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

FORM 10-Q

☒ **QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the quarterly period ended June 30, 2015

or

☐ **TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the transition period from _____ to _____

Commission File Number 001-33216

OCULUS INNOVATIVE SCIENCES, INC.
(Exact name of registrant as specified in its charter)

Delaware	**68-0423298**
(State or other jurisdiction of incorporation or organization)	(I.R.S Employer Identification No.)

1129 North McDowell Blvd.
Petaluma, CA 94954
(Address of principal executive offices) (Zip Code)

(707) 283-0550
Registrant's telephone number, including area code

Indicate by checkmark 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, or a smaller reporting company. See the definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer	Accelerated filer	Non-accelerated filer	Smaller reporting company
☐	☐	(Do not check if a smaller reporting company) ☐	☒

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

As of August 4, 2015, the number of shares outstanding of the registrant's common stock, $0.0001 par value, was 16,405,665.

OCULUS INNOVATIVE SCIENCES, INC.

Index

	Page
PART I — FINANCIAL INFORMATION	3
Item 1. Financial Statements	3
Condensed Consolidated Balance Sheets	3
Condensed Consolidated Statements of Comprehensive Loss	4
Condensed Consolidated Statements of Cash Flows	5
Notes to Condensed Consolidated Financial Statements	6
Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations	15
Item 3. Quantitative and Qualitative Disclosures About Market Risk	20
Item 4. Controls and Procedures	20
PART II — OTHER INFORMATION	20
Item 1. Legal Proceedings	20
Item 1A. Risk Factors	20
Item 2. Unregistered Sales of Equity Securities and Use of Proceeds	20
Item 3. Defaults Upon Senior Securities	20
Item 4. Mine Safety Disclosures (Not applicable.)	21
Item 5. Other Information	21
Item 6. Exhibits	21

PART I — FINANCIAL INFORMATION

Item 1. Financial Statements

OCULUS INNOVATIVE SCIENCES, INC. AND SUBSIDIARIES
Condensed Consolidated Balance Sheets
(In thousands, except share and per share amounts)

	June 30, 2015	March 31, 2015
	(Unaudited)	
ASSETS		
Current assets:		
Cash and cash equivalents	$ 8,752	$ 6,136
Accounts receivable, net	2,540	1,517
Inventories, net	1,466	1,402
Prepaid expenses and other current assets	506	592
Total current assets	13,264	9,647
Property and equipment, net	863	795
Long-term investment	–	4,538
Other assets	79	68
Total assets	$ 14,206	$ 15,048
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Accounts payable	$ 1,202	$ 932
Accrued expenses and other current liabilities	916	782
Deferred revenue	507	769
Current portion of long-term debt	35	87
Derivative liabilities	70	11
Total current liabilities	2,730	2,581
Deferred revenue, less current portion	339	413
Total liabilities	3,069	2,994
Commitments and Contingencies (Note 6)		
Stockholders' Equity		
Convertible preferred stock, $0.0001 par value; 714,286 shares authorized, none issued and outstanding at June 30, 2015 and March 31, 2015, respectively	–	–
Common stock, $0.0001 par value; 30,000,000 shares authorized, 15,681,565 and 15,045,080 shares issued and outstanding at June 30, 2015 and March 31, 2015, respectively	2	2
Additional paid-in capital	159,266	157,772
Accumulated deficit	(144,553)	(142,213)
Accumulated other comprehensive loss	(3,578)	(3,507)
Total stockholders' equity	11,137	12,054
Total liabilities and stockholders' equity	$ 14,206	$ 15,048

The accompanying footnotes are an integral part of these condensed consolidated financial statements.

OCULUS INNOVATIVE SCIENCES, INC. AND SUBSIDIARIES
Condensed Consolidated Statements of Comprehensive Loss
(In thousands, except per share amounts)
(Unaudited)

	Three Months Ended June 30,	
	2015	**2014**
Revenues		
Product	$ 2,916	$ 2,121
Product licensing fees and royalties	447	1,049
Service	317	222
Total revenues	3,680	3,392
Cost of revenues		
Product	1,516	1,322
Service	291	164
Total cost of revenues	1,807	1,486
Gross profit	1,873	1,906
Operating expenses		
Research and development	467	439
Selling, general and administrative	3,717	2,981
Total operating expenses	4,184	3,420
Loss from operations	(2,311)	(1,514)
Interest expense	–	(3)
(Loss) gain due to change in fair value of derivative liabilities	(59)	1,478
Other income expense, net	30	(31)
Net loss	(2,340)	(70)
Net loss per common share: basic and diluted	$ (0.15)	$ (0.01)
Weighted-average number of shares used in common share calculations:		
Basic and diluted	15,170	8,346
Other comprehensive loss		
Net loss	$ (2,340)	$ (70)
Foreign currency translation adjustments	(71)	8
Comprehensive loss	$ (2,411)	$ (62)

The accompanying footnotes are an integral part of these condensed consolidated financial statements.

OCULUS INNOVATIVE SCIENCES, INC. AND SUBSIDIARIES
Condensed Consolidated Statements of Cash Flows
(In thousands)
(Unaudited)

	Three Months Ended June 30,	
	2015	2014
	(In thousands)	
Cash flows from operating activities		
Net loss	$ (2,340)	$ (70)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation and amortization	62	64
Stock-based compensation	412	451
Service provider expenses settled with common stock	107	–
Loss (gain) due to change in fair value of derivative liabilities	59	(1,478)
Foreign currency transaction (gain) loss	(9)	8
Gain on disposal of property and equipment	–	(13)
Changes in operating assets and liabilities:		
Accounts receivable, net	(1,052)	(160)
Due from affiliate	–	537
Inventories, net	(82)	(213)
Prepaid expenses and other current assets	84	226
Accounts payable	284	206
Accrued expenses and other current liabilities	265	(41)
Deferred revenue	(375)	(863)
Net cash used in operating activities	(2,585)	(1,346)
Cash flows from investing activities:		
Purchases of property and equipment	(148)	(21)
Proceeds from sale of long-term investment	4,538	–
Long-term deposits	(12)	13
Net cash provided by (used in) investing activities	4,378	(8)
Cash flows from financing activities:		
Proceeds from issuance of common stock, net of offering costs	879	952
Principal payments on long-term debt	(52)	(83)
Net cash provided by financing activities	827	869
Effect of exchange rate on cash and cash equivalents	(4)	(8)
Net increase (decrease) in cash and cash equivalents	2,616	(493)
Cash and cash equivalents, beginning of period	6,136	5,480
Cash and cash equivalents, end of period	$ 8,752	$ 4,987
Supplemental disclosure of cash flow information:		
Cash paid for interest	$ –	$ 3
Non-cash operating and financing activities:		
Issuance of common stock to settle obligation	$ 96	$ –

The accompanying footnotes are an integral part of these condensed consolidated financial statements.

OCULUS INNOVATIVE SCIENCES, INC. AND SUBSIDIARIES
Notes to Condensed Consolidated Financial Statements
(Unaudited)

Note 1. Organization and Summary of Significant Accounting Policies

Organization

Oculus Innovative Sciences, Inc. (the "Company") was incorporated under the laws of the State of California in April 1999 and was reincorporated under the laws of the State of Delaware in December 2006. The Company's principal office is located in Petaluma, California. The Company is a global specialty pharmaceutical company that develops, produces, and markets solutions for the treatment of dermatological conditions and advanced tissue care in the United States and 39 countries around the world. The Company is pioneering innovative products for the dermatology, surgical, advanced tissue and skin care, and animal healthcare markets. The Company's key proprietary technology platform is called Microcyn® Technology. This technology is based on electrically charged oxychlorine small molecules designed to target a wide range of organisms that cause disease (pathogens). Several Microcyn® Technology tissue care products are designed to treat infections and enhance healing while reducing the need for antibiotics.

Basis of Presentation

The accompanying unaudited condensed consolidated financial statements as of June 30, 2015 and for the three months then ended have been prepared in accordance with the accounting principles generally accepted in the United States of America for interim financial information and pursuant to the instructions to Form 10-Q and Article 8 of Regulation S-X of the Securities and Exchange Commission ("SEC") and on the same basis as the Company prepares its annual audited consolidated financial statements. The condensed consolidated balance sheet as of June 30, 2015 and the condensed consolidated statements of comprehensive loss and cash flows for the three months ended June 30, 2015 and 2014 are unaudited, but include all adjustments, consisting only of normal recurring adjustments, which the Company considers necessary for a fair presentation of the consolidated financial position, operating results and cash flows for the periods presented. The results for the three months ended June 30, 2015 are not necessarily indicative of results to be expected for the year ending March 31, 2016 or for any future interim period. The condensed consolidated balance sheet at March 31, 2015 has been derived from audited consolidated financial statements. However, it does not include all of the information and notes required by accounting principles generally accepted in the United States of America for complete consolidated financial statements. The accompanying condensed consolidated financial statements should be read in conjunction with the consolidated financial statements for the year ended March 31, 2015, and notes thereto included in the Company's annual report on Form 10-K, which was filed with the SEC on June 16, 2015.

Note 2. Liquidity and Financial Condition

The Company reported a net loss of $2,340,000 and a loss from operations of $2,311,000 for the three months ended June 30, 2015. At June 30, 2015 and March 31, 2015, the Company's accumulated deficit amounted to $144,553,000 and $142,213,000, respectively. The Company had working capital of $10,534,000 and $7,066,000 as of June 30, 2015 and March 31, 2015, respectively. The Company expects to continue incurring losses for the foreseeable future and may need to raise additional capital to pursue its product development initiatives, penetrate markets for the sale of its products and continue as a going concern.

Pursuant to an At-the-Market Issuance Sales Agreement with MLV & Co. LLC dated April 2, 2014, the Company may issue and sell shares of common stock having an aggregate offering price of up to $9,159,000 from time to time through MLV acting as the Company's sales agent. During the three months ended June 30, 2015, the Company sold 500,000 shares of common stock for gross proceeds of $907,000 and net proceeds of $879,000 after deducting commissions and other offering expenses. Shares of common stock sold subsequent to June 30, 2015 are disclosed in Note 12.

During the three months ended June 30, 2015, the Company sold the 1,650,000 shares of Ruthigen common stock for proceeds of $4,537,500. Additionally, during the three months ended June 30, 2015, the Company paid a banker a $165,000 a finder fee related to the sale transaction (Note 3).

The Company currently anticipates that its cash and cash equivalents will be sufficient to meet its working capital requirements to continue its sales and marketing and research and development efforts through at least twelve months. However, in order to execute the Company's long-term Microcyn® product development strategy and to penetrate new and existing markets, the Company may need to raise additional funds through public or private equity offerings, debt financings, corporate collaborations or other means and potentially reduce operating expenditures.

Management believes that the Company has access to capital resources through possible public or private equity offerings, debt financings, corporate collaborations or other means. If the Company is unable to secure additional capital, it may be required to curtail its research and development initiatives and take additional measures to reduce costs in order to conserve its cash.

