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

FORM 10-Q

☑ QUARTERLY REPORT PURSUAN	NT TO SECTION 13 OR 15	5(D) OF THE SECURITIES EXCH	IANGE ACT OF 1934
Fo	or the quarterly period ende	d March 31, 2017	
☐ TRANSITION REPORT PURSUAN	NT TO SECTION 13 OR 1:	5(d) OF THE SECURITIES EXCH	ANGE ACT OF 1934
F	For the transition period from	n to	
	Commission File Numb	er 000-55334	
(Exa	COHBAR,		
Delaware		26-12999	52
(State or other jurisdiction o incorporation or organization		(I.R.S. Empl Identification N	oyer 'umber)
(Ad	1455 Adams Drive, S Menlo Park, CA Idress of principal executive	94025	
(Regi	(650) 446-78istrant's telephone number,		
Indicate by check mark whether the reging Exchange Act of 1934 during the preceding reports). Yes ☑ No □			
Indicate by check mark whether the reg interactive Data File required to be submitted a preceding 12 months (or for such shorter period	and posted pursuant to Rul	e 405 of Regulation S-T (§ 232.40	05 of this chapter) during the
Indicate by check mark whether the regi- reporting company, or an emerging growth com- company," and "emerging growth company" in	pany. See the definitions of	"large accelerated filer," "accelera	
Large accelerated filer □ Accelerated filer □	Non-accelerated filer ☐ (Do not check if a smaller reporting company)	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.
As of May 9, 2017 the registrant had outst	anding 35,857,701 shares o	f common stock.	

COHBAR, INC. FORM 10-Q For the Quarterly Period Ended March 31, 2017

		Page Number
	PART I – FINANCIAL INFORMATION	
Item 1	Financial Statements	1
Item 2	Management's Discussion and Analysis of Financial Condition and Results of Operations	10
Item 4	Controls and Procedures	13
	PART II – OTHER INFORMATION	
Item 1	Legal Proceedings	14
Item 2	Unregistered Sales of Equity Securities and Use of Proceeds	14
Item 3	Defaults Upon Senior Securities	14
Item 4	Mine Safety Disclosures	14
Item 5	Other Information	14
Item 6	Exhibits	15
	SIGNATURES	16

PART I. FINANCIAL INFORMATION

Item 1. Financial Statements

CohBar, Inc. Condensed Balance Sheets

	As of			
	1	March 31, 2017		ecember 31, 2016
	(1	unaudited)		
ASSETS				
Current assets:				
Cash	\$, ,	\$	3,257,458
Investments		6,323,323		5,428,962
Subscription receivable		- 		522,326
Prepaid expenses and other current assets		172,005		110,822
Total current assets		9,362,382		9,319,568
Property and equipment, net		214,523		230,512
Other assets		36,810		36,810
Total assets	\$	9,613,715	\$	9,586,890
	_			
LIABILITIES AND STOCKHOLDERS' EQUITY				
Current liabilities:				
Accounts payable	\$	384,525	\$	103,294
Accrued liabilities		161,467		132,780
Accrued payroll and other compensation		182,224		447,641
Note payable, net of debt discount of \$0 and \$59 as of March 31, 2017 and December 31, 2016,				
respectively		102,630		205,201
Total liabilities		830,846		888,916
Commitments and contingencies				
Stockholders' equity:				
Preferred stock, \$0.001 par value, Authorized 5,000,000 shares;				
No shares issued and outstanding as of March 31, 2017 and December 31, 2016, respectively		-		-
Common stock, \$0.001 par value, Authorized 75,000,000 shares;				
Issued and outstanding 35,857,701 shares as of March 31, 2017 and 34,807,881 as of December 31,				
2016		35,858		34,808
Additional paid-in capital		25,388,657		23,072,702
Accumulated deficit		(16,641,646)		(14,409,536)
Total stockholders' equity		8,782,869		8,697,974
Total liabilities and stockholders' equity	\$	9,613,715	\$	9,586,890

The accompanying notes are an integral part of these condensed financial statements

CohBar, Inc. Condensed Statements of Operations (unaudited)

	For The Thr Ended Ma	
	2017	2016
Revenues	<u>\$</u> -	\$ -
Operating expenses:		
Research and development	1,292,780	737,100
General and administrative	940,089	479,932
Total operating expenses	2,232,869	1,217,032
Operating loss	(2,232,869)	(1,217,032)
Other income (expense):		
Interest income	2,163	2,656
Interest expense	(1,345)	(1,875)
Amortization of debt discount	(59)	(49)
Total other income (expense)	759	732
Net loss	\$ (2,232,110)	\$ (1,216,300)
Basic and diluted net loss per share	\$ (0.06)	\$ (0.04)
Weighted average common shares outstanding - basic and diluted	35,788,158	32,331,345

