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

FORM 10-Q

(Mark one)

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

For the quarterly period ended December 31, 2012

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

For the transition period from to

Commission File No: 0-11740

MESA LABORATORIES, INC.

(Exact name of registrant as specified in its charter)

Colorado
(State or other jurisdiction of
incorporation or organization)

84-0872291
(I.R.S. Employer
Identification number)

12100 West Sixth Avenue
Lakewood, Colorado
(Address of principal executive offices)

80228
(Zip Code)

Registrant's telephone number, including area code: **(303) 987-8000**

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

Indicate by check mark whether the registrant has submitted electronically and posted on its corporate Web site, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (Section 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 definitions of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act. (Check one):

Large accelerated filer

Accelerated filer

Non-accelerated filer
(Do not check if a smaller reporting company)

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

Indicate the number of shares outstanding of each of the Issuer's classes of common stock, as of the latest practicable date:

There were 3,379,518 shares of the Issuer's common stock, no par value, outstanding as of January 31, 2013.

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Certification of Chief Executive Officer Pursuant to Rule 13a-14(a)

Certification of Chief Financial Officer Pursuant to Rule 13a-14(a)

Certification of Chief Executive Officer Pursuant to Rule 13a-14(b) and 18 U.S.C. Section 1350

Certification of Chief Financial Officer Pursuant to Rule 13a-14(b) and 18 U.S.C. Section 1350

Part I. Financial Information**Item 1. Financial Statements**

Mesa Laboratories, Inc.
Condensed Balance Sheets
(In thousands, except share amounts)

	December 31, 2012 (Unaudited)	March 31, 2012
ASSETS		
Current assets:		
Cash and cash equivalents	\$ 2,722	\$ 7,191
Accounts receivable, net	7,595	6,486
Inventories, net	5,281	4,438
Prepaid expenses and other	960	1,046
Total current assets	<u>16,558</u>	<u>19,161</u>
Property, plant and equipment, net	7,328	7,266
Intangibles and other, net	16,091	9,819
Goodwill	<u>23,640</u>	<u>14,450</u>
Total assets	<u>\$ 63,617</u>	<u>\$ 50,696</u>
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Accounts payable	\$ 883	\$ 573
Accrued salaries and payroll taxes	2,219	2,134
Other accrued expenses	488	504
Income taxes payable	269	1,051
Total current liabilities	<u>3,859</u>	<u>4,262</u>
Deferred income taxes payable	2,519	2,519
Long-term debt	5,000	—
Contingent consideration	2,140	—
Total liabilities	<u>13,518</u>	<u>6,781</u>
Stockholders' equity:		
Common stock, no par value; authorized 25,000,000 shares; issued and outstanding, 3,370,819 shares (December 31, 2012) and 3,321,965 shares (March 31, 2012)	7,613	6,699
Employee loans to purchase stock	(226)	(396)
Retained earnings	<u>42,712</u>	<u>37,612</u>
Total stockholders' equity	<u>50,099</u>	<u>43,915</u>
Total liabilities and stockholders' equity	<u>\$ 63,617</u>	<u>\$ 50,696</u>

See accompanying notes to condensed financial statements.

Mesa Laboratories, Inc.
Condensed Statements of Income
(Unaudited)
(In thousands except per share data)

	<u>Three months ended December 31,</u>		<u>Nine months ended December 31,</u>	
	<u>2012</u>	<u>2011</u>	<u>2012</u>	<u>2011</u>
Revenues	\$ 11,361	\$ 9,650	\$ 33,627	\$ 28,648
Cost of revenues	4,414	3,765	12,976	11,601
Gross profit	<u>6,947</u>	<u>5,885</u>	<u>20,651</u>	<u>17,047</u>
Operating expenses				
Selling	1,223	949	3,297	2,926
General and administrative	2,759	1,432	6,782	3,924
Research and development	501	342	1,399	1,056
Total operating expenses	<u>4,483</u>	<u>2,723</u>	<u>11,478</u>	<u>7,906</u>
Operating income	2,464	3,162	9,173	9,141
Other expense, net	38	31	109	130
Earnings before income taxes	2,426	3,131	9,064	9,011
Income taxes	883	1,144	3,173	3,291
Net income	<u>\$ 1,543</u>	<u>\$ 1,987</u>	<u>\$ 5,891</u>	<u>\$ 5,720</u>
Net income per share:				
Basic	\$ 0.46	\$ 0.60	\$ 1.76	\$ 1.74
Diluted	0.44	0.57	1.67	1.66
Weighted average common shares outstanding:				
Basic	3,360	3,286	3,349	3,280
Diluted	3,542	3,498	3,527	3,454

See accompanying notes to condensed financial statements.

Mesa Laboratories, Inc.
Condensed Statements of Cash Flows
(Unaudited)
(In thousands)

	<u>Nine months ended December 31,</u>	
	<u>2012</u>	<u>2011</u>
Cash flows from operating activities:		
Net income	\$ 5,891	\$ 5,720
Depreciation and amortization	2,585	1,636
Stock based compensation	897	349
Change in assets and liabilities, net of acquisitions		
Accounts receivable, net	(631)	914
Inventories, net	67	857
Prepaid expenses and other	114	134
Accounts payable	310	(119)
Accrued liabilities	(782)	(582)
Net cash flows provided by operating activities	<u>8,451</u>	<u>8,909</u>
Cash flows from investing activities:		
Acquisition of net assets of a business	(16,660)	(712)
Purchases of property, plant and equipment	(656)	(545)
Net cash used in investing activities	<u>(17,316)</u>	<u>(1,257)</u>
Cash flows from financing activities:		
Proceeds from the issuance of debt	11,000	—
Payments on debt	(6,000)	(4,750)
Dividends	(1,341)	(1,216)
Purchases of stock for treasury	(57)	(60)
Proceeds from the exercise of stock options	794	519
Net cash provided by (used in) financing activities	<u>4,396</u>	<u>(5,507)</u>
Net (decrease) increase in cash and cash equivalents	<u>(4,469)</u>	<u>2,145</u>
Cash and cash equivalents at beginning of period	<u>7,191</u>	<u>3,546</u>
Cash and cash equivalents at end of period	<u>\$ 2,722</u>	<u>\$ 5,691</u>
Cash paid for:		
Income taxes	\$ 4,125	\$ 3,280
Interest	120	134
Supplemental non-cash activity:		
Employee loans issued for the exercise of stock options	\$ 177	\$ 235
Repayment of employee loans related to the exercise of stock options	347	301
Contingent consideration as part of the acquisition of net assets of a business	\$ 2,140	\$ —

See accompanying notes to condensed financial statements.

