive one or more offers from large pharmaceutical companies to buyout or license our drug at a significant premium. However, there are no assurances that our Phase I Trials will be successful, or if successful, that any pharmaceutical companies will make an acceptable offer to us. In the event we do not consummate such a transaction, we will require significant capital in order to complete the requisite additional clinical trials towards a potential full marketing approval, of which there can be no assurance.



A Space-Filling Model of Our Anticancer Compound, Adva-27a

Intellectual Property

Effective October 8, 2015, we executed a Patent Purchase Agreement (the "October Purchase Agreement"), with Advanomics, a related party, pursuant to which we acquired all of the right, title and interest in and to U.S. Patent Number 8,236,935 (the "US Patent") for our anticancer compound, Adva-27a. On December 28, 2015, we executed a second Patent Purchase Agreement (the "December Purchase Agreement"), with Advanomics, pursuant to which we acquired all of the right, title and interest in and to all of the remaining worldwide rights covered by issued and pending patents under PCT/FR2007/000697 and PCT/CA2014/000029 (the "Worldwide Patents") for our anticancer compound, Adva-27a.

Effective December 28, 2015, we entered into amendments (the "Amendments") of these Purchase Agreements pursuant to which the total purchase price was reduced from $17,142,499 to $618,810, the book value of this intellectual property on the financial statements of Advanomics. Further, the Amendments provided for automatic conversion of the promissory notes representing the new purchase price into an aggregate of 321,305,415 shares of our Common Stock once we increase our authorized capital such that these shares can be issued. In July 2016 we increased our authorized capital and issued the 321,305,415 Common shares to Advanomics thereby completing all aspects of the patent purchase arrangements and securing direct ownership of all worldwide patents and rights pertaining to Adva-27a.

In addition, in 2016 we signed Cross Referencing Agreements with a major pharmaceutical company for four (4) prescription generic drugs for the treatment of Breast Cancer, Prostate Cancer and Enlarged Prostate. These agreements give us the right to register the four (4) generic products, Anastrozole, Letrozole, Bicalutamide and Finasteride in Canada under our own label and obtain a DIN for each in order to be able to place them on the market.

While no assurances can be provided, we are also planning to expand our product line through acquisitions and/or in-licensing as well as in-house research and development.

Liquidity and Capital Resources

As of March 31, 2017, we had cash or cash equivalents of $17,143.

Net cash used in operating activities was $108,037 during the three month period ended March 31, 2017, compared to $81,598 for the three month period ended March 31, 2016. We anticipate that overhead costs and other expenses will increase in the future as we move forward with our proprietary drug development activities and our efforts to become a fully integrated pharmaceutical company by offering generic pharmaceutical products as discussed above.

Cash flows from financing activities were $66,218 for the three month periods ended March 31, 2017, compared to $164,128 during the three months ended March 31, 2016. Cash flows used by investing activities were $0 for the three month periods ended March 31, 2017 and 2016.

During the three months ended March 31, 2017, we issued a total of 48,428,125 shares of our Common Stock. Of these, 42,428,125 shares valued at $128,451 were issued upon conversion of outstanding notes payable, reducing the debt by $48,500 and interest payable by $3,022 and generating a loss on conversion of $76,929. In addition, we issued 6,000,000 shares of our Common Stock valued at $14,400 for $15,000 Canadian ($11,278 US). The $15,000 Canadian was paid directly to a firm which was engaged to conduct a valuation of our patents. We believe that having an independent valuation of our patents will be helpful in connection with our ongoing financing efforts.

On March 29, 2016, we issued 10,000,000 shares of our Common Stock having a market value of $100,000 or $0.01 per share, to an unaffiliated company that is assisting us in the development of manufacturing, marketing, sales and distribution contracts for various generic pharmaceuticals and biomedical products in Canada. These services are for a two year period but the value of these shares ($100,000) was expensed in the three month period ended March 31, 2016.

In March 2016, our Board of Directors authorized a private offering of our Common Stock in Canada pursuant to Regulation S promulgated under the Securities Act of 1933, as amended, wherein we offered up to 60,000,000 shares of our Common Stock at an offering price of $0.015 Canadian per share for aggregate gross proceeds of up to $900,000 Canadian. We accepted two subscriptions each of 10,000,000 shares of our Common Stock for $150,000 Canadian, aggregating $300,000 Canadian or approximately $236,550 US.

We are not generating revenue from our operations, and our ability to implement our business plan as set forth herein will depend on the future availability of financing. Such financing will be required to enable us to further develop our generic pharmaceuticals business and proprietary drug development program. We intend to raise funds through private placements of our Common Stock and/or debt financing. We estimate that we will require approximately $6 million ($1 million for the generic pharmaceutical operations and $5 million for the proprietary drug development program) to fully implement our business plan in the future and there are no assurances that we will be able to raise this capital. While we are engaged in discussions with various investment banking firms and venture capitalists to provide us these funds, as of the date of this report we have not reached any agreement with any party that has agreed to provide us with the capital necessary to effectuate our business plan. Our inability to obtain sufficient funds from external sources when needed will have a material adverse effect on our plan of operation, results of operations and financial condition.