Note 3. Summary of Significant Accounting Policies

Use of Estimates

The preparation of condensed consolidated financial statements in conformity with accounting principles generally accepted in the United States of America requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosures of contingent liabilities at the dates of the condensed consolidated financial statements and the reported amounts of revenues and expenses during the reporting periods. Actual results could differ from these estimates. Significant estimates and assumptions include reserves and write-downs related to receivables and inventories, the recoverability of long-lived assets, the valuation allowance relating to the Company's deferred tax assets, valuation of equity and derivative instruments, debt discounts, valuation of investments, and the estimated amortization periods of upfront product licensing fees received from customers. Periodically, the Company evaluates and adjusts estimates accordingly. The allowance for doubtful accounts represents probable credit losses at June 30, 2015 and March 31, 2015 in the amounts of $19,000 and $20,000, respectively. Additionally at June 30, 2015 and March 31, 2015 the Company has allowances of $99,000 and $183,000, respectively, related to potential discounts, returns, distributor fees and rebates. The allowances are included in Accounts Receivable, net in the accompanying condensed consolidated balance sheets.

Net Loss per Share

The Company computes basic net loss per share by dividing net loss per share available to common stockholders by the weighted average number of common shares outstanding for the period and excludes the effects of any potentially dilutive securities. Diluted earnings per share, if presented, would include the dilution that would occur upon the exercise or conversion of all potentially dilutive securities into common stock using the "treasury stock" and/or "if converted" methods as applicable. The computation of basic loss per share for the three months ended June 30, 2015 and 2014 excludes the potentially dilutive securities summarized in the table below because their inclusion would be anti-dilutive.

	June 30,	
	2015	**2014**
Options to purchase common stock	2,979,000	2,536,000
Warrants to purchase common stock	7,740,000	2,300,000
	10,719,000	4,836,000

Common Stock Purchase Warrants and Other Derivative Financial Instruments

The Company classifies common stock purchase warrants and other free standing derivative financial instruments as equity if the contracts (i) require physical settlement or net-share settlement or (ii) give the Company a choice of net-cash settlement or settlement in its own shares (physical settlement or net-share settlement). The Company classifies any contracts that (i) require net-cash settlement (including a requirement to net cash settle the contract if an event occurs and if that event is outside the control of the Company), (ii) give the counterparty a choice of net-cash settlement or settlement in shares (physical settlement or net-share settlement), or (iii) contain reset provisions as either an asset or a liability. The Company assesses classification of its freestanding derivatives at each reporting date to determine whether a change in classification between assets and liabilities is required. The Company determined that its freestanding derivatives, which principally consist of warrants to purchase common stock, satisfied the criteria for classification as equity instruments, other than certain warrants that contained reset provisions and certain warrants that required net-cash settlement that the Company classified as derivative liabilities as more fully described in Note 5.

Reclassifications

Certain prior period amounts have been reclassified for comparative purposes to conform to the fiscal 2016 presentation. These reclassifications have no impact on the Company's previously reported net loss.

Revenue Recognition and Accounts Receivable

The Company generates revenue from sales of its products to hospitals, medical centers, doctors, pharmacies, and distributors. The Company sells products directly to third parties and to distributors through various cancelable distribution agreements. The Company also entered into agreements to license its technology and products.

The Company also provides regulatory compliance testing and quality assurance services to medical device and pharmaceutical companies.

The Company records revenue when (i) persuasive evidence of an arrangement exists, (ii) delivery has occurred, (iii) the fee is fixed or determinable, and (iv) collectability of the sale is reasonably assured.

The Company requires all product sales to be supported by evidence of a sale transaction that clearly indicates the selling price to the customer, shipping terms and payment terms. Evidence of an arrangement generally consists of a contract or purchase order approved by the customer. The Company has ongoing relationships with certain customers from which it customarily accepts orders by telephone in lieu of purchase orders.

The Company recognizes revenue at the time it receives confirmation that the goods were either tendered at their destination, when shipped "FOB destination," or transferred to a shipping agent, when shipped "FOB shipping point." Delivery to the customer is deemed to have occurred when the customer takes title to the product. Generally, title passes to the customer upon shipment, but could occur when the customer receives the product based on the terms of the agreement with the customer.

The selling prices of all goods are fixed, and agreed to with the customer, prior to shipment. Selling prices are generally based on established list prices. The Company does not customarily permit customers to return any products for monetary refunds or credit against completed or future sales. The Company may, from time to time, replace expired goods on a discretionary basis. The Company records these types of adjustments, when made, as a reduction of revenue.

The Company evaluates the creditworthiness of new customers and monitors the creditworthiness of its existing customers to determine whether events or changes in their financial circumstances would raise doubt as to the collectability of a sale at the time in which a sale is made. Payment terms on sales made in the United States are generally 30 days and internationally, generally range from 30 days to 90 days.

In the event a sale is made to a customer under circumstances in which collectability is not reasonably assured, the Company either requires the customer to remit payment prior to shipment or defers recognition of the revenue until payment is received. The Company maintains a reserve for amounts which may not be collectible due to risk of credit losses.

Additionally, the Company defers recognition of revenue related to distributors that are unable to provide inventory or product sell-through reports on a timely basis, until payment is received. The Company believes the receipt of payment is the best indication of product sell-through.

When the Company receives letters of credit and the terms of the sale provide for no right of return except to replace defective product, revenue is recognized when the letter of credit becomes effective and the product is shipped.

Product license revenue is generated through agreements with strategic partners for the commercialization of Microcyn® products. The terms of the agreements sometimes include non-refundable upfront fees. The Company analyzes multiple element arrangements to determine whether the elements can be separated. Analysis is performed at the inception of the arrangement and as each product is delivered. If a product or service is not separable, the combined deliverables are accounted for as a single unit of accounting and recognized over the performance obligation period.

When appropriate, the Company defers recognition of non-refundable upfront fees. If the Company has continuing performance obligations then such up-front fees are deferred and recognized over the period of continuing involvement.

The Company recognizes royalty revenues from licensed products upon the sale of the related products.

Revenue from consulting contracts is recognized as services are provided. Revenue from testing contracts is recognized as tests are completed and a final report is sent to the customer.

Inventories

Inventories are stated at the lower of cost, cost being determined on a standard cost basis (which approximates actual cost on a first-in, first-out basis), or market.

Due to changing market conditions, estimated future requirements, age of the inventories on hand and production of new products, the Company regularly reviews inventory quantities on hand and records a provision to write down excess and obsolete inventory to its estimated net realizable value. The Company recorded reserves to reduce the carrying amounts of inventories to their net realizable value in the amounts of $121,000 and $87,000 at June 30, 2015 and March 31, 2015, respectively, which is included in cost of product revenues on the Company's accompanying condensed consolidated statements of comprehensive loss.

Income Taxes

The Company is required to determine the aggregate amount of income tax expense or loss based upon tax statutes in jurisdictions in which it conducts business. In making these estimates, the Company adjusts its results determined in accordance with generally accepted accounting principles for items that are treated differently by the applicable taxing authorities. Deferred tax assets and liabilities resulting from these differences are reflected on its balance sheet for temporary differences in loss and credit carryforwards that will reverse in subsequent years. The Company also establishes a valuation allowance against deferred tax assets when it is more likely than not that some or all of the deferred tax assets will not be realized. Valuation allowances are based, in part, on predictions that management must make as to the results in future periods. The outcome of events could differ over time which would require that the Company makes changes in its valuation allowance.

Financial Assets and Liabilities

Financial instruments, including cash and cash equivalents, accounts receivable, accounts payable and accrued liabilities are carried at cost, which management believes approximates fair value due to the short-term nature of these instruments. The fair value of capital lease obligations and equipment loans approximates their carrying amounts as a market rate of interest is attached to their repayment. The Company measures the fair value of financial assets and liabilities based on the exchange price that would be received for an asset or paid to transfer a liability (an exit price) in the principal or most advantageous market for the asset or liability in an orderly transaction between market participants on the measurement date. The Company maximizes the use of observable inputs and minimizes the use of unobservable inputs when measuring fair value. The Company uses three levels of inputs that may be used to measure fair value:

Level 1 – quoted prices in active markets for identical assets or liabilities

Level 2 – quoted prices for similar assets and liabilities in active markets; quoted prices for identical or similar instruments in markets that are not active; and model-derived valuations in which all significant inputs and significant value drivers are observable in active markets.

Level 3 – inputs that are unobservable (for example cash flow modeling inputs based on assumptions)

	Fair Value Measurements at June 30, 2015 Using			
	Total June 30, 2015	**Quoted prices in active markets for identical assets (Level 1)**	**Significant other observable inputs (Level 2)**	**Significant other unobservable inputs (Level 3)**
Liabilities:				
Derivative liabilities – warrants	$ 70,000	–	–	$ 70,000

	Fair Value Measurements at March 31, 2015 Using			
	Total March 31, 2015	**Quoted prices in active markets for identical assets (Level 1)**	**Significant other observable inputs (Level 2)**	**Significant other unobservable inputs (Level 3)**
Liabilities:				
Derivative liabilities – warrants	$ 11,000	–	–	$ 11,000

Level 3 liabilities are valued using unobservable inputs to the valuation methodology that are significant to the measurement of the fair value of the liabilities. For fair value measurements categorized within Level 3 of the fair value hierarchy, the Company's accounting and finance department, who report to the Chief Financial Officer, determine its valuation policies and procedures. The development and determination of the unobservable inputs for Level 3 fair value measurements and fair value calculations are the responsibility of the Company's accounting and finance department and are approved by the Chief Financial Officer.

Level 3 Valuation Techniques:

Level 3 financial liabilities consist of the derivative liabilities for which there is no current market for these securities such that the determination of fair value requires significant judgment or estimation. Changes in fair value measurements categorized within Level 3 of the fair value hierarchy are analyzed each period based on changes in estimates or assumptions and recorded as appropriate.

The Company uses the Black-Scholes option valuation model to value Level 3 derivatives at inception and on subsequent valuation dates. This model incorporates transaction details such as the Company's stock price, contractual terms, maturity, risk free rates, as well as volatility. A significant decrease in the volatility or a significant decrease in the Company's stock price, in isolation, would result in a significantly lower fair value measurement. Changes in the values of the derivative liabilities are recorded in "Loss (gain) due to change in fair value of derivative liabilities" in the Company's condensed consolidated statements of comprehensive loss.

As of June 30, 2015 and March 31, 2015, there were no transfers in or out of Level 3 from other levels in the fair value hierarchy.

Long-Term Investments

The Company accounted for its ownership of shares of Ruthigen common stock at cost in accordance with Accounting Standards Codification ("ASC") 325-20 as a result of (a) the restrictions on voting the shares held as disclosed above, (b) the Company having no representation on the Ruthigen Board of Directors, (c) the Company's inability to set policy at Ruthigen (d) the Company having no further commitments for funding the operations of Ruthigen and (e) the restrictions on transferability of its shares.

The Company's long-term investments consisted of the Company's ownership of 1,650,000 shares of Ruthigen common stock at March 31, 2015. During the three months ended June 30, 2015, the Company sold its remaining 1,650,000 shares of Ruthigen common stock for proceeds of $4,537,500 pursuant to a securities purchase agreement with two investors. Additionally, during the three months ended June 30, 2015, the Company paid a $165,000 banker fee related to the sale transaction.