Mesa Laboratories, Inc. Notes to Condensed Financial Statements

Note 1 - Description of Business and Summary of Significant Accounting Policies

Description of Business

Mesa Laboratories, Inc. (we, us, our, the “Company” or “Mesa”) was incorporated under the laws of the State of Colorado on March 26, 1982. We pursue a strategy of focusing primarily on quality control products, which are sold into niche markets that are driven by regulatory requirements. We prefer markets that have limited competition where we can establish a commanding presence and achieve high gross margins. We are organized into two divisions across four physical locations. Our Instruments division designs, manufactures and markets quality control instruments and disposable products utilized in connection with the healthcare, pharmaceutical, food and beverage, medical device, industrial hygiene, semiconductor and petrochemical industries. Our Biological Indicators division manufactures and markets biological indicators and distributes chemical indicators used to assess the effectiveness of sterilization processes, including steam, gas, hydrogen peroxide and radiation, in the hospital, dental, medical device and pharmaceutical industries.

Basis of Presentation

The accompanying condensed balance sheet as of March 31, 2012, has been derived from audited financial statements. The accompanying unaudited interim condensed financial statements of Mesa have been prepared on the same basis as the annual audited financial statements and in accordance with accounting principles generally accepted in the United States (“US GAAP”) for interim financial information and the rules and regulations of the Securities and Exchange Commission (“SEC”) for interim financial statements. In the opinion of management, such unaudited information includes all adjustments (consisting only of normal recurring accruals) necessary for a fair presentation of this interim information. Operating results and cash flows for interim periods are not necessarily indicative of results that can be expected for the entire year. The information included in this report should be read in conjunction with our audited financial statements and notes thereto included in our Annual Report on Form 10-K for the year ended March 31, 2012.

On October 1, 2012 our articles of incorporation were amended to increase the number of authorized shares of common stock from 8,000,000 to 25,000,000.

Approximately \$390,000 and \$1,210,000, respectively, of customer payments for shipping services were reclassified from cost of revenues to revenues in the condensed statements of income for the three and nine months ended December 31, 2011. This reclassification affected revenues and cost of revenues, but had no other impact on amounts in the accompanying condensed statements of income.

The summary of our significant accounting policies is incorporated by reference to our annual report on Form 10-K as of March 31, 2012.

Recently Issued Accounting Pronouncements

In July 2012, the Financial Accounting Standards Board (“FASB”) issued Accounting Standards Update (“ASU”) No. 2012-02, “Testing Indefinite-Lived Intangible Assets for Impairment.” We do not have indefinite-lived intangible assets, as a result, this guidance does not have an impact on our financial statements or disclosures.

In October 2012, the FASB issued ASU 2012-04, “Technical Corrections and Improvements.” This standard includes: 1) source literature amendments to conform the language between current accounting literature and legacy source literature; 2) clarification of guidance and reference corrections; and 3) relocation of guidance to a more appropriate location. The adoption of this standard did not have an impact on our financial statements and disclosures.

Note 2 - Acquisition of Net Assets of a Business

On May 15, 2012, we completed a business combination (the “Bios Acquisition”) by acquiring specific assets and assuming certain liabilities of Bios International Corporation (“Bios”), a New Jersey corporation. The asset acquisition agreement (the “Bios Agreement”) includes a provision for contingent consideration based on revenue growth over a three year earn-out period. The Bios Acquisition further diversifies and grows our Instruments segment, and we believe that it will maintain our historic profitability measures.

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The contingent consideration arrangement requires us to pay Bios if cumulative revenues related to the acquisition for the three years subsequent to the acquisition exceed \$22,127,000. The potential undiscounted future payment that we could be required to make ranges from \$0 to \$6,710,000. The fair value of the contingent consideration arrangement included in the purchase price below was estimated based on the historic revenue growth rates of Bios.

We expect to achieve significant savings and income growth as we integrate the Bios operations and marketing functions. These factors, among others, contributed to a purchase price in excess of the estimated fair value of Bios' net identifiable assets and, as a result, we recorded goodwill in connection with this transaction. The goodwill is expected to be deductible for tax purposes. All of the goodwill was assigned to our Instruments segment.

The Bios Acquisition constitutes the acquisition of a business and was recognized at fair value. We determined the estimated fair values using discounted cash flow analyses and estimates made by management. The following reflects our allocation of the consideration, subject to customary purchase price adjustments in accordance with the Bios Agreement (in thousands):

Cash consideration	\$	16,660
Contingent purchase price liability		2,140
Aggregate consideration	\$	<u>18,800</u>
The purchase price was allocated as follows:		
Accounts receivable, net	\$	478
Inventory		910
Other current assets		28
Property, plant and equipment		63
Intangibles		8,200
Goodwill		9,190
Current liabilities		(69)
Total purchase price allocation	\$	<u>18,800</u>

The accompanying condensed statements of income include the results of the Bios Acquisition from the acquisition date of May 15, 2012. The pro forma effects of the acquisition on the results of operations as if the acquisition had been completed on April 1, 2012 and 2011, are as follows (in thousands, except per share data):

	Nine months ended December 31,	
	2012	2011
Total net revenues	\$ 34,408	\$ 34,036
Net income	5,912	5,863
Net income per common share:		
Basic	\$ 1.77	\$ 1.79
Diluted	1.68	1.70

The above pro forma results include adjustments for amortization of acquired intangibles, interest expense and income tax expense. The pro forma information as presented above is for informational purposes only and is not necessarily indicative of results of operations that would have been achieved if the acquisition had taken place at the dates identified.