Our cost to continue operations as they are now conducted is nominal, but these are expected to increase as we move forward with implementation of our business plan. We do not have sufficient funds to cover the anticipated increase in the relevant expenses. We need to raise additional funds in order to continue our existing operations, to initiate R&D activities, and to finance our plans to expand our operations for the next year. If we are successful in raising additional funds, we expect our operations and business efforts to continue and expand. There are no assurances this will occur.

Subsequent Events

In April 2017 we engaged in the following transactions:

- We received monies in exchange for a convertible note payable having a face value of $100,000 Canadian (approximately $75,190 US).
- We sold 34,000,000 shares of our Common Stock valued at $98,600 for $85,000 Canadian (approximately $63,912 US).
- We issued 6,666,667 shares of our Common Stock for the purchase of laboratory equipment valued at $22,000.
- We received monies in exchange for a convertible note payable having a face value of $65,000. In connection with this note, 100,000,000 shares of our Common Stock have been reserved for issuance.

INFLATION

Although our operations are influenced by general economic conditions, we do not believe that inflation had a material effect on our results of operations during the three month period ended March 31, 2017.

CRITICAL ACCOUNTING ESTIMATES

The discussion and analysis of our financial condition and results of operations are based upon our consolidated financial statements, which have been prepared in accordance with accounting principles generally accepted in the United States. The preparation of these consolidated financial statements requires us to make estimates and judgments that affect the amounts of assets, liabilities, revenues and expenses, and related disclosure of contingent assets and liabilities. On an on-going basis, we evaluate our estimates based on historical experience and on various other assumptions that are believed to be reasonable under the circumstances, the results of which form the basis for making judgments about the carrying values of assets and liabilities that are not readily apparent from other sources. Actual results may differ from these estimates under different assumptions or conditions. The following represents a summary of our critical accounting policies, defined as those policies that we believe are the most important to the portrayal of our financial condition and results of operations and that require management's most difficult, subjective or complex judgments, often as a result of the need to make estimates about the effects of matters that are inherently uncertain.

ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK.

We are a smaller reporting company and are not required to provide the information under this item pursuant to Regulation S-K.

ITEM 4. CONTROLS AND PROCEDURES.

Disclosure Controls and Procedures – Our management, with the participation of our Chief Executive Officer and Chief Financial Officer, has evaluated the effectiveness of our disclosure controls and procedures (as such term is defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, as amended (the "Exchange Act") as of the end of the period covered by this Report.

These controls are designed to ensure that information required to be disclosed in the reports we file or submit pursuant to the Securities Exchange Act of 1934 is recorded, processed, summarized and reported within the time periods specified in the rules and forms of the Securities and Exchange Commission, and that such information is accumulated and communicated to our management, including our CEO/CFO to allow timely decisions regarding required disclosure.

Based on this evaluation, our CEO and CFO have concluded that our disclosure controls and procedures were not effective as of March 31, 2017, at reasonable assurance level, for the following reasons:

- Ineffective control environment and lack of qualified full-time CFO who has SEC experience to focus on our financial affairs;
- Lack of qualified and sufficient personnel, and processes to adequately and timely identify making any and all required public disclosures;
- Deficiencies in the period-end reporting process and accounting policies;
- Inadequate internal controls over the application of new accounting principles or the application of existing accounting principles to new transactions;
- Inadequate internal controls relating to the authorization, recognition, capture, and review of transactions, facts, circumstances, and events that could have a material impact on the company's financial reporting process;
- Deficient revenue recognition policies;
- Inadequate internal controls with respect to inventory transactions; and
- Improper and lack of timely accounting for accruals such as prepaid expenses, accounts payable and accrued liabilities.

Our Board of Directors has assigned a priority to the short-term and long-term improvement of our internal control over financial reporting. We are reviewing various potential solutions to remedy the processes that would eliminate the issues that may arise due to the absence of separation of duties within the financial reporting functions. Additionally, the Board of Directors will work with management to continuously review controls and procedures to identified deficiencies and implement remediation within our internal controls over financial reporting and our disclosure controls and procedures.

We believe that our financial statements presented in this annual report on Form 10-K fairly present, in all material respects, our financial position, results of operations, and cash flows for all periods presented herein.

Inherent Limitations – Our management, including our Chief Executive Officer and Chief Financial Officer, does not expect that our disclosure controls and procedures will prevent all error and all fraud. A control system, no matter how well conceived and operated, can provide only reasonable, not absolute, assurance that the objectives of the control system are met. The design of any system of controls is based in part upon certain assumptions about the likelihood of future events, and there can be no assurance that any design will succeed in achieving its stated goals under all potential future conditions. Further, the design of a control system must reflect the fact that there are resource constraints, and the benefits of controls must be considered relative to their costs. Because of the inherent limitations in all control systems, no evaluation of controls can provide absolute assurance that all control issues and instances of fraud, if any, within our company have been detected. These inherent limitations include the realities that judgments in decision-making can be faulty, and that breakdown can occur because of simple error or mistake. In particular, many of our current processes rely upon manual reviews and processes to ensure that neither human error nor system weakness has resulted in erroneous reporting of financial data.