The accompanying notes are an integral part of these condensed financial statements

CohBar, Inc. Condensed Statements of Cash Flows (unaudited)

		For The Three Months Ended March 31,		
	2017	2016		
Cash flows from operating activities:				
Net loss	\$ (2,232,110)	\$ (1,216,300)		
Adjustments to reconcile net loss to net cash used in operating activities:				
Depreciation	15,989	12,273		
Stock-based compensation	414,513	123,646		
Amortization of debt discount	59	49		
Changes in operating assets and liabilities:				
Prepaid expenses and other current assets	(61,183)			
Accounts payable	281,231	90,975		
Accrued liabilities	28,687	(73,016)		
Accrued payroll and other compensation	(265,417)	(94,072)		
Net cash used in operating activities	(1,818,231)	(1,191,935)		
Cash flows from investing activities:				
Purchases of property and equipment	-	(8,422)		
Payment for security deposit	-	(2,152)		
Purchases of investments	(5,327,361)	(4,008,518)		
Proceeds from redemptions of investments	4,433,000	4,742,000		
Net cash (used in) provided by investing activities	(894,361)	722,908		
Cash flows from financing activities:				
Proceeds from exercise of warrants	2,404,993	-		
Repayment of note payable	(102,630)	-		
Proceeds from exercise of compensation options	-	6,650		
Proceeds from exercise of employee stock options	19,825	2,600		
Net cash provided by financing activities	2,322,188	9,250		
Too onship promised by mannering non-times	2,322,100	7,230		
Net decrease in cash	(390,404)	(459,777)		
Cash at beginning of period	3,257,458	4,803,687		
Cash at end of period	\$ 2,867,054	\$ 4,343,910		
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Supplemental disclosure of cash flow information:				
Income taxes paid	\$ -	\$ 1,300		
Interest paid	\$ 12,418	\$ -		

The accompanying notes are an integral part of these condensed financial statements

Note 1 - Business Organization

CohBar, Inc. ("CohBar" or the "Company") is an innovative biotechnology company and a leader in the research and development of mitochondria based therapeutics (MBTs), an emerging class of drugs with the potential to treat a wide range of diseases associated with aging and metabolic dysfunction, including obesity, fatty liver disease (NAFLD) and non-alcoholic steatohepatitis (NASH), type 2 diabetes mellitus (T2D), cancer, atherosclerosis, cardiovascular disease, and neurodegenerative diseases such as Alzheimer's.

The Company's primary activities include research and development of its MBT pipeline, securing intellectual property protection, managing collaborations with contract research organizations ("CROs") and academic institutions, expanding its scientific leadership and raising capital. To date, the Company has not generated any revenues from operations and does not expect to generate any revenues in the near future. The Company has financed its operations primarily through proceeds from its initial public offering ("IPO") and concurrent private offering, private placements of its preferred stock, the exercise of outstanding warrants and stock options and, to a lesser extent, from a grant from a research foundation.

The unaudited interim condensed financial statements of the Company have been prepared in accordance with United States generally accepted accounting principles ("U.S. GAAP") for interim financial information and the rules and regulations of the Securities and Exchange Commission ("SEC"). They do not include all information and footnotes required by U.S. GAAP for complete financial statements. Except as disclosed herein, there have been no material changes in the information disclosed in the notes to the financial statements for the year ended December 31, 2016, included in the Company's Annual Report on Form 10-K (the "2016 Form 10-K"), filed with the SEC on March 31, 2017. The interim unaudited condensed financial statements should be read in conjunction with those audited financial statements included in the 2016 Form 10-K. In the opinion of management, all adjustments considered necessary for fair presentation, consisting solely of normal recurring adjustments, have been made. Operating results for the three months ended March 31, 2017 are not necessarily indicative of the results that may be expected for the year ending December 31, 2017, or any other period.

Note 2 - Management's Liquidity Plans

As of March 31, 2017, the Company had working capital and stockholders' equity of \$8,531,536 and \$8,782,869 respectively. During the three months ended March 31, 2017, the Company incurred a net loss of \$2,232,110. The Company has not generated any revenues, has incurred net losses since inception and does not expect to generate revenues in the near term.