Subsequent Events

Management has evaluated subsequent events or transactions occurring through the date the condensed consolidated financial statements were issued (Note 12).

Recent Accounting Pronouncements

Accounting standards that have been issued or proposed by the Financial Accounting Standards Board ("FASB"), SEC and/or other standards-setting bodies that do not require adoption until a future date are not expected to have a material impact on the condensed consolidated financial statements upon adoption.

Note 4. Condensed Consolidated Balance Sheets

Inventories, net

Inventories, net consist of the following:

	June 30, 2015	March 31, 2015
Raw materials	$ 1,011,000	$ 865,000
Finished goods	455,000	537,000
	$ 1,466,000	$ 1,402,000

Note 5. Derivative Liabilities

The Company deems financial instruments, which require net-cash settlement, as either an asset or a liability. The common stock purchase warrants issued in conjunction with the Company's December 9, 2013 and February 26, 2014 registered direct offerings contain net-cash settlement features, which give the warrant holder the right to a net-cash settlement in the event certain transactions occur. Pursuant to the terms of the warrants, if such a transaction occurs, the warrant holder will be entitled to a net-cash settlement value calculated using the Black-Scholes valuation model using an expected volatility equal to the greater of 100% and the 30 day volatility obtained from the HVT function on Bloomberg, an expected term equal to the remaining term of the warrants, and applicable risk-free interest rate corresponding to the U.S. Treasury.

The derivative liabilities relating to the warrants with net-cash settlement provisions were valued using the Black-Scholes option valuation model and the following assumptions on the following dates:

	Measurement Date	Warrants	Remaining Contract Term in Years	Exercise Price	Volatility	Risk-free Interest Rate	Fair Value
Warrant							
Placement Agent Warrants	March 31, 2014	16,500	2.09	$ 5.00	128%	0.44%	$ 37,000
Investor - Series A Warrants	March 31, 2014	1,000	1.41	$ 3.00	128%	0.44%	1,000
Investor - Series B Warrants	March 31, 2014	1,400,000	1.41	$ 3.63	128%	0.44%	2,958,000
Placement Agent Warrants	March 31, 2014	69,037	2.09	$ 3.00	128%	0.44%	179,000
							$ 3,175,000
Warrant							
Placement Agent Warrants	June 30, 2014	16,500	1.84	$ 5.00	100%	0.47%	$ 19,000
Investor - Series A Warrants	June 30, 2014	1,000	1.16	$ 3.00	100%	0.11%	1,000
Investor - Series B Warrants	June 30, 2014	1,400,000	1.16	$ 3.63	100%	0.11%	1,568,000
Placement Agent Warrants	June 30, 2014	69,037	1.84	$ 3.00	100%	0.47%	109,000
							$ 1,697,000
Warrant							
Placement Agent Warrants	March 31, 2015	16,500	1.09	$ 5.00	100%	0.26%	$ 1,000
Investor - Series A Warrants	March 31, 2015	1,000	0.41	$ 3.00	100%	0.14%	–
Investor - Series B Warrants	March 31, 2015	1,400,000	0.41	$ 3.63	100%	0.14%	5,000
Placement Agent Warrants	March 31, 2015	69,037	1.09	$ 3.00	100%	0.26%	5,000
							$ 11,000
Warrant							
Placement Agent Warrants	June 30, 2015	16,500	0.84	$ 5.00	143%	0.28%	$ 5,000
Investor - Series A Warrants	June 30, 2015	1,000	0.16	$ 3.00	143%	0.01%	–
Investor - Series B Warrants	June 30, 2015	1,400,000	0.16	$ 3.63	143%	0.01%	35,000
Placement Agent Warrants	June 30, 2015	69,037	0.84	$ 3.00	143%	0.28%	30,000
							$ 70,000

The following table sets forth a summary of the changes in the fair value of our Level 3 financial liabilities that are measured at fair value on a recurring basis:

	Three Months Ended June 30,	
	2015	2014
Beginning balance	$ 11,000	$ 3,175,000
Mark to market net unrealized loss (gain)	59,000	$ (1,478,000)
Ending balance	$ 70,000	$ 1,697,000

Note 6. Commitments and Contingencies

Legal Matters

The Company, on occasion, may be involved in legal matters arising in the ordinary course of our business including matters involving proprietary technology. While management believes that such matters are currently insignificant, matters arising in the ordinary course of business for which the Company is or could become involved in litigation may have a material adverse effect on its business, financial condition or results of operations.

On November 13, 2014, the Company received a letter from Exeltis USA Dermatology, Inc. formerly known as Quinnova Pharmaceuticals, Inc., and herein referred to as "Exeltis", claiming that the Company breached its Exclusive Sales and Distribution Agreement with Exeltis. Specifically, Exeltis claimed that the marketing and selling of its Alevicyn gel product violates the terms of the Exclusive Sales and Distribution Agreement and demanded the Company cease and desist from any further marketing or sales. The Company believes that the marketing and selling of its Alevicyn gel is not in violation of the Exclusive Sales and Distribution Agreement and that the claims made by Exeltis are without merit. Exeltis continues to purchase products from the Company under a new, non-exclusive distribution agreement for sale to their customers under their own brand. The Company intends to defend this matter vigorously and does not believe an accrual for a potential loss relating to this matter is necessary at this time. While the Company believes this claim is without merit, there can be no assurances provided that the outcome of this matter will be favorable to the Company or will not have a negative impact on its condensed consolidated financial position or results of operations.

Employment Agreements

As of June 30, 2015, the Company had employment agreements in place with four of its key executives. The agreements provide, among other things, for the payment of nine to twenty-four months of severance compensation for terminations under certain circumstances. With respect to these agreements, at June 30, 2015, potential severance amounted to $1,130,000 and aggregated annual salaries amounted to $935,000.

Note 7. Stockholders' Equity

Authorized Capital

The Company is authorized to issue up to 30,000,000 shares of common stock with a par value of $0.0001 per share and 714,286 shares of convertible preferred stock with a par value of $0.0001 per share.

On June 29, 2015, the stockholders of the Company approved a reverse stock split of the Company's outstanding common stock and to proportionally decrease the total number of shares that the Company is authorized to issue at a whole number ratio in the range of 1-for-5 to 1-for-9, such ratio to be determined in the discretion of the Company's Board of Directors, and authorized the Company's Board of Directors to effect the reverse stock split, if their judgment it is necessary, at any time until June 29, 2016, upon which date the resolution lapses. To date, no reverse stock split has been authorized by the Board of Directors.

Sale of Common Stock

Pursuant to an At-the-Market Issuance Sales Agreement with MLV & Co. LLC dated April 2, 2014, the Company may issue and sell shares of common stock having an aggregate offering price of up to $9,159,000 from time to time through MLV acting as the Company's sales agent. During the three months ended June 30, 2015, the Company sold 500,000 shares of common stock for gross proceeds of $907,000 and net proceeds of $879,000 after deducting commissions and other offering expenses. As of June 30, 2015, the Company has sold an aggregate 967,934 shares of common stock to date, for gross proceeds of $2,350,000 and net proceeds of $2,220,000 after deducting commissions and other offering expenses. The Company pays MLV a commission rate equal to 3.0% of the gross proceeds from the sale of any shares of common stock sold through MLV as agent. Shares of common stock sold subsequent to June 30, 2015 are disclosed in Note 12.

Common Stock Issued to Settle Fees for Services Provided

On April 24, 2009, the Company entered into an agreement with Advocos LLC, a contract sales organization that serves as part of the Company's sales force, for the sale of the Company's wound care products in the United States. Pursuant to the agreement, the Company agreed to pay the contract sales organization a monthly fee and potential bonuses that will be based on achievement of certain levels of sales. The Company agreed to issue the contract sales organization cash or shares of common stock to settle fees for its services. During the three months ended June 30, 2015, the Company issued 135,485 shares of common stock, with a fair market value of $203,000, in connection with this agreement. The Company has determined that the fair value of the common stock was more readily determinable than the fair value of the services rendered. During the three months ended June 30, 2015, the Company recorded $107,000 of expense related

to this agreement and settled $96,000 of fees accrued in prior periods. The expense was recorded as selling, general and administrative expense in the accompanying condensed consolidated statement of comprehensive loss for the three months ended June 30, 2015.

Note 8. Stock-Based Compensation

On June 1, 2015, the Board of Directors of the Company approved an increase of 250,000 shares authorized for issuance under the 2006 Stock Plan as of April 1, 2015, and an increase of 2,256,762 shares authorized for issuance under the 2011 Stock Plan as of April 1, 2015.

The Company estimated the fair value of employee and non-employee stock options using the Black-Scholes option pricing model. The fair values of employee and non-employee stock options are being amortized on a straight-line basis over the requisite service periods of the respective awards. Compensation expense includes the impact of an estimate for forfeitures for all stock options.

The expected term of stock options represents the average period the stock options are expected to remain outstanding and is based on the expected term calculated using the approach prescribed by the SEC's Staff Accounting Bulletin No. 110 for "plain vanilla" options. The expected stock price volatility for the Company's stock options was determined by using an average of the historical volatilities of the Company and its industry peers for non-employee grants and was determined by using the historical volatilities of the Company for employee options. The Company will continue to analyze the stock price volatility and expected term assumptions as more data for the Company's common stock and exercise patterns become available. The risk-free interest rate assumption is based on the U.S. Treasury instruments whose term was consistent with the expected term of the Company's stock options. The expected dividend assumption is based on the Company's history and expectation of dividend payouts. The Company estimates forfeitures based on historical experience and reduces compensation expense accordingly. The estimated forfeiture rates used during the three months ended June 30, 2015 ranged from 0.85% to 1.81%. The estimated forfeiture rates used during the three months ended June 30, 2014 ranged from 0.36% to 0.37%.

The fair value of the stock options granted was calculated using the Black-Scholes option-pricing model using the following weighted-average assumptions:

	Three Months Ended June 30, 2015
Expected life	8.96 years
Risk-free interest rate	2.18%
Dividend yield	0.00%
Volatility	86%
Fair value of options granted	$ 1.08

The Company did not grant stock options during the three months ended June 30, 2014.

Stock-based compensation expense is as follows:

	Three Months Ended June 30,	
	2015	2014
Cost of revenues	$ 55,000	$ 64,000
Research and development	73,000	96,000
Selling, general and administrative	284,000	291,000
Total stock-based compensation	$ 412,000	$ 451,000

The aggregate intrinsic value is calculated as the difference between the exercise price of the underlying stock options and the fair value of the Company's common stock ($1.43) for stock options.

At June 30, 2015, there were unrecognized compensation costs of $2,473,000 related to stock options which is expected to be recognized over a weighted-average amortization period of 1.74 years.

The Company did not capitalize any cost associated with stock-based compensation.

The Company issues new shares of common stock upon exercise of stock options.

No income tax benefit has been recognized relating to stock-based compensation expense and no tax benefits have been realized from exercised stock options.