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Note 3 - Inventories

Inventories consist of the following (in thousands):

	December 31, 2012	March 31, 2012
Raw materials	\$ 3,890	\$ 3,242
Work-in-process	446	331
Finished goods	1,360	1,090
Less: reserve	(415)	(225)
	<u>\$ 5,281</u>	<u>\$ 4,438</u>

Note 4 - Long-Term Debt

Long-term debt consists of the following (in thousands):

	December 31, 2012	March 31, 2012
Line of credit (1.5% at December 31, 2012)	\$ 5,000	\$ —
Less: current portion	—	—
Long-term portion	<u>\$ 5,000</u>	<u>\$ —</u>

In February 2012, we entered into a three year agreement (the “Credit Facility”) for a \$20,000,000 revolving line of credit (“Line of Credit”) and up to \$1,000,000 of letters of credit, maturing in February 2015. Funds from the Credit Facility may be used for general working capital and corporate needs, retiring existing debt, or to support acquisitions and capital expenditures.

Under the Credit Facility, indebtedness bears interest at either: (1) LIBOR, as defined, plus an applicable margin ranging from 1.25% to 2.00%; or (2) the bank’s commercial bank floating rate (“CBFR”), which is the greater of the bank’s prime rate or one month LIBOR + 2.50%, adjusted down, from 1.25% to 0.50%. We elect the interest rate with each borrowing under the line of credit. In addition, there is an unused capacity fee of 0.15% to 0.30%. The adjustments and unused capacity fee depend on the ratio of funded debt to our trailing four quarters of EBITDA, as defined, with four tiers ranging from a ratio of less than one to greater than two. Letter of credit fees are based on the applicable LIBOR rate.

The Credit Facility is secured by all of our assets and requires us to maintain a ratio of funded debt to our trailing four quarters of EBITDA, as defined, of 2.5 to 1.0, and a minimum fixed charge coverage ratio of 1.5 to 1.0. We were in compliance with these covenants at December 31, 2012.

In order to facilitate the Bios Acquisition, in May 2012 we borrowed \$11,000,000 under the terms of the Line of Credit. During the three and nine months ended December 31, 2012 we made principal repayments of \$3,000,000 and \$6,000,000, respectively. As a result, the amount outstanding under the Line of Credit was \$5,000,000 as of December 31, 2012. In January 2013, we made an additional principal payment of \$1,000,000.

Future contractual maturities of debt are as follows (in thousands):

<u>Year ending March 31,</u>	
2013	\$ —
2014	—
2015	5,000
	<u>\$ 5,000</u>

In April 2010, we entered into a credit facility consisting of: a) 36 month reducing line of credit for \$3,000,000 and maturing at April 27, 2013, requiring quarterly principal payments of \$250,000 beginning July 27, 2010, which was retired in February 2012; and b) revolving line of credit for \$4,000,000 maturing on December 23, 2011, which was retired in December 2011. Both of these lines of credit were subject to a variable rate of interest and a rate floor.

Note 5 - Stock-Based Compensation

Stock-based compensation costs for award grants to employees and directors are valued at fair value and recognized on a straight line basis over the service period for the entire award, with the amount of compensation cost recognized at any date equaling at least the portion of the award that is vested. We estimate forfeiture rates based on historical experience. Amounts recognized in the condensed financial statements related to stock-based compensation are as follows (in thousands, except per share data):

	Three months ended December 31,		Nine months ended December 31,	
	2012	2011	2012	2011
Total cost of stock-based compensation charged against income before income taxes	\$ 459	\$ 157	\$ 897	\$ 349
Amount of income tax benefit recognized in earnings	163	57	314	127
Amount charged against net income	<u>\$ 296</u>	<u>\$ 100</u>	<u>\$ 583</u>	<u>\$ 222</u>
Impact on net income per common share:				
Basic	\$ 0.09	\$ 0.03	\$ 0.17	\$ 0.07
Diluted	0.08	0.03	0.17	0.06

Stock-based compensation expense is included in cost of revenues and general and administrative expense in the accompanying condensed statements of income.

The fair value of each option grant is estimated on the date of grant using the Black-Scholes option-pricing model (“Black-Scholes”). We use historical data to estimate the expected price volatility, the expected option life and expected forfeiture rate. The risk-free interest rate is based on the United States Treasury yield curve in effect at the time of grant for the estimated life of the option. The dividend yield is calculated based upon the dividend payments made during the prior four quarters as a percent of the average stock price for that period. The following summarizes the Black-Scholes assumptions used for option grants:

	Nine months ended December 31,	
	2012	2011
Volatility	29.3%	33.5%
Risk-free interest rate	0.6-1.0%	0.9-2.2%
Expected option life (years)	5-10	5-10
Dividend yield	1.4%	1.8%

The following is a summary of option activity:

	Number of Shares	Weighted- average Exercise Price per Share	Weighted- average Remaining Contractual Term	Aggregate Intrinsic Value (000s)
Outstanding at March 31, 2012	433,785	\$ 22.77	3.9	
Options granted	116,080	50.01	6.0	
Options forfeited	(36,110)	33.04		
Options expired	(40)	18.98		
Options exercised	<u>(63,375)</u>	20.17		
Outstanding at December 31, 2012	<u>450,340</u>	29.33		\$ 9,392
Exercisable at December 31, 2012	180,925	21.30	3.0	5,212

We issue new shares of common stock upon exercise of stock options. The total intrinsic value of options exercised was \$1,800,000 and \$738,000 for the nine months ended December 31, 2012 and 2011, respectively.

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A summary of the status of our unvested option shares as of December 31, 2012 is as follows:

	Number of Shares	Weighted- average Grant-Date Fair Value
Unvested at March 31, 2012	284,875	\$ 7.23
Options granted	116,080	12.45
Options forfeited	(36,110)	8.97
Options vested	(95,430)	6.80
Unvested at December 31, 2012	<u>269,415</u>	9.40

As of December 31, 2012, there was approximately \$2,084,000 of total unrecognized compensation cost related to unvested stock options granted under our plans, which is expected to be recognized over a weighted-average period of 3 years.