Based on current budget assumptions and with the cash and investments on hand as of March 31, 2017, the Company believes that it has sufficient capital to meet its operating expenses and obligations for the next twelve months from the date of this filing. However, if unanticipated difficulties or circumstances arise the Company may require additional capital sooner to support its operations. If the Company is unable to raise additional capital whenever necessary it may be forced to decelerate or curtail its research and development activities and delay planned FDA filings and clinical activities until such time as additional capital becomes available. Such limitation of the Company's activities would allow the Company to slow its rate of spending and extend its use of cash until additional capital is raised. There can be no assurance that such a plan will be successful. There is no assurance that additional financing will be available when needed or that the Company will be able to obtain such financing on reasonable terms.

Note 3 - Summary of Significant Accounting Policies

Basis of Presentation

All amounts are presented in U.S. Dollars.

Use of Estimates

The preparation of financial statements in conformity with U.S. GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent liabilities at dates of the financial statements and the reported amounts of revenue and expenses during the periods. Actual results could differ from these estimates. The Company's significant estimates and assumptions include the fair value of financial instruments, stock-based compensation and the valuation allowance relating to the Company's deferred tax assets.

INVESTMENTS

Investments consist of U.S. Treasury Bills of \$4,331,283, which are classified as held-to-maturity, and Certificates of Deposit of \$1,992,040. The Company determines the appropriate balance sheet classification of its investments at the time of purchase and evaluates the classification at each balance sheet date. All of the Company's U.S. Treasury Bills and Certificates of Deposit mature within the next twelve months. Unrealized gains and losses are *de minimis*. As of March 31, 2017, the carrying value of the Company's U.S. Treasury Bills approximates their fair value, due to their short-term maturities.

SHARE-BASED PAYMENT

The Company accounts for share-based payments using the fair value method. For employees and directors, the fair value of the award is measured, as discussed below, on the grant date. For non-employees, fair value is generally valued based on the fair value of the services provided or the fair value of the equity instruments on the measurement date, whichever is more readily determinable, and remeasured on each financial reporting date until the service is complete. The Company has granted stock options at exercise prices equal to the higher of (i) the closing price of the Company's common stock as reported on the OTCQX marketplace or (ii) the closing price of the Company's common stock as reported by the TSX Venture Exchange on the date of grant.

The weighted-average fair value of options and warrants has been estimated on the date of grant using the Black-Scholes pricing model. The fair value of each instrument is estimated on the date of grant utilizing certain assumptions for a risk free interest rate, volatility and expected remaining lives of the awards. Since the Company has a limited history of being publicly traded, the fair value of stock-based payment awards issued was estimated using a volatility derived from an index of comparable entities. The assumptions used in calculating the fair value of share-based payment awards represent management's best estimates, but these estimates involve inherent uncertainties and the application of management judgment. As a result, if factors change and the Company uses different assumptions, the Company's stock-based compensation expense could be materially different in the future. In addition, the Company is required to estimate the expected forfeiture rate and only recognize expense for those shares expected to vest. In estimating the Company's forfeiture rate, the Company analyzed its historical forfeiture rate, the remaining lives of unvested options, and the number of vested options as a percentage of total options outstanding. If the Company's actual forfeiture rate is materially different from its estimate, or if the Company reevaluates the forfeiture rate in the future, the stock-based compensation expense could be significantly different from what the Company has recorded in the current period.

Note 3 - Summary of Significant Accounting Policies (continued)

The weighted-average Black-Scholes assumptions are as follows:

Ended March 31, 2017 2016 Expected life 8 years 6 years Risk free interest rate 2.16% 1.27% Expected volatility 79% 79% Expected dividend yield 0% 0% Forfeiture rate 0% 0%

For the Three Months

As of March 31, 2017, total unrecognized stock option compensation expense is \$1,821,693, which will be recognized as those options vest over a period of approximately four years. The amount of future stock option compensation expense could be affected by any future option grants or by any option holders leaving the Company before their grants are fully vested.

NET LOSS PER SHARE OF COMMON STOCK

Basic net loss per share is computed by dividing net loss attributable to common stockholders by the weighted average number of common shares outstanding during the period. Diluted net earnings per share reflects the potential dilution that could occur if securities or other instruments to issue common stock were exercised or converted into common stock. Potentially dilutive securities are excluded from the computation of diluted net loss per share as their inclusion would be anti-dilutive and consist of the following:

	As of Ma	rch 31,
	2017	2016
Options	5,618,497	5,376,995
Warrants	951,635	8,064,716
Totals	6,570,132	13,441,711

Note 4 - Accrued Liabilities

Accrued liabilities consist of:

	M	As of arch 31, 2017	Dec	As of cember 31, 2016
Lab services & supplies	\$	100,225	\$	87,100
Professional fees		46,340		17,760
Consultant fees		2,500		2,500
Interest		12,402		25,420
Total accrued liabilities	\$	161,467	\$	132,780

Note 5 - Commitments and Contingencies

LITIGATIONS, CLAIMS AND ASSESSMENTS

The Company may from time to time be party to litigation and subject to claims incident to the ordinary course of business. As the Company grows and gains prominence in the marketplace it may become party to an increasing number of litigation matters and claims. The outcome of litigation and claims cannot be predicted with certainty, and the resolution of these matters could materially affect the Company's future results of operations, cash flows or financial position. The Company is not currently a party to any legal proceedings.