A summary of all option activity as of June 30, 2015 and changes during the three months then ended is presented below:

	Number of Shares	Weighted-Average Exercise Price	Weighted-Average Contractual Term	Aggregate Intrinsic Value
Outstanding at April 1, 2015	2,877,000	$ 6.96		
Options granted	155,000	1.32		
Options exercised	–	–		
Options forfeited or expired	(53,000)	(4.97)		
Outstanding at June 30, 2015	2,979,000	$ 6.70	7.19	$ 38,000
Exercisable at June 30, 2015	1,781,000	$ 8.91	6.05	$ 36,000
Options available for grant as of June 30, 2015	3,858,000			

Note 9. Income Taxes

The Company has completed a study to assess whether a change in control has occurred or whether there have been multiple changes of control since the Company's formation. The Company determined, based on the results of the study, no change in control occurred for purposes of Internal Revenue Code section 382. The Company, after considering all available evidence, fully reserved its deferred tax assets since it is more likely than not, such benefits, will not be realized in future periods. The Company incurred losses for both financial reporting and income tax purposes for the year ended March 31, 2015. Accordingly, the Company is continuing to fully reserve for its deferred tax assets. The Company will continue to evaluate its deferred tax assets to determine whether any changes in circumstances could affect the realization of their future benefit. If it is determined in future periods that portions of the Company's deferred income tax assets satisfy the realization standards, the valuation allowance will be reduced accordingly.

As a result of certain realization requirements of Accounting Standards Codification Topic 718, the Company's deferred tax assets and liabilities do not include certain deferred tax assets at June 30, 2015 that arose directly from tax deductions related to equity compensation in excess of compensation recognized for financial reporting purposes. Equity will be increased by approximately $533,000 if and when such deferred tax assets are ultimately realized.

Note 10. Segment and Geographic Information

The Company generates product revenues from wound care products that are sold into the human and animal healthcare markets, and the Company generates service revenues from laboratory testing services, which are provided to medical device manufacturers.

The Company operates a single segment business for product revenues, which consists of three geographical sales territories as follows:

	Three Months Ended June 30,	
Product related revenues	**2015**	**2014**
United States	$ 787,000	$ 355,000
Latin America	1,558,000	1,093,000
Europe and Rest of the World	571,000	673,000
	2,916,000	2,121,000
Product license fees and royalties	447,000	1,049,000
Total product related revenues	$ 3,363,000	$ 3,170,000

The following table shows the Company's product license fees and royalties revenues by partner:

	Three Months Ended June 30,	
Product license fees and royalties	**2015**	**2014**
Exeltis (formerly Quinnova)	$ 54,000	$ 146,000
Innovacyn	20,000	528,000
Laboratorios Sanfer (formerly More Pharma)	373,000	375,000
Total product license fees and royalties	$ 447,000	$ 1,049,000

The Company's service revenues amounted to $317,000 and $222,000 for the three months ended June 30, 2015 and 2014, respectively.

Note 11. Significant Customer Concentrations

For the three months ended June 30, 2015, one customer represented 53% of the quarter's revenue. For the three months ended June 30, 2014, one customer represented 43%, and one customer represented 16% of the quarter's revenue.

At June 30, 2015, one customer represented 57%, and one customer represented 11% of the net accounts receivable balance. At March 31, 2015, one customer represented 56%, and one customer represented 14% of the net accounts receivable balance.

Note 12. Subsequent Events

Pursuant to an At-the-Market Issuance Sales Agreement with MLV & Co. LLC dated April 2, 2014, the Company may issue and sell shares of common stock having an aggregate offering price of up to $9,159,000 from time to time through MLV acting as the Company's sales agent. Subsequent to June 30, 2015, the Company sold 724,100 shares of common stock for gross proceeds of $1,157,000 and net proceeds of $1,121,000 after deducting commissions and other offering expenses.

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

The following discussion of our financial condition and results of operations should be read in conjunction with the condensed consolidated financial statements and notes to those statements included elsewhere in this Quarterly Report on Form 10-Q as of June 30, 2015 and our audited consolidated financial statements for the year ended March 31, 2015 included in our Annual Report on Form 10-K, filed with the Securities and Exchange Commission on June 16, 2015.

This report contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. When used in this report, the words "anticipates," "suggests," "estimates," "plans," "projects," "continue," "ongoing," "potential," "expect," "predict," "believe," "intend," "may," "will," "should," "could," "would," "proposal," and similar expressions are intended to identify forward-looking statements.

Forward-looking statements are subject to risks and uncertainties that could cause our actual results to differ materially from those projected. These risks and uncertainties include, but are not limited to the risks described in our Annual Report on Form 10-K including: our ability to become profitable; the progress and timing of our development programs and regulatory approvals for our products; the benefits and effectiveness of our products; the ability of our products to meet existing or future regulatory standards; the progress and timing of clinical trials and physician studies; our expectations related to the use of our cash reserves; our expectations and capabilities relating to the sales and marketing of our current products and our product candidates; our ability to gain sufficient reimbursement from third-party payors; our ability to compete with other companies that are developing or selling products that are competitive with our products; the establishment of strategic partnerships for the development or sale of products; the risk our research and development efforts do not lead to new products; the timing of commercializing our products; our ability to penetrate markets through our sales force, distribution network, and strategic business partners to gain a foothold in the market and generate attractive margins; the expansion of our sales force and distribution network; the ability to attain specified revenue goals within a specified time frame, if at all, or to reduce costs; the outcome of discussions with the U.S. Food and Drug Administration, or FDA, and other regulatory agencies; the content and timing of submissions to, and decisions made by, the FDA and other regulatory agencies, including demonstrating to the satisfaction of the FDA the safety and efficacy of our products; our ability to manufacture sufficient amounts of our product candidates for clinical trials and products for commercialization activities; our ability to protect our intellectual property and operate our business without infringing on the intellectual property of others; our ability to continue to expand our intellectual property portfolio; our expectations about the outcome of litigation and controversies with third parties; the risk we may need to indemnify our distributors or other third parties; our ability to attract and retain qualified directors, officers and employees; our expectations relating to the concentration of our revenue from international sales; our ability to expand to and commercialize products in markets outside the wound care market; and the impact of the Sarbanes-Oxley Act of 2002 and any future changes in accounting regulations or practices in general with respect to public companies. These forward-looking statements speak only as of the date hereof. We expressly disclaim any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements contained herein to reflect any change in our expectations with regard thereto or any change in events, conditions or circumstances on which any such statement is based, except as required by law.

Additional Information

Investors and others should note that we announce material financial information using our company website (www.oculusis.com), our investor relations website (ir.oculusis.com), SEC filings, press releases, public conference calls and webcasts. Information about Oculus, our business, and our results of operations may also be announced by posts on the following social media channels:

- Oculus corporate blog (http://oculusis.com/dialogue/)
- Oculus Facebook page (www.facebook.com/oculusinnovativesciences)
- Dan McFadden's Twitter feed (http://twitter.com/dmcfaddenocls). Mr. McFadden is the Vice President of Public and Investor Relations of our Company.

The information that we post on these social media channels could be deemed to be material information. Therefore, we encourage investors, the media, and others interested in Oculus to review the information that we post on these social media channels. These social media channels may be updated from time to time on Oculus' investor relations website. The information on, or accessible through, our websites and social media channels is not incorporated by reference in this Quarterly Report on Form 10-Q.

Our Business

We are a specialty pharmaceutical company that develops and markets solutions for the treatment of dermatological conditions and advanced tissue care. Our products, which are sold throughout the United States and 39 countries around the world, have improved patient outcomes for more than five million patients globally by reducing infections, itch, pain, scarring, odor and harmful inflammatory responses.

We initially built our business by developing and promoting products via partnerships. Our key proprietary technology, Microcyn®, is based on electrically charged oxychlorine small molecules designed to target a wide range of pathogens that cause disease. These pathogens include viruses, fungi, spores and bacteria, including antibiotic-resistant strains such as methicillin-resistant *Staphylococcus aureus*, or MRSA, and vancomycin-resistant *Enterococcus,* or VRE, as well as *Clostridium difficile,* or C. diff, a highly contagious bacteria spread by human contact. Several Microcyn® Technology advanced tissue care products are designed to treat infections and enhance healing while reducing the need for antibiotics.

To date, we have obtained eleven clearances from the U.S. Food and Drug Administration, or FDA, that permit us to sell our Microcyn®-based products as medical devices for Section 510(k) of the Federal Food, Drug and Cosmetic Act in the United States. However, we do not have the necessary regulatory approvals to market Microcyn® as a drug.

Our clinical trials from around the world suggest that our Microcyn® Technology helps reduce a wide range of pathogens while curing or improving infection. Our clinical studies suggest that our Microcyn® Technology is safe, easy to use and complementary to many existing treatment methods in dermatology and advanced tissue care. These clinical studies and usage of our products in the United States also suggest that our 510(k)-cleared products may shorten hospital stays, lower aggregate patient care costs and, in certain cases, reduce the need for systemic antibiotics.

Outside of the United Sates, we sell products for dermatological and advanced tissue care with a European Conformity marking (known as Conformité Européenne or CE) covering ten of our products, 14 approvals from the Mexican Ministry of Health, and various approvals in Central America, China, Southeast Asia, and the Middle East.

In 2013 and 2014, we added new members to our Board of Directors, thus enhancing our expertise in sales, marketing, strategy and dermatology, and we hired new managers to complement our executive team. Our new team commenced a strategic realignment of our business with a sharp focus on dermatology markets. Our decision to focus on dermatology was based on our already strong presence in this market and the ability of our core hypochlorous acid-based technology, Microcyn®, to address other dermatological indications including acne, atopic dermatitis, anti-itch and scar management.

Building upon our commercialization experience selling our Microcyn® Technology-based products, we believe we can significantly increase our revenue growth by focusing on our own dermatology efforts. Key aspects of our dermatology growth strategy are set forth below:

Expand our Internal U.S. Sales Force: We recently hired and intend to hire an additional experienced dermatology management team and sales force, most of who are seasoned sales veterans that have established relationships with dermatologists in their territories.

Develop and Launch New Dermatology Products: In October 2014, we launched two prescription dermatology products in the United States, an antipruritic gel and dermal spray. We also have a strong product pipeline, including our new product for treatment of scars that we intend to launch over the next 12 months. We have licensed several proprietary dermatology products from two European dermatology companies that we believe we can bring to market in the near term.

Create a Competitive Pricing Strategy: We have and will continue to develop a unique product pricing strategy, which we believe solves many of the challenges associated with the prescription dermatology market's current pricing and rebate programs.

Develop a Pharmaceutical Line: We plan to acquire or develop pharmaceutical products with affordable clinical trials to increase our market presence and create innovator patent protection.

Generate International Growth: In Europe, we received clearance for four new dermatology products during the year for acne, atopic dermatitis, scar reduction, three of which we launched in the fall of 2014 and spring of 2015 and we are in the process of contracting experienced, country-specific dermatology distributors to sell three products with these indications. We intend to launch a new product for post-laser procedures in the fall of 2015.

Our plan is to evolve into a leading dermatology and advanced tissue care company, providing innovative and cost-effective solutions to patients, while generating strong,consistent revenue growth and maximizing long-term shareholder value.