Effective November 30, 2012, as part of our Chief Financial Officer transition, 14,400 unvested options were modified to a) extend the expiration date to 10 years following the original grant date, b) allow them to be exercised through their expiration date, and c) accelerate the vesting such that all options will vest by November 30, 2014. This is a modification of the terms of an equity award and, accordingly, we have treated this as an exchange of the original award for a new award. We recorded incremental compensation expense of approximately \$240,000 during the three and nine months ended December 31, 2012, included in general and administrative expense on the accompanying condensed statements of income.

Note 6 - Net Income Per Share

Basic net income per share is computed by dividing net income by the weighted average number of shares of common stock outstanding during the period. Diluted net income per share uses the treasury stock method to compute the weighted average common stock outstanding assuming the conversion of potentially dilutive common shares.

The following table presents a reconciliation of the denominators used in the computation of net income per share - basic and diluted (in thousands, except per share data):

	Three Months Ended December 31,		Nine Months Ended December 31,	
	2012	2011	2012	2011
Net income available for stockholders	\$ 1,543	\$ 1,987	\$ 5,891	\$ 5,720
Weighted average outstanding shares of common stock	3,360	3,286	3,349	3,280
Dilutive effect of stock options	182	212	178	174
Common stock and equivalents	<u>3,542</u>	<u>3,498</u>	<u>3,527</u>	<u>3,454</u>
Net income per share:				
Basic	\$ 0.46	\$ 0.60	\$ 1.76	\$ 1.74
Diluted	0.44	0.57	1.67	1.66

For the three and nine months ended December 31, 2012, 81,000 outstanding stock options were excluded from the calculation of diluted earnings per share because the exercise prices of the stock options were greater than or equal to the average price of the common shares and, therefore, their inclusion would have been anti-dilutive. No stock options were excluded for the three and nine months ended December 31, 2011.

Note 7 - Contingencies

As part of the Bios Acquisition, the Bios Agreement includes a provision for contingent consideration based on revenue growth over a three year earn-out period. The contingent consideration arrangement requires us to pay Bios if the cumulative revenues from the acquisition for the three years subsequent to the acquisition exceed \$22,127,000. The potential undiscounted future payment that we could be required to make ranges from \$0 to \$6,710,000. The fair value of the contingent consideration arrangement included in the purchase price was estimated based on the historic revenue growth of Bios. We recorded a contingent consideration liability of \$2,140,000 on the accompanying condensed balance sheet as of December 31, 2012. Any changes to the contingent consideration ultimately paid would result in additional income or expense on the condensed statements of income. There has been no material change to the contingent consideration liability as of December 31, 2012. The contingent consideration is payable in the first quarter of our year ending March 31, 2016.

During the three months ended December 31, 2012, we determined that we have a potential liability for state sales taxes. The ultimate amount due will depend upon a number of factors, including the amount of sales that were made to customers who already paid the tax or who are exempt, and any penalties and interest. We recorded an estimate of \$100,000, which is included in other accrued expenses on the accompanying condensed balance sheets, and general and administrative expense in the accompanying condensed statements of income. This estimate may change as further analysis is completed and sales tax returns are filed. During the three months ended December 31, 2011, we determined that we had a separate liability for state sales taxes and recorded an estimate of \$250,000. During the three months ended June 30, 2012, we settled the liability and determined that no additional liability was required. We continue to evaluate our exposure in additional states, but at this time the amount of the potential liability, if any, is not estimable.

Note 8 - Segment Information

Our operations are organized into two segments: Biological Indicators and Instruments. The following tables set forth our segment information (in thousands):

	Three Months Ended December 31, 2012			Three Months Ended December 31, 2011		
	Biological Indicators	Instruments	Total	Biological Indicators	Instruments	Total
Revenues	\$ 5,154	\$ 6,207	\$ 11,361	\$ 4,742	\$ 4,908	\$ 9,650
Gross profit	\$ 2,936	\$ 4,011	\$ 6,947	\$ 2,540	\$ 3,345	\$ 5,885
Selling expenses	385	838	1,223	391	558	949
	\$ 2,551	\$ 3,173	5,724	\$ 2,149	\$ 2,787	4,936
Reconciling items (1)			(3,298)			(1,805)
Earnings before income taxes			\$ 2,426			\$ 3,131

	Nine Months Ended December 31, 2012			Nine Months Ended December 31, 2011		
	Biological Indicators	Instruments	Total	Biological Indicators	Instruments	Total
Revenues	\$ 15,590	\$ 18,037	\$ 33,627	\$ 14,600	\$ 14,048	\$ 28,648
Gross profit	\$ 8,890	\$ 11,761	\$ 20,651	\$ 7,756	\$ 9,291	\$ 17,047
Selling expenses	1,165	2,132	3,297	1,194	1,732	2,926
	\$ 7,725	\$ 9,629	17,354	\$ 6,562	\$ 7,559	14,121
Reconciling items (1)			(8,290)			(5,110)
Earnings before income taxes			\$ 9,064			\$ 9,011

(1) Reconciling items include general and administrative, research and development, and other expenses.

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	<u>December 31,</u> <u>2012</u>	<u>March 31,</u> <u>2012</u>
Total assets		
Biological Indicators	\$ 27,392	\$ 28,887
Instruments	32,511	13,572
Corporate and administrative	3,714	8,237
	<u>\$ 63,617</u>	<u>\$ 50,696</u>

All long-lived assets are located in the United States.

Revenues from external customers are attributed to individual countries based upon locations to which the product is shipped or exported, as follows (in thousands):

	<u>Three Months Ended</u> <u>December 31,</u>		<u>Nine Months Ended</u> <u>December 31,</u>	
	<u>2012</u>	<u>2011</u>	<u>2012</u>	<u>2011</u>
Net revenues from unaffiliated customers:				
United States	\$ 7,013	\$ 5,684	\$ 20,504	\$ 17,497
Foreign	4,348	3,966	13,123	11,151
	<u>\$ 11,361</u>	<u>\$ 9,650</u>	<u>\$ 33,627</u>	<u>\$ 28,648</u>

No foreign country exceeds 10% of total revenues.