OPERATING LEASE

The Company is a party to a lease agreement for a shared laboratory facility leased on a month-to-month basis in Menlo Park, California.

Rent expense was \$55,215 and \$34,396 for the three months ended March 31, 2017 and 2016, respectively.

Note 6 - Stockholders' Equity

STOCK OPTIONS

The Company has an incentive stock plan, the Amended and Restated 2011 Equity Incentive Plan (the "2011 Plan"), and has granted stock options to employees, non-employee directors and consultants from the 2011 Plan. Options granted under the 2011 Plan may be Incentive Stock Options or Non-statutory Stock Options, as determined by the Administrator at the time of grant. In January 2016 the Company issued a warrant to purchase 125,000 shares of the Company's common stock to an investor relations firm as partial compensation for consulting services it will provide the Company over a two year period. Pursuant to applicable policies of the TSX-V, the shares issuable under the warrant will be counted against the limit of shares authorized for issuance under the 2011 Plan, notwithstanding that the warrant was not issued under the 2011 Plan. After giving effect to this limitation there are 683,322 shares remaining available for issuance under the 2011 Plan at March 31, 2017.

During the three months ended March 31, 2017, the Company granted stock options to employees to purchase 1,031,000 shares of the Company's common stock at \$2.40 per share. Of the 1,031,000 stock options granted, 300,000 are subject to vesting based on continuous service over periods between zero and four years from the date of grant and have terms of ten years. The balance of the total grant, or 731,000, is performance based and will be valued at the time the milestones are reached. The stock options have an aggregate grant date fair value of \$528,580.

In February 2017, 16,250 stock options were exercised for cash proceeds of \$19,825 and the Company cancelled 48,750 stock options.

The Company recorded \$414,513 and \$123,646 of stock based compensation in the three months ended March 31, 2017 and 2016, respectively.

Note 6 - Stockholders' Equity (continued)

The following table represents stock option activity for the three months ended March 31, 2017:

				Weighted Average								
	Stock O	ptions		Exercis	e Pri	ce	r Value	Contractual	Intrinsic			
	Outstanding	Exercisable	Outs	standing	Exe	Exercisable		Vested Life (Yea		Value		
Balance – December 31, 2016	4,652,497	1,908,883	\$	0.92	\$	0.41	\$	0.41	8.24	\$ -		
Granted	1,031,000	100,000		2.40		2.40		6.25	6.25	-		
Exercised	(16,250)	-		-		-		-	-	-		
Cancelled	(48,750)	-		-		-		-	-	-		
Balance – March 31, 2017	5,618,497	2,511,008	\$	1.01	\$	0.58	\$	0.58	6.72	\$4,034,869		

The following table summarizes information on stock options outstanding and exercisable as of March 31, 2017:

	Exercise Price	Number Outstanding	Weighted Average Remaining Contractual Term	 Weighted Average Exercise Price	Number Exercisable	_	Weighted Average Exercise Price
\$	0.05	72,876	5.01 years	\$ 0.05	72,876	\$	0.05
\$	0.26	1,024,810	7.03 years	\$ 0.26	948,901	\$	0.26
\$	0.73	1,475,687	7.63 years	\$ 0.73	860,817	\$	0.73
\$	1.00	313,124	8.31 years	\$ 1.00	146,457	\$	1.00
\$	1.10	10,000	8.78 years	\$ 1.10	3,333	\$	1.10
\$	1.17	70,000	8.62 years	\$ 1.17	27,500	\$	1.17
\$	1.22	125,000	8.85 years	\$ 1.22	36,458	\$	1.22
\$	1.50	40,000	8.92 years	\$ 1.50	10,833	\$	1.50
\$	1.55	1,456,000	8.94 years	\$ 1.55	283,000	\$	1.55
\$	2.40	1,031,000	9.84 years	\$ 2.40	120,833	\$	2.40
Totals	1	5,618,497			2,511,008		

WARRANTS

In January 2017, a total of 926,588 warrants were exercised for cash proceeds of \$1,853,176. An additional \$522,326 was received in January 2017 which was the proceeds from warrants exercised in December 2016. During the three months ended March 31, 2017, 4,695,846 warrants expired.