Comparison of the Quarters Ended June 30, 2015 and 2014

Revenues

Total revenues of $3,680,000 increased by $288,000 or 8% for the quarter ended June 30, 2015, as compared to $3,392,000 for the quarter ended June 30, 2014. The increase in product revenue was largely, led by growth in the United States and Latin America of $795,000 or 37%, which was offset by a decrease in product licensing fees and royalties of $602,000 mostly related to animal health care.

Product revenues in the United States for the quarter ended June 30, 2015 of $787,000, increased by $432,000, or 122%, when compared to the same period in the prior year as a result of the launch of our dermatology and animal health care products as well as increases in our advanced wound tissue products. In October 2014, we hired a direct sales force focused on dermatology and launched four new dermatology products, one of which was just recently approved by the FDA. Additionally, at the end of the quarter ended March 31, 2015, we launched our animal health care products with our new animal health care partner, SLA Brands.

Product revenue in Latin America for the quarter ended June 30, 2015 of $1,558,000 increased by $465,000, or 43%, when compared to the same period in the prior year. This increase was dampened by an 18% decline in the peso from the same period in prior year. The sales growth in local currency was 68% with strong increases in all product categories. The increase in revenue is primarily attributed to the larger distribution network of our new partner Laboratorios Sanfer, S.A. de C.V.

Product revenue in Europe and the Rest of the World for the quarter ended June 30, 2015 of $571,000, decreased by $102,000, or 15%, as compared to the same period in the prior year, with decreases in Europe, Middle East, Singapore and India. The decrease in Europe is the result of the 23% decline in the Euro with a local currency growth of 16% for the quarter. The sales to the Middle East are sporadic and the sales from our new partner in India will not start until the quarter ending September 2015.

The following table shows our product revenues by geographic region:

	Three Months Ended June 30,			
	2015	2014	$ Change	% Change
United States	$ 787,000	$ 355,000	$ 432,000	122%
Latin America	1,558,000	1,093,000	465,000	43%
Europe and Rest of the World	571,000	673,000	(102,000)	(15%)
	2,916,000	2,121,000	795,000	37%
Product license fees and royalties	447,000	1,049,000	(602,000)	(57%)
Total	$ 3,363,000	$ 3,170,000	$ 193,000	6%

In the quarter ended June 30, 2015, product license fees and royalties revenue declined primarily as a result of the termination of our agreement with Innovacyn and a decline in unit volume sold by Exeltis (formerly Quinnova).

The following table shows our product license fees and royalties revenue by partner:

	Three Months Ended June 30,			
Product license fees and royalties	**2015**	**2014**	**$ Change**	**% Change**
Exeltis (formerly Quinnova)	$ 54,000	$ 146,000	$ (92,000)	(63)%
Innovacyn	20,000	528,000	(508,000)	(96)%
Laboratorios Sanfer (formerly More Pharma)	373,000	375,000	(2,000)	(1)%
Total product license fees and royalties	$ 447,000	$ 1,049,000	$ (602,000)	(57)%

Service revenues were $317,000 and $222,000 for the quarters ended June 30, 2015 and 2014, respectively, due to an increase in the number of tests and services provided by our lab services business.

Gross Profit

We reported gross profit related to our products of $1,847,000 or 55% of product related revenues, during the quarter ended June 30, 2015, compared to a gross profit of $1,848,000, or 58% of product related revenues, for the same period in the prior year. Licensing fees and royalties revenues are included in our calculation of product related revenues and gross profit for the quarter ended June 30, 2015 and 2014.

Research and Development Expense

We reported research and development expense of $467,000 for the quarter ended June 30, 2015, an increase of $28,000, or 6%, when compared to the same period in the prior year. The increase is largely due to higher costs related to studies.

We expect our research and development expenses will remain relatively flat over the next year.

Selling, General and Administrative Expense

We reported selling, general and administrative of $3,717,000 for the quarter ended June 30, 2015, an increase of $736,000, or 25%, when compared to the same period in the prior year. The increase for the quarter ended June 30, 2015 was primarily due to higher sales and marketing expenses of $700,000 incurred primarily in the United States and Europe due to the recent hiring of a direct dermatology sales force, and higher costs related to the launch of four new dermatology products.

We expect selling, general and administrative expenses to remain within this range over the next several quarters.

Interest Expense and Interest Income

Interest expense and interest income were negligible for the quarters ended June 30, 2015 and 2014.

(Loss) gain due to Change in Fair Value of Derivative Liabilities

In connection with our December 9, 2013 and February 26, 2014 registered direct offerings we issued a series of common stock purchase warrants, which contain cash settlement provisions. During the quarter ended June 30, 2015, we recorded a loss due to an increase in the fair value of our derivative liabilities of $59,000, primarily due to an increase in our common stock price, partially offset by the decreasing contractual term of the warrants. During the quarter ended June 30, 2014, we recorded a gain due to a decrease in the fair value of our derivative liabilities of $1,478,000, primarily due to a decrease in our common stock price, partially offset by the decreasing contractual term of the warrants.

Other Income Expense, Net

Other income, net of $30,000 for the quarter ended June 30, 2015, increased $61,000, from other expense, net of $31,000 for the same period in the prior year. The increase in other income, net for the quarter ended June 30, 2015 was primarily related to foreign exchange gains and losses and a decrease in franchise tax expenses.

Net Loss

Net loss for the quarter ended June 30, 2015 was $2,340,000, an increase of $2,270,000, as compared to net loss of $70,000 for the same period in the prior year. The increase in net loss is primarily due to an increase of $736,000 in our selling general and administrative expenses, and the change in the fair value of our derivative liabilities of $1,537,000 when compared to the prior period.

Liquidity and Capital Resources

We incurred a net loss of $2,340,000 for the three months ended June 30, 2015. At June 30, 2015 and March 31, 2015, our accumulated deficit amounted to $144,553,000 and $142,213,000, respectively. At June 30, 2015 and March 31, 2015, our working capital amounted to $10,534,000 and $7,066,000, respectively. In the future, we may raise additional capital from external sources in order to continue the longer term efforts contemplated under our business plan. We expect to continue incurring losses for the foreseeable future and may need to raise additional capital to pursue our product development initiatives, to penetrate markets for the sale of our products and continue as a going concern. We cannot provide any assurances that we will be able to raise additional capital. Our management believes that we have access to capital resources through possible public or private equity offerings, debt financings, corporate collaborations or other means.

Sources of Liquidity

As of June 30, 2015, we had cash and cash equivalents of $8,752,000. Since our inception, substantially all of our operations have been financed through sales of equity securities. Other sources of financing that we have used to date include our revenues, as well as various loans.

Since July 1, 2013, substantially all of our operations have been financed through the following transactions:

- proceeds of $1,295,000 received from the exercise of common stock purchase warrants;
- net proceeds of $2,002,000 received from a registered direct offering on December 9, 2013;
- net proceeds of $1,187,000 received from a registered direct offering on February 26, 2014;
- net proceeds of $5,444,000 received from an underwritten public offering on January 26, 2015;
- net proceeds of $5,300,000 received from the sale of 2,000,000 Ruthigen shares; and
- net proceeds of $3,341,000 received from At the Market Issuances of common stock through August 4, 2015.

Cash Flows

As of June 30, 2015, we had cash and cash equivalents of $8,752,000, compared to $6,136,000 as of March 31, 2015.

Net cash used in operating activities during the three months ended June 30, 2015 was $2,585,000, primarily due to our net loss of $2,340,000 for the period and an increase in accounts receivable of $1,052,000 due to increased sales on payment terms, offset by an increase of $549,000 in accounts payable and accrued liabilities

Net cash used in operating activities during the three months ended June 30, 2014 was $1,346,000, primarily due to our net loss of $70,000 for the period which was offset by a $1,478,000 non-cash gain due to a change in fair value of derivative liabilities. Additionally, we had $451,000 of stock-based compensation expenses and we received $537,000 from our affiliate and formerly wholly-owned subsidiary Ruthigen, Inc.

Net cash provided by investing activities was $4,378,000 for the three months ended June 30, 2015, consisting of $4,538,000 received from the sale of 1,650,000 of our shares of Ruthigen common stock, offset by $148,000 related to equipment purchases and $12,000 related to changes in long-term assets.

Net cash used in investing activities was $8,000 for three months ended June 30, 2014, primarily related to the purchase of equipment in Mexico.

Net cash provided by financing activities was $827,000 for the three months ended June 30, 2015 was primarily related to $879,000 of net proceeds received from At-the-Market Issuances of common stock which was offset by principal payments on debt in the amount of $52,000.

Net cash provided by financing activities was $869,000 for the three months ended June 30, 2014 was primarily related to $952,000 of net proceeds received from At-the-Market Issuances of common stock which was offset by principal payments on debt in the amount of $83,000.

Operating Capital and Capital Expenditure Requirements

We incurred a net loss of $2,340,000 for the three months ended June 30, 2015. At June 30, 2015 and March 31, 2015, our accumulated deficit amounted to $144,553,000 and $142,213,000, respectively. At June 30, 2015 and March 31, 2015, our working capital amounted to $10,534,000 and $7,066,000, respectively.

We may need to raise additional capital from external sources in order to continue the longer term efforts contemplated under our business plan. We expect to continue incurring losses for the foreseeable future and may need to raise additional capital to pursue our product development initiatives and to penetrate markets for the sale of our products.

In order for us to potentially commercialize Microcyn® as a drug product in the United States, we must conduct clinical trials, which can be costly. Therefore, commencement of such pivotal clinical trials will be delayed until we find a strategic partner to assist with funding. Without a strategic partner or additional capital, our pivotal clinical trials will be delayed for a period of time that is currently indeterminate.

Our future funding requirements will depend on many factors, including:

- the cost and timing of establishing sales, marketing and distribution capabilities;
- the scope, rate of progress and cost of our clinical trials and other research and development activities;
- future clinical trial results;
- the terms and timing of any collaborative, licensing and other arrangements that we may establish;
- the cost and timing of regulatory approvals;
- the cost and delays in product development as a result of any changes in regulatory oversight applicable to our products;
- the effect of competing technological and market developments;
- the cost of filing, prosecuting, defending and enforcing any patent claims and other intellectual property rights; and
- the extent to which we acquire or invest in businesses, products and technologies.

Use of Estimates

The preparation of condensed consolidated financial statements in conformity with accounting principles generally accepted in the United States requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosures of contingent liabilities at the dates of the condensed consolidated financial statements and the reported amounts of revenues and expenses during the reporting periods. Actual results could differ from these estimates. Significant estimates and assumptions include reserves and write-downs related to receivables and inventories, the recoverability of long-lived assets, the valuation allowance related to our deferred tax assets, valuation of equity and derivative instruments, debt discounts, valuation of investments and the estimated amortization periods of upfront product licensing fees received from customers.

Off-Balance Sheet Transactions

We currently have no off-balance sheet arrangements that have or are reasonably likely to have a current or future material effect on our financial condition, changes in financial condition, revenues or expenses, results of operations, liquidity, capital expenditures or capital resources.

Item 3. Quantitative and Qualitative Disclosures About Market Risk

As a smaller reporting company, we are electing scaled disclosure reporting obligations and therefore are not required to provide the information required by this Item.