Changes in Internal Control over Financial Reporting – There were no changes in our internal control over financial reporting during our fiscal year ended December 31, 2016, which were identified in conjunction with management's evaluation required by paragraph (d) of Rules 13a-15 and 15d-15 under the Exchange Act, that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

PART II. OTHER INFORMATION

ITEM 1. LEGAL PROCEEDINGS

Other than the matter discussed above under "GMP Manufacturing" we are not party to any material legal proceedings, nor have any such actions been threatened against us.

ITEM 1A. RISK FACTORS

We are a smaller reporting company and are not required to provide the information under this item pursuant to Regulation S-K.

ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

During the three months ended March 31, 2017, we issued a total of 48,428,125 shares of our Common Stock. Of these 42,428,125 shares valued at $128,451 were issued upon conversion of outstanding notes payable, reducing the debt by $48,500 and interest payable by $3,022 and generating a loss on conversion of $76,929.

In addition, we issued 6,000,000 shares of our Common Stock valued at $14,400 for $15,000 Canadian ($11,278 US). The $15,000 Canadian was paid directly to a firm which was engaged to conduct a valuation of our patents.

Subsequent Events

In April 2017 we engaged in the following transactions:

- We received monies in exchange for a convertible note payable having a face value of $100,000 Canadian (approximately $75,190 US).
- We sold 34,000,000 shares of our Common Stock valued at $98,600 for $85,000 Canadian (approximately $63,912 US).
- We issued 6,666,667 shares of our Common Stock for the purchase of laboratory equipment valued at $22,000.
- We received monies in exchange for a convertible note payable having a face value of $65,000.In connection with this note, 100,000,000 shares of our Common Stock have been reserved for issuance.

The funds obtained from these transactions were used for working capital, including the development of our new business described above under “Plan of Operation.”

ITEM 3. DEFAULTS UPON SENIOR SECURITIES

None

ITEM 4. MINE SAFETY DISCLOSURE

Not Applicable

ITEM 5. OTHER INFORMATION

None

ITEM 6. EXHIBITS

Exhibit No.	Description
31.1	Certification of Chief Executive Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
31.2	Certification of Chief Financial Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
32	Certification of Chief Executive Officer and Chief Financial Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002
101.INS	XBRL Instance Document*
101.SCH	XBRL Schema Document*
101.CAL	XBRL Calculation Linkbase Document*
101.DEF	XBRL Definition Linkbase Document*
101.LAB	XBRL Label Linkbase Document*
101.PRE	XBRL Presentation Linkbase Document*

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* Pursuant to Rule 406T of Regulation S-T, these interactive data files are not deemed filed or part of a registration statement or prospectus for purposes of Section 11 or 12 of the Securities Act or Section 18 of the Securities Exchange Act and otherwise not subject to liability.

SIGNATURES

Pursuant to the requirements of Section 12 of the Securities and Exchange Act of 1934, the Registrant has duly caused this Report to be signed on its behalf by the undersigned, thereunto duly authorized on May 12, 2017.

SUNSHINE BIOPHARMA, INC.

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

By: s/ Camille Sebaaly
Camille Sebaaly,
Principal Financial Officer and
Principal Accounting Officer

EXHIBIT 31.1

CERTIFICATION PURSUANT TO 18 USC, SECTION 1350, AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES OXLEY ACT OF 2002

I, Steve N. Slilaty, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Sunshine Biopharma, Inc.;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary in order to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal controls over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

   a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedure to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

   b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

   c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based upon such evaluation; and

   d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

   a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

   b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Dated: May 12, 2017

/s/ Steve N. Slilaty
Steve N. Slilaty, Chief Executive Officer

EXHIBIT 31.2

CERTIFICATION PURSUANT TO 18 USC, SECTION 1350, AS ADOPTED PURSUANT TO SECTION 302 OF THE SARBANES OXLEY ACT OF 2002

I, Camille Sebaaly, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Sunshine Biopharma, Inc.;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary in order to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal controls over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

   a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedure to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

   b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

   c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based upon such evaluation; and

   d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

   a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

   b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Dated: May 12, 2017

/s/ Camille Sebaaly
Camille Sebaaly, Chief Financial Officer

EXHIBIT 32

CERTIFICATION PURSUANT TO 18 USC, SECTION 1350, AS ADOPTED PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

In connection with this quarterly report of Sunshine Biopharma, Inc. (the "Company") on Form 10-Q for the nine month period ended September 30, 2016, as filed with the Securities and Exchange Commission on May 12, 2017 (the "Report"), we, the undersigned, in the capacities and on the date indicated below, certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that to the best of our knowledge:

1. The Report fully complies with the requirements of Rule 13(a) or 15(d) of the Securities Exchange Act of 1934; and

2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Dated: May 12, 2017

/s/ Steve N. Slilaty
Steve N. Slilaty, Chief Executive Officer

Dated: May 12, 2017

/s/ Camille Sebaaly
Camille Sebaaly, Chief Financial Officer