We have changed our presentation of segments to correspond to the discrete financial information reviewed by our chief operating decision maker and have stated all periods consistently.

Note 9 - Subsequent Event

In February 2013, our Board of Directors declared a quarterly cash dividend of \$0.14 per share of common stock, payable on March 15, 2013, to shareholders of record at the close of business on February 27, 2013.

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

Forward Looking Statements

This report contains information that may constitute "forward-looking statements." Generally, the words "believe," "expect," "intend," "estimate," "anticipate," "project," "will" and similar expressions identify forward-looking statements, which generally are not historical in nature. However, the absence of these words or similar expressions does not mean that a statement is not forward-looking. All statements that address operating performance, events or developments that we expect or anticipate will occur in the future — including statements relating to revenue growth and statements expressing general views about future operating results — are forward-looking statements. Management believes that these forward-looking statements are reasonable as and when made. However, caution should be taken not to place undue reliance on any such forward-looking statements because such statements speak only as of the date when made. We undertake no obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events or otherwise, except as required by law. In addition, forward-looking statements are subject to certain risks and uncertainties that could cause actual results to differ materially from our historical experience and our present expectations or projections. These risks and uncertainties include, but are not limited to, those described in Part II, "Item 1A. Risk Factors" and elsewhere in this report and in our Annual Report on Form 10-K for the year ended March 31, 2012, and those described from time to time in our subsequent reports filed with the Securities and Exchange Commission.

General Discussion

We pursue a strategy of focusing primarily on quality control products, which are sold into niche markets that are driven by regulatory requirements. We prefer markets that have limited competition where we can establish a commanding presence and achieve high gross margins. Mesa Laboratories, Inc. has two divisions — Our Instruments division manufactures and markets quality control instruments and disposable products utilized in connection with the healthcare, pharmaceutical, food and beverage, medical device, industrial hygiene, semiconductor and petrochemical industries. Our Biological Indicators division manufactures and markets biological indicators and distributes chemical indicators used to assess the effectiveness of sterilization processes in the hospital, dental, medical device and pharmaceutical industries. We follow a philosophy of manufacturing a high quality product and providing a high level of on-going service for those products.

Our revenues come from two main sources — products sales, and parts and services. Product sales are dependent on several factors, including general economic conditions, both domestic and international, customer capital spending trends, competition, introduction of new products and acquisitions. Biological indicator products are disposable and are used on a routine basis for quality control, thus product sales are less sensitive to general economic conditions. Instrument products have a longer life, and their purchase by our customers is somewhat discretionary, so sales are more sensitive to general economic conditions. Parts and service demand is driven by our customers' quality control and regulatory environments, which require periodic repair and recalibration or certification of our instrument products. We typically evaluate costs and pricing annually. Our policy is to price our products competitively and, where possible, we try to pass along cost increases in order to maintain our margins. As part of the integration of our previous biological indicator acquisitions we have been adjusting prices to achieve price parity for similar products.

Gross profit is affected by our product mix, manufacturing efficiencies and price competition. Historically, as we have integrated our acquisitions and taken advantage of manufacturing efficiencies, our gross margins for some of the products have improved. There are, however differences in gross margins between different product lines, and ultimately the mix of sales between different segments may continue to impact our overall gross margin.

Selling expense is driven primarily by labor costs, including salaries and commissions. Accordingly, it may vary with sales levels. Labor costs and amortization of intangible assets drive 70-80% of general and administrative expense. Research and development expense is predominantly comprised of labor costs.

In May 2012, we completed the Bios Acquisition by acquiring specific assets and assuming certain liabilities of Bios, a New Jersey corporation. The purchase price for the acquired net assets was \$16,660,000 and potential contingent consideration based on revenue growth over a three year earn-out period. The contingent consideration arrangement requires us to pay Bios if cumulative revenues related to the acquisition for the three years subsequent to the acquisition exceed \$22,127,000. The potential undiscounted future payment that we could be required to make ranges from \$0 to \$6,710,000. We borrowed \$11,000,000 under our Line of Credit to finance the acquisition, with the balance being paid from available cash. Our New Jersey operations were not significantly impacted by Hurricane Sandy.

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Results of Operations

The following table sets forth, for the periods indicated, condensed statements of income data. The table and the discussion below should be read in conjunction with the accompanying condensed financial statements and the notes thereto appearing elsewhere in this report (in thousands, except percent data):

	Three months ended December 31,		Change	Percent Change
	2012	2011		
Revenues	\$ 11,361	\$ 9,650	\$ 1,711	18%
Cost of revenues	4,414	3,765	649	17%
Gross profit	6,947	5,885	1,062	18%
Gross margin	61%	61%	—%	
Operating expenses	4,483	2,723	1,760	65%
Net income	1,543	1,987	(444)	(22)%
Net profit margin	14%	21%	(7)%	

	Nine months ended December 31,		Change	Percent Change
	2012	2011		
Revenues	\$ 33,627	\$ 28,648	\$ 4,979	17%
Cost of revenues	12,976	11,601	1,375	12%
Gross profit	20,651	17,047	3,604	21%
Gross margin	61%	60%	1%	
Operating expenses	11,478	7,906	3,572	45%
Net income	5,891	5,720	171	3%
Net profit margin	18%	20%	(2)%	

Revenues

The following table summarizes our revenues by source (in thousands, except percent data):

	Three months ended December 31,		Change	Percent Change
	2012	2011		
Biological Indicators — Product	\$ 5,154	\$ 4,742	\$ 412	9%
Instruments:				
Product	4,316	3,305	1,011	31%
Parts and disposables	698	719	(21)	(3)%
Service	1,193	884	309	35%
Total Instruments	6,207	4,908	1,299	26%
	\$ 11,361	\$ 9,650	\$ 1,711	18%

	Nine months ended December 31,		Change	Percent Change
	2012	2011		
Biological Indicators — Product	\$ 15,590	\$ 14,600	\$ 990	7%
Instruments:				
Product	12,399	9,084	3,315	36%
Parts and disposables	2,191	2,220	(29)	(1)%
Service	3,447	2,744	703	26%
Total Instruments	18,037	14,048	3,989	28%
	\$ 33,627	\$ 28,648	\$ 4,979	17%

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Instruments revenues for the three and nine months ended December 31, 2012 increased as compared to the prior year as a result of the Bios Acquisition, partially offset by lower revenues from the legacy Instruments product lines. The decrease in revenue from the legacy Instrument product lines is due to cyclical variability on a quarter by quarter basis.