Item 4. Controls and Procedures

(a) ***Evaluation of Disclosure Controls and Procedures***. We maintain "disclosure controls and procedures," as such term is defined in Rule 13a-15(e) under the Securities Exchange Act of 1934, or the Exchange Act, that are designed to ensure that information required to be disclosed by us in reports that we file or submit under the Exchange Act is recorded, processed, summarized, and reported within the time periods specified in Securities and Exchange Commission rules and forms, and that such information is accumulated and communicated to our management, including our Chief Executive Officer and Chief Financial Officer, as appropriate, to allow timely decisions regarding required disclosure. In designing and evaluating our disclosure controls and procedures, our management recognized that disclosure controls and procedures, no matter how well conceived and operated, can provide only reasonable, not absolute, assurance that the objectives of the disclosure controls and procedures are met. Accordingly, our disclosure controls and procedures have been designed to meet reasonable assurance standards. Additionally, in designing disclosure controls and procedures, our management was necessarily required to apply its judgment in evaluating the cost-benefit relationship of possible disclosure controls and procedures. The design of any disclosure controls and procedures is also 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.

Under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, we have evaluated the effectiveness of our disclosure controls and procedures as required by Exchange Act Rule 13a-15(b) as of the end of the period covered by this report. Based on that evaluation, our Chief Executive Officer and Chief Financial Officer have concluded that these disclosure controls and procedures are effective at the reasonable assurance level as of June 30, 2015.

(b) ***Changes in Internal Controls.*** There were no changes in our internal control over financial reporting that occurred during the quarter ended June 30, 2015 that has materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

PART II — OTHER INFORMATION

Item 1. Legal Proceedings

On occasion, we may be involved in legal matters arising in the ordinary course of our business including matters involving proprietary technology. While management believes that such matters are currently insignificant, matters arising in the ordinary course of business for which we are or could become involved in litigation may have a material adverse effect on our business, financial condition or results of comprehensive loss.

On November 13, 2014, we received a letter from Exeltis USA Dermatology, Inc. formerly known as Quinnova Pharmaceuticals, Inc., and herein referred to as "Exeltis", claiming that we breached its Exclusive Sales and Distribution Agreement with Exeltis. Specifically, Exeltis claimed that the marketing and selling of its Alevicyn gel product violates the terms of the Exclusive Sales and Distribution Agreement and demanded that we cease and desist from any further marketing or sales. We believe that the marketing and selling of its Alevicyn gel is not in violation of the Exclusive Sales and Distribution Agreement and that the claims made by Exeltis are without merit. Exeltis continues to purchase products from us under a new, non-exclusive distribution agreement for sale to their customers under their own brand. We intend to defend this matter vigorously and does not believe an accrual for a potential loss relating to this matter is necessary at this time. While we believe this claim is without merit, there can be no assurances provided that the outcome of this matter will be favorable to us or will not have a negative impact on its consolidated financial position or results from operations.

Item 1A. Risk Factors

There have been no material changes from risk factors previously disclosed in our annual report on Form 10-K for the fiscal year ended March 31, 2015, as filed with the SEC on June 16, 2015.

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

On June 17, 2015 we issued 135,485 shares of common stock to Advocos, LLC as compensation for services provided, and such shares were valued at $203,000. We relied on the Section 4(a)(2) exemption from securities registration under the federal securities laws for transactions not involving any public offering. No advertising or general solicitation was employed in offering the securities. The securities were issued to an accredited investor. The securities were offered for investment purposes only and not for the purpose of resale or distribution. The transfer thereof was appropriately restricted by us.

Item 3. Default Upon Senior Securities

We did not default upon any senior securities during the quarter ended June 30, 2015.

Item 4. Mine Safety Disclosures

Not applicable.

Item 5. Other Information

On June 1, 2015, we granted an aggregate of 32,133 stock options to our four non-employee directors in lieu of cash. The options have an exercise price of $0.90 and were vested at the date of grant. In addition, our non-employee directors received an aggregate of approximately $26,000 of cash compensation. The compensation was paid pursuant to our Non-Employee Director Compensation Plan as compensation for services provided during the three months ended March 31, 2015.

On June 29, 2015, our stockholders approved a reverse stock split of our outstanding common stock and a proportional decrease of the total number of shares that we are authorized to issue at a whole number ratio in the range of 1-for-5 to 1-for-9, such ratio to be determined in the discretion of our Board of Directors, and authorized our Board of Directors to effect the reverse stock split, if in their judgment it is necessary, at any time until June 29, 2016, upon which date the resolution lapses.

Pursuant to an At-the-Market Issuance Sales Agreement with MLV & Co. LLC dated April 2, 2014, we may issue and sell shares of common stock having an aggregate offering price of up to $9,159,000 from time to time through MLV acting as our sales agent. During the three months ended June 30, 2015, we sold 500,000 shares of common stock for gross proceeds of $907,000 and net proceeds of $879,000 after deducting commissions and other offering expenses. Subsequent to June 30, 2015, through August 4, 2015, we sold 724,100 shares of common stock for gross proceeds of $1,157,000 and net proceeds of $1,121,000 after deducting commissions and other offering expenses. We have sold an aggregate 1,692,034 shares of common stock through August 4, 2015, for gross proceeds of 3,507,000 and net proceeds of $3,341,000 after deducting commissions and other offering expenses. We pay MLV a commission rate equal to 3.0% of the gross proceeds from the sale of any shares of common stock sold through MLV as agent.

Item 6. Exhibits

Exhibit Index

Exhibit No.	Description
3.1	Restated Certificate of Incorporation of Oculus Innovative Sciences, Inc. (included as Exhibit 3.1 of the Company's Annual Report on Form 10-K filed June 20, 2007, and incorporated herein by reference).
3.2	Certificate of Amendment of Restated Certificate of Incorporation of Oculus Innovative Sciences, Inc. (included as Exhibit A in the Company's Definitive Proxy Statement on Schedule 14A filed July 21, 2008, and incorporated herein by reference).
3.3	Amended and Restated Bylaws, as Amended of Oculus Innovative Sciences, Inc., effective November 3, 2010 (included as Exhibit 3.3 to the Company's Quarterly Report on Form 10-Q filed November 4, 2010, and incorporated herein by reference).
3.4	Certificate of Amendment of Restated Certificate of Incorporation of Oculus Innovative Sciences, Inc., as amended (included as Exhibit 3.1 to the Company's Current Report on Form 8-K filed March 22, 2013, and incorporated herein by reference).
3.5	Certificate of Amendment of Restated Certificate of Incorporation of Oculus Innovative Sciences, Inc., as amended (included as Exhibit 3.1 to the Company's Current Report on Form 8-K filed December 8, 2014, and incorporated herein by reference).
4.1	Specimen Common Stock Certificate (included as Exhibit 4.1 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
4.2	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 4.4 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
4.3	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 4.5 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
4.4	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 4.11 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
4.5	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 4.12 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
4.6	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 10.3 to the Company's Current Report on Form 8-K filed August 13, 2007, and incorporated herein by reference).
4.7	Form of Warrant to Purchase Common Stock of Oculus Innovative Sciences, Inc. (included as Exhibit 4.1 to the Company's Current Report on Form 8-K filed March 28, 2008, and incorporated herein by reference).
4.8	Warrant issued to Dayl Crow, dated March 4, 2009 (included as Exhibit 4.16 to the Company's Annual Report on Form 10-K filed June 11, 2009, and incorporated herein by reference).
4.9	Form of Common Stock Purchase Warrant for April 2009 offering (included as Exhibit 4.15 to the Company's Registration Statement on Form S-1 (File No. 333-158539) declared effective on July 24, 2009, and incorporated herein by reference).
4.10	Form of Common Stock Purchase Warrant for July 2009 offering (included as Exhibit 4.15 to the Company's Registration Statement on Form S-1 (File No. 333-158539), as amended, declared effective on July 24, 2009, and incorporated herein by reference).
4.11	Certificate of Designation of Preferences, Rights and Limitations of Series A 0% Convertible Preferred Stock, filed with the Delaware Secretary of State on April 24, 2012 (included as Exhibit 4.2 to the Company's Current Report on Form 8-K, filed

April 25, 2012, and incorporated herein by reference).

4.12	Form of Common Stock Purchase Warrant for April 2012 offering (included as Exhibit 4.1 to the Company's Current Report on Form 8-K, filed April 25, 2012, and incorporated herein by reference).
4.13	Form of Underwriters Warrant to be issued to the Underwriters in connection with the March 2013 Offering (included as Exhibit 4.1 to the Company's Current Report on Form 8-K, filed March 7, 2013, and incorporated herein by reference).
4.14	Warrant issued to Dawson James Securities, Inc., dated December 9, 2013 (included as exhibit 4.14 to the Company's 10-Q filed February 14, 2014 and incorporated herein by reference).
4.15	Form of Series A Common Stock Purchase Warrant for February 2014 offering (included as exhibit 4.1 to the Company's Current Report on Form 8-K filed February 26, 2014 and incorporated herein by reference).
4.16	Form of Series B Common Stock Purchase Warrant for February 2014 offering (included as exhibit 4.2 to the Company's Current Report on Form 8-K filed February 26, 2014 and incorporated herein by reference).
4.17	Warrant issued to Dawson James Securities, Inc., dated February 26, 2014 (included as exhibit 4.3 to the Company's Current Report on Form 8-K filed February 26, 2014 and incorporated herein by reference).
4.18	Warrant Agreement, including Form of Warrant entered into by and between Oculus Innovative Sciences, Inc. and Computershare, Inc. and Computershare Trust Company, N.A., dated January 20, 2015 (included as exhibit 4.1 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
4.19	Underwriters Warrant issued to Maxim Partners LLC on January 26, 2015 (included as exhibit 4.2 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
4.20	Underwriters Warrant issued to Robert D. Keyser, Jr. on January 26, 2015 (included as exhibit 4.3 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
4.21	Underwriters Warrant issued to R. Douglas Armstrong on January 26, 2015 (included as exhibit 4.4 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
4.22	Underwriters Warrant issued to Dawson James Securities, Inc. on January 26, 2015 (included as exhibit 4.5 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
4.23	Underwriters Warrant issued to Dawson James Securities, Inc. on January 26, 2015 (included as exhibit 4.6 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
10.1	Form of Indemnification Agreement between Oculus Innovative Sciences, Inc. and its officers and directors (included as Exhibit 10.1 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.2	Office Lease Agreement, dated October 26, 1999, between Oculus Innovative Sciences, Inc. and RNM Lakeville, L.P. (included as Exhibit 10.7 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.3	Amendment No. 1 to Office Lease Agreement, dated September 15, 2000, between Oculus Innovative Sciences, Inc. and RNM Lakeville L.P. (included as Exhibit 10.8 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.4	Amendment No. 2 to Office Lease Agreement, dated July 29, 2005, between Oculus Innovative Sciences, Inc. and RNM Lakeville L.P. (included as Exhibit 10.9 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.5	Amendment No. 3 to Office Lease Agreement, dated August 23, 2006, between Oculus Innovative Sciences, Inc. and RNM Lakeville L.P. (included as Exhibit 10.23 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.6	Office Lease Agreement, dated May 18, 2006, between Oculus Technologies of Mexico, S.A. de C.V. and Antonio Sergio Arturo Fernandez Valenzuela (translated from Spanish) (included as Exhibit 10.10 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.7	Office Lease Agreement, dated July 2003, between Oculus Innovative Sciences, B.V. and Artikona Holding B.V. (translated from Dutch) (included as Exhibit 10.11 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.8	Form of Director Agreement (included as Exhibit 10.20 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.9	Framework Agreement, dated June 16, 2005, by and among Javier Orozco Gutierrez, Quimica Pasteur, S de R.L., Jorge Paulino Hermosillo Martin, Oculus Innovative Sciences, Inc. and Oculus Technologies de Mexico, S.A. de C.V. (included as Exhibit 10.25 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.10	Mercantile Consignment Agreement, dated June 16, 2005, between Oculus Technologies de Mexico, S.A. de C.V., Quimica Pasteur, S de R.L. and Francisco Javier Orozco Gutierrez (included as Exhibit 10.26 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).