In January and February 2017, consultants to the Company exercised a total of 106,982 warrants for cash proceeds of \$29,491.

As of March 31, 2017, the Company has 951,635 warrants outstanding and exercisable to purchase common stock. Such warrants have a weighted average exercise price of \$0.39, a weighted average remaining contractual life of 6.25 years and an aggregate intrinsic value of \$1,339,531.

Note 7 - Related Party Transactions

Two of the Company's directors, Pinchas Cohen and Nir Barzilai, provide consulting services to the Company pursuant to agreements that provide for annual compensation to each director of \$42,000. Each agreement provides for an annual service term and can be extended by mutual consent of both parties. The service terms under the agreements expired in 2015. The Company continues to compensate Dr. Cohen and Dr. Barzilai for their ongoing services under the terms of the original agreements. Payments of \$10,500 were made to each Director during each of the three months ended March 31, 2017 and 2016. As of March 31, 2017 and December 31, 2016, no amounts were owed to either Director.

Note 8 - Subsequent Events

Management has evaluated subsequent events to determine if events or transactions occurring through the date on which the condensed financial statements were issued require adjustment or disclosure in the Company's condensed financial statements.

Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations

The following discussion and analysis is based upon our financial statements as of the dates and for the periods presented in this section. You should read this discussion and analysis in conjunction with the financial statements and notes thereto found in Part I, Item 1 of this Form 10-Q and our financial statements and notes thereto included in our Annual Report on Form 10-K for the year ended December 31, 2016 (the "2016 Form 10-K"). All references to the first quarter and first three months of 2017 and 2016 mean the three-month periods ended March 31, 2017 and 2016, respectively. Unless the context otherwise requires, "CohBar," "we," "us" and "our" refer to CohBar, Inc.

Special Note Regarding Forward-Looking Statements

This report, including the "Management's Discussion and Analysis of Financial Condition and Results of Operations," contains forward-looking statements regarding future events and our future results that are based on our current expectations, estimates, forecasts, and projections about our business, our potential drug candidates, our capital resources and ability to fund our operations, our results of operations, the industry in which we operate and the beliefs and assumptions of our management. Words such as "expect," "anticipate," "target," "goal," "project," "would," "could," "intend," "plan," "believe," "seek" and "estimate," variations of these words, and similar expressions are intended to identify those forward-looking statements. These forward-looking statements are only predictions and are subject to risks, uncertainties and assumptions that are difficult to predict. Therefore, actual results may differ materially from those expressed in any forward-looking statements. Factors that might cause or contribute to such differences include, but are not limited to, those discussed in this report under the section entitled "Risk Factors" in Item 1A of Part I of the 2016 Form 10-K, as supplemented or modified in our quarterly reports on Form 10-Q. We undertake no obligation to revise or update publicly any forward-looking statements for any reason, whether as a result of new information, future events or otherwise, except as may be required by law.

Overview

We are an innovative biotechnology company and a leader in the research and development of mitochondria based therapeutics (MBTs), an emerging class of drugs with the potential to treat a wide range of diseases associated with aging and metabolic dysfunction, including obesity, fatty liver disease (NAFLD) and non-alcoholic steatohepatitis (NASH), type 2 diabetes mellitus (T2D), cancer, atherosclerosis, cardiovascular disease, and neurodegenerative diseases such as Alzheimer's.

MBTs originate from almost two decades of research by our founders, resulting in their discovery of a novel group of peptides called mitochondrial-derived peptides (MDPs) encoded within the genome of mitochondria, the powerhouses of the cell. These naturally occurring MDPs and related analogs have demonstrated a range of biological activity and therapeutic potential in pre-clinical models across multiple diseases associated with aging.

We believe CohBar is a first mover in exploring the mitochondrial genome for therapeutically relevant peptides, and have developed a proprietary MBT technology platform which uses cell based assays and animal models of disease to rapidly identify mitochondrial peptides with promising biological activity. Once identified, we deploy optimization techniques to improve the drug-like properties of our MBT candidates, enabling us to match the most biologically promising peptides to disease indications that have substantial unmet medical needs.