Biological Indicator revenues for the three and nine months ended December 31, 2012 increased as compared to the prior year as a result of continued organic growth, achieved through existing customers and expansion into new markets.

Effective January 1, 2013, we are subject to a 2.3% medical device excise tax on the domestic sales of a majority of our medical instruments and biological indicators. Wherever possible, we renegotiated prices with our customers to recover this additional cost. We can offer no assurance that we will be able to successfully recover the full amounts paid as medical device excise tax.

Gross Profit

The following summarizes our gross profit by source (in thousands, except percent data):

	Three months ended December 31,		Change	Percent Change
	2012	2011		
Biological Indicators	\$ 2,936	\$ 2,540	\$ 396	16%
Instruments	4,011	3,345	666	20%
	<u>\$ 6,947</u>	<u>\$ 5,885</u>	<u>\$ 1,062</u>	18%

	Nine months ended December 31,		Change	Percent Change
	2012	2011		
Biological Indicators	\$ 8,890	\$ 7,756	\$ 1,134	15%
Instruments	11,761	9,291	2,470	27%
	<u>\$ 20,651</u>	<u>\$ 17,047</u>	<u>\$ 3,604</u>	21%

Instruments gross profit for the three and nine months ended December 31, 2012 increased as compared to the prior year as a result of the Bios Acquisition, partially offset by lower gross profit from the legacy Instrument product lines. The decrease in gross profit from the legacy Instrument product lines is due to cyclical variability of sales on a quarter by quarter basis which also affects the absorption of fixed costs in any given quarter.

Biological Indicator gross profit for the three and nine months ended December 31, 2012 increased as compared to the prior year as a result of improved manufacturing efficiencies and increased sales.

[Table of Contents](#)**Operating Expenses**

Operating expenses increased (decreased) as compared to the prior year as follows (in thousands):

	Increase (Decrease)	
	Three months ended December 31, 2012	Nine months ended December 31, 2012
Selling	<u>\$ 274</u>	<u>\$ 371</u>
General and administrative		
Chief Financial Officer transition	526	526
ERP system upgrade and SOX compliance	55	220
Bios Acquisition — professional fees	—	150
Amortization — Bios Acquisition	260	650
Amortization — Trademarks	40	120
Stock option expense	50	230
Sales tax accrual	(150)	(150)
Personnel costs	360	680
Other, net	186	432
	<u>1,327</u>	<u>2,858</u>
Research and development	<u>159</u>	<u>343</u>
Operating expenses	<u>\$ 1,760</u>	<u>\$ 3,572</u>

Selling — The Bios Acquisition resulted in an increase of our selling costs, with negligible increase in other product lines. As a percent of revenues, selling expense remained relatively flat.

General and administrative — As part of our Chief Financial Officer transition, certain unvested options were modified, resulting in incremental stock option expense of approximately \$240,000 for the three and nine months ended December 31, 2012. The balance of the Chief Financial officer transition impact includes a severance package and miscellaneous other costs. All costs associated with the transition were expensed in the three months ended December 31, 2012. We have upgraded our ERP system and implemented computer-based controls as part of our Sarbanes-Oxley compliance efforts, which we believe makes us better prepared for any future growth we may experience. Amortization expense increased due to the Bios Acquisition, in May 2012, and the amortization of trademarks, which began in February 2012. We recorded estimated tax liabilities of \$100,000 and \$250,000, respectively, for the three months ended December 31, 2012 and 2011. Personnel costs increased primarily due to the Bios Acquisition, but also for additional personnel and salary adjustments. The remaining increase primarily consists of general and administrative expenses associated with the acquired Bios operations.

Research and development — The increase is due to additional internal personnel added as a result of the Bios Acquisition and external research and development consulting costs, as we continue our commitment to research and development. The cost of intangible assets that are purchased from others for use in research and development activities and have alternative future uses, however, are capitalized and amortized over their expected useful life. During the year ended March 31, 2012, we capitalized \$175,000 of Biological Indicator research as an intangible asset, as it had alternative future uses, and are amortizing it through research and development expense over ten years. That project is ongoing during our year ending March 31, 2013.

Net Income

Net income varied consistently with the growth in revenues, gross profit and operating expenses. Other expenses, consisting primarily of interest expense, remained relatively flat. Our effective income tax rate decreased period over period as the goodwill associated with the Bios Acquisition is deductible for tax purposes. After filing our tax returns for the year ended March 31, 2012, however, we revised our estimate of the annualized effective income tax rate in the three month period ended December 31, 2012. This resulted in an increase to our effective income tax rate from other previously reported periods in the year ending March 31, 2013. As of January 1, 2013, we will begin paying a 2.3% medical device excise tax on the domestic sales of a majority of our medical instruments and biological indicators.

Liquidity and Capital Resources

Our sources of liquidity may include cash generated from operations, working capital, capacity under our Credit Facility and potential equity and debt offerings. We believe that cash generated from these sources will be sufficient to meet our short-term and long-term needs. Our more significant uses of resources include quarterly dividends to shareholders, payment of debt obligations, long-term capital equipment expenditures and potential acquisitions.

Working capital is the amount by which current assets exceed current liabilities. We had working capital of \$12,699,000 and \$14,899,000, respectively, at December 31, 2012 and March 31, 2012. The decrease in working capital is due to the use of cash for the Bios Acquisition and repayment of long-term debt, offset by cash flows from operations.