10.11	Partnership Interest Purchase Option Agreement, dated June 16, 2005, by and between Oculus Innovative Sciences, Inc. and Javier Orozco Gutierrez (included as Exhibit 10.27 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.12	Termination of Oculus Innovative Sciences, Inc. and Oculus Technologies de Mexico, S.A. de C.V.'s Agreements with Quimica Pasteur, S de R.L. by Jorge Paulino Hermosillo Martin (translated from Spanish) (included as Exhibit 10.28 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.13	Termination of Oculus Innovative Sciences, Inc. and Oculus Technologies de Mexico, S.A. de C.V.'s Agreements with Quimica Pasteur, S de R.L. by Francisco Javier Orozco Gutierrez (translated from Spanish) (included as Exhibit 10.29 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.14	Employment Agreement by and between Oculus Innovative Sciences, Inc. and Hojabr Alimi, dated January 1, 2004 (included as Exhibit 10.14 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.15	Employment Agreement by and between Oculus Innovative Sciences, Inc. and Jim Schutz, dated January 1, 2004 (included as Exhibit 10.15 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.16	Employment Agreement by and between Oculus Innovative Sciences, Inc. and Robert Miller, dated June 1, 2004 (included as Exhibit 10.16 to the Company's Registration Statement on Form S-1 (File No. 333-135584), as amended, declared effective on January 24, 2007, and incorporated herein by reference).
10.17	Amended and Restated Oculus Innovative Sciences, Inc. 2006 Stock Incentive Plan and related form stock option plan agreements (included as Exhibit 10.2 to the Company's Current Report on Form 8-K filed May 2, 2007, and incorporated herein by reference).
10.18	Amendment No. 4 to Office Lease Agreement, dated September 13, 2007, by and between Oculus Innovative Sciences, Inc. and RNM Lakeville L.P. (included as Exhibit 10.43 to the Company's Annual Report on Form 10-K filed June 13, 2008, and incorporated herein by reference).
10.19	Amendment to Office Lease Agreement, effective February 15, 2008, by and between Oculus Innovative Sciences Netherlands B.V. and Artikona Holding B.V. (translated from Dutch) (included as Exhibit 10.44 to the Company's Annual Report on Form 10-K filed June 13, 2008, and incorporated herein by reference).
10.20	Purchase Agreement by and between Oculus Innovative Sciences, Inc. and Robert Burlingame, dated January 26, 2009 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K filed January 29, 2009, and incorporated herein by reference).
10.21	Purchase Agreement by and between Oculus Innovative Sciences, Inc. and Non-Affiliated Investors, dated January 26, 2009 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K filed January 29, 2009, and incorporated herein by reference).
10.22	Revenue Sharing Distribution Agreement by and between Oculus Innovative Sciences, Inc. and VetCure, Inc., dated January 26, 2009 (included as Exhibit 10.3 to the Company's Current Report on Form 8-K filed January 29, 2009, and incorporated herein by reference).
10.23	Purchase Agreement by and between Oculus Innovative Sciences, Inc., Robert Burlingame and Seamus Burlingame, dated February 24, 2009 (included as Exhibit 10.4 to the Company's Current Report on Form 8-K filed February 27, 2009, and incorporated herein by reference).
10.24	Amendment No. 1 to Revenue Sharing Distribution Agreement by and between Oculus Innovative Sciences, Inc. and VetCure, Inc., dated February 24, 2009 (included as Exhibit 10.5 to the Company's Current Report on Form 8-K filed February 27, 2009, and incorporated herein by reference).
10.25	Consultant Agreement by and between Oculus Innovative Sciences, Inc. and Robert C. Burlingame, dated April 1, 2009 (included as Exhibit 10.52 to the Company's Annual Report on Form 10-K filed June 11, 2009, and incorporated herein by reference).
10.26	Microcyn U.S. Commercial Launch Agreement by and between Oculus Innovative Sciences, Inc. and Advocos, dated April 24, 2009 (included as Exhibit 10.53 to the Company's Annual Report on Form 10-K filed June 11, 2009, and incorporated herein by reference).
10.27	Amendment No. 5 to Office Lease Agreement by and between Oculus Innovative Sciences, Inc. and RNM Lakeville, LLC, dated May 18, 2009 (included as Exhibit 10.54 to the Company's Annual Report on Form 10-K filed June 11, 2009, and incorporated herein by reference).
10.28	Engagement Agreement by and between Oculus Innovative Sciences, Inc. and Dawson James Securities, Inc., dated April 10, 2009 (included as Exhibit 10.55 to the Company's Registration Statement on Form S-1 (File No. 333-158539), as amended, declared effective on July 24, 2009, and incorporated herein by reference).
10.29	Amendment and Clarification of Engagement Letter by and between Oculus Innovative Sciences, Inc. and Dawson James Securities, Inc., dated July 2, 2009 (included as Exhibit 10.56 to the Company's Registration Statement on Form S-1 (File No. 333-158539), as amended, declared effective on July 24, 2009, and incorporated herein by reference).
10.30	Second Amendment and Clarification of Engagement Letter by and between Oculus Innovative Sciences, Inc. and Dawson James Securities, Inc., dated July 10, 2009 (included as Exhibit 10.57 to the Company's Registration Statement on Form S-1 (File No. 333-158539), as amended, declared effective on July 24, 2009, and incorporated herein by reference).
10.31†	Warrant Purchase Agreement by and between Oculus Innovative Sciences, Inc. and Dawson James Securities, Inc., dated July 13, 2009 (included as Exhibit 10.58 to the Company's Registration Statement on Form S-1 (File No. 333-158539), as amended, declared effective on July 24, 2009, and incorporated herein by reference).
10.32	Amendment No. 2 to Revenue Sharing, Partnership and Distribution Agreement between Oculus Innovative Sciences, Inc. and Vetericyn, Inc., dated July 24, 2009 (refiled as Exhibit 10.44 to the Company's Quarterly Report on Form 10-Q/A for the quarter ended September 30, 2010 filed April 29, 2011, and incorporated herein by reference).

10.33	Loan and Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing V, Inc., dated May 1, 2010 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K filed May 6, 2010, and incorporated herein by reference).
10.34†	Supplement to the Loan and Security Agreement between Oculus Innovative Sciences, Inc., and Venture Lending & Leasing V, Inc., dated May 1, 2010 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K filed May 6, 2010, and incorporated herein by reference).
10.35†	Amendment No. 3 to Revenue Sharing, Partnership and Distribution Agreement between Oculus Innovative Sciences, Inc. and Vetericyn, Inc., dated June 1, 2010 (refiled as Exhibit 10.44 to the Company's Quarterly Report on Form 10-Q/A for the quarter ended June 30, 2010 filed April 29, 2011, and incorporated herein by reference).
10.36	Amendment No. 1 to Exhibit A to the Revenue Sharing Distribution Agreement and to the Revenue Sharing, Partnership and Distribution Agreement as Revised and Amended, June 1, 2010, dated September 1, 2010 (included as Exhibit 10.46 to the Company's Quarterly Report on Form 10-Q filed November 4, 2010, and incorporated herein by reference).
10.37	Continuous Offering Program Agreement between Oculus Innovative Sciences, Inc. and Rodman & Renshaw, LLC, dated September 3, 2010 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K filed September 17, 2010, and incorporated herein by reference).
10.38†	Purchase Agreement by and between Oculus Innovative Sciences, Inc. and accredited investors, dated February 6, 2009 (refiled as Exhibit 10.32 to the Company's Quarterly Report on Form 10-Q filed November 4, 2010, and incorporated herein by reference).
10.39†	Distribution Agreement between Oculus Innovative Sciences, Inc. and Tianjin Ascent Import and Export Company, Ltd., dated January 28, 2011 (included as Exhibit 10.47 to the Company's Quarterly Report on Form 10-Q filed February 4, 2011, and incorporated herein by reference).
10.40†	Exclusive Sales and Distribution Agreement between Oculus Innovative Sciences, Inc. and Quinnova Pharmaceuticals, Inc., dated February 14, 2011 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K filed February 18, 2011, and incorporated herein by reference).
10.41	Exclusive Co-Promotion Agreement between Oculus Innovative Sciences, Inc. and Quinnova Pharmaceuticals, Inc., dated February 14, 2011 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K filed February 18, 2011, and incorporated herein by reference).
10.42	Product Option Agreement between Oculus Innovative Sciences, Inc. and AmDerma Pharmaceuticals, LLC, dated February 14, 2011 (included as Exhibit 10.3 to the Company's Current Report on Form 8-K filed February 18, 2011, and incorporated herein by reference).
10.43	Amendment No. 6 to Office Lease Agreement by and between Oculus Innovative Sciences, Inc. and RNM Lakeville, L.P., dated April 26, 2011 (included as Exhibit 10.52 to the Company's Annual Report on Form 10-K filed June 3, 2011, and incorporated herein by reference).
10.44	Loan and Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing VI, Inc., dated June 29, 2011 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K filed July 6, 2011, and incorporated herein by reference).
10.45	Supplement to the Loan and Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing VI, Inc., dated June 29, 2011 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K filed July 6, 2011, and incorporated herein by reference).
10.46	Amendment No. 1 to the Loan and Security Agreement and Supplement to the Loan and Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing V, Inc., dated June 29, 2011 (included as Exhibit 10.4 to the Company's Current Report on Form 8-K filed July 6, 2011, and incorporated herein by reference).
10.47	Intellectual Property Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing VI, Inc., dated June 29, 2011 (included as Exhibit 10.5 to the Company's Current Report on Form 8-K filed July 6, 2011, and incorporated herein by reference).
10.48	Intellectual Property Security Agreement between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing V, Inc., dated June 29, 2011 (included as Exhibit 10.6 to the Company's Current Report on Form 8-K filed July 6, 2011, and incorporated herein by reference).
10.49†	Oculus Innovative Sciences, Inc. 2011 Stock Incentive Plan (included as Exhibit A in the Company's Definitive Proxy Statement on Schedule 14A filed July 29, 2011, and incorporated herein by reference).
10.50†	Distribution Agreement between Oculus Innovative Sciences, Inc. and Shanghai Sunvic Technology Co. Ltd., dated June 26, 2011 (included as Exhibit 10.58 to the Company's Quarterly Report on Form 10-Q filed August 4, 2011 and incorporated herein by reference).
10.51	Patent License Agreement-Exclusive between Oculus Innovative Sciences, Inc. and agencies of the United States Public Health Service within the Department of Health and Human Services, dated August 22, 2011 (included as Exhibit 10.60 to the Company's Quarterly Report on Form 10-Q filed November 3, 2011, and incorporated herein by reference).
10.52†	Securities Purchase Agreement by and between the Company and the Purchasers, dated April 22, 2012 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed April 25, 2012, and incorporated herein by reference).
10.53†	Collaboration Agreement between Oculus Innovative Sciences, Inc. and AmDerma Pharmaceuticals, LLC, dated June 21, 2012 (included as Exhibit 10.53 to the Company's Annual Report on Form 10-K filed June 21, 2012 and incorporated herein by reference).
10.54†	License, Exclusive Distribution and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Oculus Technologies of Mexico, S.A. de C.V., and, More Pharma Corporation, S. de R.L. de C.V., dated August 9, 2012 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed August 15, 2012, and incorporated herein by reference).