In September 2016, we advanced two novel, optimized analogs of our MOTS-c MDP, CB4209 and CB4211, into IND-enabling studies as our lead MBT drug candidates with potential for treatment of fatty liver disease (NAFLD), Nonalcoholic steatohepatitis (NASH), obesity, and type 2 diabetes (T2D). To date, our founders and scientific team have discovered a large number of MDPs that have demonstrated a range of biological activities and therapeutic potential. Our ongoing research and development of our pipeline MDPs is focused on identifying and advancing novel improved analogs of those MDPs that have the greatest therapeutic and commercial potential for development into drugs.

We have financed our operations primarily through proceeds from our IPO and concurrent private offering, private placements of our preferred stock, the exercise of outstanding warrants and stock options and, to a lesser extent, from a grant from a research foundation. Since our inception through December 31, 2016, our operations have been funded with an aggregate of approximately \$25.8 million, of which approximately \$0.2 million was from a grant-funding organization and approximately \$25.6 million was from the issuance of equity instruments.

Since inception, we have incurred significant operating losses. Our net losses were \$2,232,110 and \$1,216,300 for the three months ended March 31, 2017 and 2016, respectively. As of March 31, 2017, we had an accumulated deficit of \$16,641,646. Our net losses may fluctuate significantly from quarter to quarter and from year to year. We expect to continue to incur significant expenses and operating losses as we continue research, discovery and pre-clinical development efforts on our current MBT candidates and any other MDPs, expand and protect our intellectual property portfolio, and hire additional development and scientific personnel. Our MBT drug target candidates are in early stages of investigational research. Candidates in later stages of clinical development generally have higher development costs than those in earlier stages of clinical development, primarily due to the increased size and duration of later-stage clinical trials. We expect research and development costs to increase significantly for the foreseeable future as our product candidate development programs progress.

Based on current budget assumptions and with the cash and investments on hand as of March 31, 2017, we believe that we have sufficient capital to meet our operating expenses and obligations for the next twelve months from the date of this filing. However, if unanticipated difficulties or circumstances arise we may require additional capital sooner to support our operations. If we are unable to raise additional capital whenever necessary we may be forced to decelerate or curtail our research and development activities and delay planned FDA filings and clinical activities until such time as additional capital becomes available. Such limitation of our activities would allow us to slow our rate of spending and extend our use of cash until additional capital is raised. There can be no assurance that such a plan will be successful. There is no assurance that additional financing will be available when needed or that we will be able to obtain such financing on reasonable terms.

Financial Operations Review

Revenue

To date, we have not generated any revenue from product sales and do not expect to generate any revenue from the sale of products in the near future. In the future, we will seek to generate revenue from product sales, either directly or under any future licensing, development or similar relationship with a strategic partner.

Research and development expenses

Research and development expenses consist primarily of costs incurred for our research activities, including our drug discovery efforts, and the development of our product candidates, which include:

- employee-related expenses including salaries, benefits, and stock-based compensation expense;
- expenses incurred under agreements with third parties, including contract research organizations, or CROs, that conduct research and development and pre-clinical activities on our behalf, and the cost of consultants;
- the cost of laboratory equipment, supplies and manufacturing MBT test materials; and
- depreciation and other personnel-related costs associated with research and product development.

We expense all research and development expenses as incurred. We expect our research and development expenses to increase in the year ending December 31, 2017, as we continue our efforts to advance our lead MBT candidate program and to discover, evaluate and optimize other MDPs as potential MBT drug candidates.

General and administrative expenses

General and administrative expenses consist primarily of salaries and other related costs, including stock-based compensation, for personnel in executive, finance and administrative functions. Other significant costs include legal fees relating to patent and corporate matters and fees for accounting and consulting services. We anticipate that our general and administrative expenses will increase in the future to support continued research and development activities and the potential commercialization of our product candidates. These increases will likely include increased costs related to the hiring of additional personnel, and fees to outside consultants, lawyers and accountants, among other expenses.

Results of Operations

The following table sets forth our results of operations for the periods presented. The period-to-period comparison of financial results is not necessarily indicative of financial results to be achieved in future periods.

For	The Three Mon	ths Ended Marci	n 31.
2017	2016	\$	%

Operating expenses:

Research and development General and administrative	\$ 1,292,780	\$ 737,100 479,932	\$ 555,680 460,157	75%
Total operating expenses	\$ 2,232,869	\$ 1,217,032	\$ 1,015,837	83%

Comparison of Three Months Ended March 31, 2017 and 2016

Research and development expenses were \$1,292,780 in the three months ended March 31, 2017 compared to \$737,100 in the prior year period, a \$555,680 increase. The increase in research and development expenses was primarily due to the \$423,524 incurred which related to our IND-enabling activities, \$46,328 increase in laboratory supplies related to our efforts to develop additional optimized MBT candidates, a \$44,697 increase in expense primarily associated with our Scientific Advisory Board meeting held in January 2017 and a \$20,819 increase in rent associated with the expanded laboratory facility in Menlo Park. We expect research and development expenses to increase in the coming quarters as we continue to advance our lead MBT candidate program and evaluate and optimize other MDPs as potential MBT drug candidates.