In February 2012, we entered into the Credit Facility, which is comprised of a three year agreement for a \$20,000,000 revolving line of credit and up to \$1,000,000 of letters of credit. Funds from the Credit Facility may be used for general working capital and corporate needs, retiring existing debt, or to support acquisitions and capital expenditures. In February 2012, we also extinguished our obligations under our previous debt agreement. In May 2012, we borrowed \$11,000,000 against the Line of Credit to partially finance the Bios Acquisition. At December 31, 2012, we had unused capacity under our Line of Credit of \$15,000,000. In January 2013 we made an additional principal payment of \$1,000,000.

On October 1, 2012, we amended our articles of incorporation to increase the number of authorized shares of common stock from 8 million to 25 million.

We routinely evaluate opportunities for strategic acquisitions. Future material acquisitions may require that we obtain additional capital, assume third party debt or incur other long-term obligations. We have the option to utilize both equity and debt instruments as vehicles for the long-term financing of our investment activities and acquisitions.

On November 7, 2005, our Board of Directors authorized a program to repurchase up to 300,000 shares of our outstanding common stock. Under the plan, the shares may be purchased from time to time in the open market at prevailing prices or in negotiated transactions off the market. Shares purchased will be canceled and repurchases will be made with existing cash reserves. We do not maintain a set policy or schedule for our buyback program. We have purchased 156,412 shares of common stock under this program from inception through December 31, 2012.

We have been paying regular quarterly dividends since 2003. Dividends per share paid by quarter were as follows:

	Year ending March 31,	
	2013	2012
First quarter	\$ 0.13	\$ 0.12
Second quarter	0.13	0.12
Third quarter	0.14	0.13
Fourth quarter	—	0.13

In February 2013, our Board of Directors declared a quarterly cash dividend of \$0.14 per share of common stock, payable on March 15, 2013, to shareholders of record at the close of business on February 27, 2013.

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Cash Flows

Our cash flows from operating, investing and financing activities were as follows (in thousands):

	Nine months ended December 31,	
	2012	2011
Net cash provided by operating activities	\$ 8,451	\$ 8,909
Net cash used in investing activities	(17,316)	(1,257)
Net cash provided by (used in) financing activities	4,396	(5,507)

Net cash provided by operating activities changed primarily due to increases in revenues and corresponding net income, offset by the timing of certain working capital expenditures related to inventory, income taxes, and our expanding international customer base, which has extended payment terms.

Net cash used in investing activities was primarily for the \$16,660,000 Bios Acquisition in May 2012. In addition, purchases of property, plant and equipment were \$656,000 and \$545,000, respectively, for the nine months ended December 31, 2012 and 2011.

Net cash provided by financing activities for the nine months ended December 31, 2012, resulted from borrowings under our Line of Credit of \$11,000,000 and proceeds from the exercise of stock options of \$794,000, partially offset by payments on long-term debt of \$6,000,000 and the payment of dividends of \$1,341,000. Net cash used in financing activities for the nine months ended December 31, 2011, resulted from the repayment of debt of \$4,750,000, payment of dividends of \$1,216,000, partially offset by proceeds from the exercise of stock options of \$519,000.

At December 31, 2012, we had contractual obligations for open purchase orders for routine purchases of supplies and inventory, which would be payable in less than one year. In September 2011, we entered into a license agreement for certain biological indicator technology and up to \$225,000 of additional payments may be made in the future, depending on meeting certain development and performance milestones. As part of our Bios Acquisition, the Bios Agreement includes a provision for contingent consideration based on revenue growth over a three year earn-out period. The contingent consideration arrangement requires us to pay Bios if cumulative revenues from the acquisition for the three years subsequent to the acquisition exceed \$22,127,000. The potential undiscounted future payment that we could be required to make ranges from \$0 to \$6,710,000.

Critical Accounting Estimates

Our condensed financial statements and accompanying notes have been prepared in accordance with U.S. GAAP. The preparation of these financial statements requires management to make estimates, judgments and assumptions that affect reported amounts of assets, liabilities, revenues and expenses. We continually evaluate the accounting policies and estimates used to prepare the condensed financial statements. The estimates are based on historical experience and assumptions believed to be reasonable under current facts and circumstances. Actual amounts and results could differ from these estimates made by management. Certain accounting policies that require significant management estimates and are deemed critical to our results of operations or financial position are discussed in our Annual Report on Form 10-K for the year ended March 31, 2012 in the Critical Accounting Policies and Estimates section of Management's Discussion and Analysis of Financial Condition and Results of Operations.

Item 3. Quantitative and Qualitative Disclosures About Market Risk

We are exposed to a variety of market risks, currently all investments are in dollar denominated accounts, such as money market funds, with variable interest rates. In the normal course of business, we employ established policies and procedures to manage our exposure to changes in the market value of our investments.

Item 4. Controls and Procedures

Evaluation of Disclosure Controls and Procedures

We maintain disclosure controls and procedures (as defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, as amended) that are designed to reasonably ensure that information required to be disclosed by us in the reports we file or submit under the Securities Exchange Act of 1934, as amended, is recorded, processed, summarized and reported within the time periods specified in the Securities and Exchange Commission's rules and forms and that such information is accumulated and communicated to our management, including our principal executive and principal financial officers, or persons performing similar functions, as appropriate to allow timely decisions regarding required disclosure. Our management evaluated, with the participation of our Chief Executive Officer and Chief Financial Officer, the effectiveness of our disclosure controls and procedures as of December 31, 2012. Based on that evaluation, our management concluded that our disclosure controls and procedures were effective at December 31, 2012.

Our management, including our Chief Executive Officer and Chief Financial Officer, is responsible for establishing and maintaining adequate internal control over financial reporting as required by Sarbanes-Oxley (SOX) Section 404(a). Our internal control over financial reporting is a process designed to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements in accordance with generally accepted accounting principles in the United States. Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. Therefore, even those systems determined to be effective can provide only reasonable assurance of achieving their control objectives. Management evaluated the effectiveness of our internal control over financial reporting based on the framework in "Internal Control — Integrated Framework" issued by the Committee of Sponsoring Organizations of the Treadway Commission (COSO).