10.55	Exclusive Distribution and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Oculus Technologies of Mexico, S.A. de C.V., and, More Pharma Corporation, S. de R.L. de C.V., dated August 9, 2012 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K, filed August 15, 2012, and incorporated herein by reference).
10.56	Amendment No. 7 to Office Lease Agreement by and between Oculus Innovative Sciences, Inc. and 1125-1137 North McDowell, LLC, dated October 10, 2012 (included as Exhibit 10.58 to the Company's Quarterly Report on Form 10-Q filed November 8, 2012, and incorporated herein by reference).
10.57	Stock Purchase Agreement by and between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing V, LLC and Venture Lending & Leasing VI, LLC, dated October 30, 2012 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed November 1, 2012, and incorporated herein by reference).
10.58	Letter Agreement by and between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing V, Inc., dated October 30, 2012 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K, filed November 1, 2012, and incorporated herein by reference).
10.59	Letter Agreement by and between Oculus Innovative Sciences, Inc. and Venture Lending & Leasing VI, Inc., dated October 30, 2012 (included as Exhibit 10.3 to the Company's Current Report on Form 8-K, filed November 1, 2012, and incorporated herein by reference).
10.60	Side Letter Agreement to the Stock Purchase Agreement dated April 22, 2012 by and between Oculus Innovative Sciences, Inc., on one hand, and Sabby Healthcare Volatility Master Fund, Ltd. and Sabby Volatility Warrant Master Fund, Ltd. on the other hand, dated October 29, 2012 (included as Exhibit 10.4 to the Company's Current Report on Form 8-K, filed November 1, 2012, and incorporated herein by reference).
10.61	Offer of Employment Letter between Oculus Innovative Sciences, Inc. and Sameer Harish, effective as of February 1, 2013 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed February 4, 2013, and incorporated herein by reference).
10.62	Employment Agreement by and between Ruthigen, Inc. and Hojabr Alimi, dated March 21, 2013 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed March 22, 2013, and incorporated herein by reference).
10.63	License and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated May 23, 2013 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed June 7, 2013, and incorporated herein by reference).
10.64	Shared Services Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated May 23, 2013 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K, filed June 7, 2013, and incorporated herein by reference).
10.65	Amendment to Offer of Employment Letter between Oculus Innovative Sciences, Inc. and Sameer Harish, dated May 23, 2013 (included as Exhibit 10.4 to the Company's Current Report on Form 8-K, filed June 7, 2013, and incorporated herein by reference).
10.66	Employment Agreement by and between Oculus Innovative Sciences, Inc. and Jim Schutz, dated June 20, 2013 (filed as Exhibit 10.68 to the Company's Annual Report on Form 10-K, filed June 25, 2013 and incorporated herein by reference).
10.67	Employment Agreement by and between Oculus Innovative Sciences, Inc. and Robert Miller, dated June 20, 2013 (filed as Exhibit 10.69 to the Company's Annual Report on Form 10-K, filed June 25, 2013 and incorporated herein by reference).
10.68	Separation Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated August 2, 2013 (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed August 8, 2013 and incorporated herein by reference).
10.69	Amendment No. 1 to License and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated October 9, 2013 (included as exhibit 10.64 to the Company's 10-Q filed November 19, 2013 and incorporated herein by reference).
10.70	Amendment No. 2 to License and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated November 6, 2013 (included as exhibit 10.65 to the Company's 10-Q filed November 19, 2013 and incorporated herein by reference).
10.71	Letter Agreement by and between Oculus Innovative Sciences, Inc., Venture Lending & Leasing V, Inc., and Venture Lending & Leasing VI, Inc., dated November 6, 2013 (filed as exhibit 10.66 to the Company's 10-Q filed November 19, 2013 and incorporated herein by reference).
10.72	Form of Securities Purchase Agreement by and between Oculus Innovative Sciences, Inc. and the Purchasers, dated December 4, 2013 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed December 6, 2013, and incorporated herein by reference).
10.73	Funding Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated January 31, 2014 (included as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed February 6, 2014, and incorporated herein by reference).
10.74	Amended Separation Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated January 31, 2014 (included as Exhibit 10.2 to the Company's Current Report on Form 8-K, filed February 6, 2014, and incorporated herein by reference).
10.75	Amendment No. 3 to License and Supply Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated January 31, 2014 (included as exhibit 10.3 to the Company's Current Report on Form 8-K filed February 6, 2014 and incorporated herein by reference).
10.76	Amendment No. 1 to Shared Services Agreement by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc., dated January 31, 2014 (included as exhibit 10.4 to the Company's Current Report on Form 8-K filed February 6, 2014).
10.77	Letter Agreement by and between Oculus Innovative Sciences, Inc. and Hojabr Alimi, dated January 31, 2014 (included as exhibit 10.6 to the Company's Current Report on Form 8-K filed February 6, 2014).
10.78	Form of Securities Purchase Agreement by and between Oculus Innovative Sciences, Inc. and the Purchasers, dated February 21, 2014 (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed February 26, 2014 and incorporated herein by reference).

10.79	At-the-Market Issuance Sales Agreement, dated April 2, 2014, by and between Oculus Innovative Sciences, Inc. and MLV & Co. LLC (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed April 2, 2014 and incorporated herein by reference).
10.80	Lease Agreement by and between Oculus Innovative Sciences, Inc. and 2500 Investors, Inc., dated July 9, 2014.
10.81	Securities Purchase Agreement, dated January 8, 2015, by and between Oculus Innovative Sciences, Inc. and two investors, Ruthigen, Inc. and Dawson James Securities, Inc. (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed January 12, 2015 and incorporated herein by reference).
10.82	Underwriting Agreement entered into by and between Oculus Innovative Sciences, Inc. and Maxim Group LLC as representative of the underwriters named on Schedule A thereto, dated January 20, 2015 (included as exhibit 1.1 to the Company's Current Report on Form 8-K filed January 26, 2015 and incorporated herein by reference).
10.83†	Sales Representation Contract, dated February 1, 2015, by and between Oculus Innovative Sciences, Inc. and SLA Brands, Inc. (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed March 2, 2015 and incorporated herein by reference).
10.84	Securities Purchase Follow-Up Agreement, dated March 13, 2015, by and between Oculus Innovative Sciences, Inc., two investors, Ruthigen, Inc. and Dawson James Securities, Inc. (included as exhibit 10.1 to the Company's Current Report on Form 8-K filed March 16, 2015 and incorporated herein by reference).
10.85	Securities Purchase Agreement, dated March 13, 2015, by and between Oculus Innovative Sciences, Inc., several investors, Ruthigen, Inc. and Dawson James Securities, Inc. (included as exhibit 10.2 to the Company's Current Report on Form 8-K filed March 16, 2015 and incorporated herein by reference).
10.86	Agreement, dated March 13, 2015, by and between Oculus Innovative Sciences, Inc. and Pulmatrix, Inc. (included as exhibit 10.3 to the Company's Current Report on Form 8-K filed March 16, 2015 and incorporated herein by reference).
10.87	Agreement, dated March 13, 2015, by and between Oculus Innovative Sciences, Inc. and Ruthigen, Inc. (included as exhibit 10.4 to the Company's Current Report on Form 8-K filed March 16, 2015 and incorporated herein by reference).
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.
32.1*	Certification of Officers 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.
101.CAL*	XBRL Taxonomy Extension Calculation Linkbase.
101.DEF*	XBRL Taxonomy Extension Definition Linkbase.
101.LAB*	XBRL Taxonomy Extension Label Linkbase.
101.PRE*	XBRL Taxonomy Extension Presentation Linkbase.

* Filed herewith.

† Confidential treatment has been granted with respect to certain portions of this agreement.

Copies of above exhibits not contained herein are available to any stockholder, upon payment of a reasonable per page fee, upon written request to: Chief Financial Officer, Oculus Innovative Sciences, Inc., 1129 N. McDowell Blvd., Petaluma, California 94954.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

OCULUS INNOVATIVE SCIENCES, INC.

Date: August 7, 2015

By: /s/ Jim Schutz
Jim Schutz
Chief Executive Officer
(Principal Executive Officer)

Date: August 7, 2015

By: /s/ Robert Miller
Robert Miller
Chief Financial Officer
(Principal Financial Officer and Principal Accounting Officer)

Exhibit 31.1

CERTIFICATION OF CHIEF EXECUTIVE OFFICER PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002 (18 U.S.C. SECTION 1350)

I, Jim Schutz, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Oculus Innovative Sciences, Inc. for the quarter ended June 30, 2015;

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)) and internal control 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 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 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 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; and

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):

(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.

By: /s/ Jim Schutz
Jim Schutz
Chief Executive Officer
(Principal Executive Officer)

Date: August 7, 2015

Exhibit 31.2

CERTIFICATION OF CHIEF FINANCIAL OFFICER PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002 (18 U.S.C. SECTION 1350)

I, Robert Miller, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Oculus Innovative Sciences, Inc. for the quarter ended June 30, 2015;

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)) and internal control 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 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 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 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; and

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):

(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.

By: /s/ Robert Miller
Robert Miller
Chief Financial Officer
(Principal Financial Officer and Principal Accounting Officer)

Date: August 7, 2015

Exhibit 32.1

CERTIFICATION PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002
(18 U.S.C. SECTION 1350)

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 Oculus Innovative Sciences, Inc., a Delaware corporation (the "Company"), do hereby certify, to such officers' knowledge, that:

The Quarterly Report on Form 10-Q for the quarter ended June 30, 2015 (the "Form 10-Q") of the Company fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, and the information contained in the Form 10-Q fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: August 7, 2015

By: /s/ Jim Schutz
Jim Schutz
Chief Executive Officer
(Principal Executive Officer)

Date: August 7, 2015

By: /s/ Robert Miller
Robert Miller
Chief Financial Officer
(Principal Financial Officer and Principal Accounting Officer)