General and administrative expenses were \$940,089 in the three months ended March 31, 2017 compared to \$479,932 in the prior year period, a \$460,157 increase. The increase was primarily due to a \$412,410 increase in wages and stock based compensation due to the increase in head count with a full quarter's expense associated with our CEO and Director of Investor Relations and the expensing of the additional stock options grants during the current year quarter.

Liquidity and Capital Resources

As of March 31, 2017 we had a cash balance of \$2,867,054. We maintain our cash in a checking and savings account on deposit with a banking institution in the United States. We also maintain a portfolio of short-term highly liquid securities investing in U.S. Treasury Bills and Certificate of Deposits. As of March 31, 2017, we had an investments balance of \$6,323,323.

Cash Flows from Operating Activities

Net cash used in operating activities for the three months ended March 31, 2017 and 2016 was \$1,818,231 and \$1,191,935, respectively. The cash used in operations for the three months ended March 31, 2017 was primarily due to our reported net loss of \$2,232,110 offset by \$430,561 in stock based compensation, depreciation and amortization expense in the current year quarter. The cash used in operations for the three months ended March 31, 2016 was primarily due to our reported net loss of \$1,216,300 and a \$94,072 decrease in accrued payroll as bonuses recognized at the end of the previous year were paid in the current year quarter, offset by \$123,646 in stock based compensation.

Cash Flows from Investing Activities

Net cash used in investing activities in the three months ended March 31, 2017 was \$894,361 which was due to the purchases of our investments in certificates of deposit and treasury bills as compared to the timing of maturities of those investments. The cash used provided by investing activities of \$722,908 in the three months ended March 31, 2016 was primarily due to the timing of maturities of our investments when compared to the purchases of those investments.

Cash Flows from Financing Activities

Net cash provided by financing activities in the three months ended March 31, 2017 and 2016 was \$2,322,188 and \$9,250, respectively. Cash provided by financing activities in the three months ended March 31, 2017 was due to the exercise of warrants and employee stock options, partially offset by payment to the Alzheimer's Drug Development Foundation of \$102,630, representing the first installment of a note obligation associated with a grant from that organization. Cash provided by financing activities of \$9,250 in the three months ended March 31, 2016 was due to the exercise of employee and Compensation Options.

Contractual Obligations

The Company is a party to a lease agreement for a shared laboratory facility leased on a month-to-month basis in Menlo Park, California.

Rent expense was \$55,215 and \$34,396 for the three months ended March 31, 2017 and 2016, respectively.

Item 4. Evaluation of Disclosure Controls and Procedures

As of the end of the period covered by this report, we conducted an evaluation under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, of the effectiveness of the design and operation of our disclosure controls and procedures. The term "disclosure controls and procedures," as defined in Rules 13a-15(e) and 15d-15(e) under the Securities and Exchange Act of 1934, as amended ("Exchange Act"), means controls and other procedures of a company that are designed to ensure that information required to be disclosed by the company in the reports it files or submits under the Exchange Act is recorded, processed, summarized and reported, within the time periods specified in the SEC's rules and forms. Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information required to be disclosed by a company in the reports that it files or submits under the Exchange Act is accumulated and communicated to the company's management, including its principal executive and principal financial officers, or persons performing similar functions, as appropriate, to allow timely decisions regarding required disclosure. Based on this evaluation, our Chief Executive Officer and Chief Financial Officer concluded as of the end of the period covered by this report that our disclosure controls and procedures were not effective due to a material weakness.

The material weakness relates to our having one employee assigned to positions that involve processing financial information, resulting in a lack of segregation of duties so that all journal entries and account reconciliations are reviewed by someone other than the preparer, heightening the risk of error or fraud. If we are unable to remediate the material weakness, or other control deficiencies are identified, we may not be able to report our financial results accurately, prevent fraud or file our periodic reports as a public company in a timely manner. Due to our small size and the early stage of our business, segregation of duties may not always be possible and may not be economically feasible. We have limited capital resources and have given priority in the use of those resources to our research and development efforts. As a result, we have not been able to take steps to improve our internal controls over financial reporting during the quarter ended March 31, 2017. However, we continue to evaluate the effectiveness of internal controls and procedures on an on-going basis. As our operations grow and become more complex, we intend to hire additional personnel in financial reporting and other areas. However, there can be no assurance of when, if ever, we will be able to remediate the identified material weaknesses.