Our management evaluated, with the participation of our Chief Executive Officer and Chief Financial Officer, the effectiveness of our internal control over financial reporting as of December 31, 2012. Based on that evaluation, our management concluded that our internal control over financial reporting was effective at December 31, 2012. As allowed, this evaluation excludes the operations of Bios due to the recency of the acquisition. Revenues related to the Bios Acquisition were approximately 12% of total revenues for the nine months ended December 31, 2012.

Changes in Internal Control Over Financial Reporting

During the three months ended December 31, 2012, we implemented numerous computer-based controls. These additional computer-based controls have strengthened our internal controls and positioned us to more easily maintain effective internal controls as we implement our growth strategy.

Part II. Other Information

Item 1A. Risk factors

We are affected by risks specific to us as well as factors that affect all businesses operating in a global market. The significant factors known to us that could materially adversely affect our business, financial condition or operating results are described in our annual report on Form 10-K for the year ended March 31, 2012, under the heading "Part I — Item 1A. Risk Factors." There have been no material changes to those risk factors other than the following:

Global economic conditions could materially adversely affect the Company

Our operations and performance depend on worldwide economic conditions. Uncertainty about global economic conditions poses a risk as businesses postpone spending in response to tighter credit, unemployment, negative financial news and/or declines in income or asset values. Other factors that could influence demand include increases in fuel and other energy costs, conditions in the real estate, labor and healthcare costs, access to credit and other macroeconomic factors affecting spending behavior. These worldwide and regional economic conditions could have a material adverse effect on demand for our products and services as our customers reduce or delay capital equipment and other types of purchases.

In the event of financial turmoil affecting the banking system and financial markets, additional consolidation of the financial services industry, or significant financial service institution failures, there could be a new or incremental tightening in the credit markets, low liquidity, and extreme volatility in fixed income, credit, currency, and equity markets. This could have a number of effects on our business, including the insolvency or financial instability of our customers.

Our business may be subject to sales tax

The application of indirect taxes, such as sales tax, is a complex and evolving issue. A company must collect and remit state sales tax from its customers, if the company has “nexus” in that particular state. The determination of nexus varies by state and often requires knowledge of each state’s sales tax case law. The application of existing, new or future laws could change the states in which we have nexus, which could have an adverse effect on our business.

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

There were no repurchases of our common stock, including settlement of loans to employees for the exercise of stock options, for the three months ended December 31, 2012.

On November 7, 2005, the Board of Directors of Mesa Laboratories, Inc. adopted a share repurchase plan which allows for the repurchase of up to 300,000 of our common shares. This plan will continue until the maximum is reached or the plan is terminated by further action of the Board. As of December 31, 2012, we had purchased 156,412 shares, leaving 143,588 remaining under the plan.

Item 6. *Exhibits*

a) Exhibits:

- 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 Chief Executive Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002
- 32.2 Certification of Chief Financial Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002
- 101 The following financial information from the quarterly report on Form 10-Q of Mesa Laboratories, Inc. for the quarter ended December 31, 2012, formatted in XBRL (eXtensible Business Reporting Language): (i) Condensed Statements of Income, (ii) Condensed Balance Sheets, (iii) Condensed Statements of Cash Flows, and (iv) Notes to the Condensed Financial Statements.

b) Reports on Form 8-K:

On December 14, 2012, the Registrant filed a Report on Form 8-K, under Item 5.02, reporting that the Registrant entered into employment agreements with certain officers.

On November 30, 2012, the Registrant filed a Report on Form 8-K, under Item 5.02, reporting the departure of the Registrant’s Chief Financial Officer and the appointment of a new Chief Financial Officer.

On November 8, 2012, the Registrant filed a Report on Form 8-K, under Item 2.02, reporting the issuance of a press release reporting revenues and earnings for the quarter ended September 30, 2012.

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.

MESA LABORATORIES, INC.
(Registrant)

DATED: February 8, 2013

BY: /s/ John J. Sullivan, Ph.D.
John J. Sullivan, Ph.D.
Chief Executive Officer

DATED: February 8, 2013

BY: /s/ John V. Sakys
John V. Sakys
Chief Financial Officer

CERTIFICATIONS PURSUANT TO RULE 13a-14(a)

I, John J. Sullivan, Ph.D., certify that:

1. I have reviewed this quarterly report on Form 10-Q of Mesa Laboratories, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal 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(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - (a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - (b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: February 8, 2013

/s/ John J. Sullivan, Ph.D
John J. Sullivan, Ph.D.
Chief Executive Officer

CERTIFICATIONS PURSUANT TO RULE 13a-14(a)

I, John V. Sakys, certify that:

1. I have reviewed this quarterly report on Form 10-Q of Mesa Laboratories, Inc.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal 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(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - (a) All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - (b) Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: February 8, 2013

/s/ John V. Sakys
John V. Sakys
Chief Financial Officer

CERTIFICATIONS PURSUANT TO RULE 13a-14(b) AND 18 U.S.C SECTION 1350

In connection with the Quarterly Report of Mesa Laboratories, Inc. (the "Company") on Form 10-Q for the fiscal quarter ended December 31, 2012, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, John J. Sullivan, Ph.D., Chief Executive Officer of the Company, certify, pursuant to Rule 13a-14(b) and 18 U.S.C. § 1350, that:

- (1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
- (2) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: February 8, 2013

/s/ John J. Sullivan, Ph.D.

John J. Sullivan, Ph.D.

Chief Executive Officer

CERTIFICATIONS PURSUANT TO RULE 13a-14(b) AND 18 U.S.C SECTION 1350

In connection with the Quarterly Report of Mesa Laboratories, Inc. (the "Company") on Form 10-Q for the fiscal quarter ended December 31, 2012, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, John V. Sakys, Chief Financial Officer of the Company, certify, pursuant to Rule 13a-14(b) and 18 U.S.C. § 1350, that:

- (1) The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
- (2) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: February 8, 2013

/s/John V. Sakys

John V. Sakys
Chief Financial Officer

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