Changes in Internal Control over Financial Reporting

There were no changes in our internal control over financial reporting identified in connection with the evaluation required by Rule 13a-15(d) and 15d-15(d) of the Exchange Act that occurred during the three months ended March 31, 2017 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

We may from time to time be party to litigation and subject to claims incident to the ordinary course of business. As we grow and gain prominence in the marketplace we may become party to an increasing number of litigation matters and claims. The outcome of litigation and claims cannot be predicted with certainty, and the resolution of these matters could materially affect our future results of operations, cash flows or financial position. We are not currently a party to any legal proceedings.

Item 2. Unregistered Sale of Equity Securities and Use of Proceeds

Sales of Unregistered Securities

None.

Use of Proceeds from Registered Securities

There has been no material change in the planned use of proceeds from our IPO as described in our final prospectus filed with the Securities and Exchange Commission on December 16, 2014 pursuant to Rule 424(b). As of March 31, 2017, we have used proceeds from our IPO for working capital and general corporate purposes.

Item 3. Defaults Upon Senior Securities

None.

Item 4. Mine Safety Disclosures

None.

Item 5. Other Information

None.

Item 6. Exhibits

Exhibit

LAIIIOIT	
Number	Description
31.1	Certification of Principal Executive Officer Pursuant to Rule 13-14(a) or Rule 15d-14(a) of the Securities Exchange Act of
	1934 as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
31.2	Certification of Principal Financial Officer Pursuant to Rule 13-14(a) or Rule 15d-14(a) of the Securities Exchange Act of
	1934 as Adopted Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002
32.1	Certification of Principal Executive Officer and Principal Financial Officer Pursuant to 18 U.S.C. Section 1350, as Adopted
	Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002
101.INS	XBRL Instance Document
101.SCH	XBRL Taxonomy Extension Schema Document
101.CAL	XBRL Taxonomy Extension Calculation Linkbase Document
101.DEF	XBRL Taxonomy Extension Definition Linkbase Document
101.LAB	XBRL Taxonomy Extension Label Linkbase Document
101.PRE	XBRL Taxonomy Extension Presentation Linkbase Document

^{*} Management contract or compensatory agreement or arrangement in which our directors or executive officers may participate.

SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized on.

COHBAR, INC.

Date: May 15, 2017

By: /s/ Jeffrey F. Biunno
Jeffrey F. Biunno
Chief Financial Officer, Treasurer and Secretary
(Principal Financial Officer)

CERTIFICATION OF PRINCIPAL EXECUTIVE OFFICER PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Simon Allen, certify that:

- 1. I have reviewed this Quarterly Report on Form 10-Q of CohBar, 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 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 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)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures 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 general 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 on 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 fiscal 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.
- 5. The registrant's other certifying officer 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):
 - 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;
 - 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.

May 15, 2017	By: /s/ Simon Allen
Date	Simon Allen
	Chief Executive Officer
	(Principal Executive Officer)

CERTIFICATION OF PRINCIPAL FINANCIAL OFFICER PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Jeffrey F. Biunno, certify that:

- 1. I have reviewed this Quarterly Report on Form 10-Q of CohBar, 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 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 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)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures 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 general 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 on 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 fiscal 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.
- 5. The registrant's other certifying officer 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):
 - 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;
 - 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.

May 15, 2017	By: /s/ Jeffrey F. Biunno
Date	Jeffrey F. Biunno
	Chief Financial Officer
	(Principal Financial Officer)

CERTIFICATION PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002 (Subsection (a) and (b) of Section 1350, Chapter 63 of Title 18, United States Code)

Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 (Subsections (a) and (b) of Section 1350, Chapter 63 of Title 18, United States Code), the undersigned officers of CohBar, Inc., a Delaware corporation (the "Company"), do hereby certify that:

- 1. To our knowledge, the Quarterly Report on Form 10-Q for the quarter ended March 31, 2017 (the "Form 10-Q") of the Company fully complies with the requirements of Section 13(a) or 15(d) of the Securities Act of 1934; and
- 2. The information contained in the Form 10-Q fairly presents, in all material respects, the financial condition and results of operations of the Company.

May 15, 2017	By: /s/ Simon Allen
Date	Simon Allen
	Chief Executive Officer
	(Principal Executive Officer)
May 15, 2017	By: /s/ Jeffrey F. Biunno
Date	Jeffrey F. Biunno
	Chief Financial Officer
	(Principal Financial and Accounting Officer)