

BIOMERICA INC
Form 10-Q
October 15, 2018

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 AUGUST 31, 2018

OR

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

Commission File Number: 0-8765

BIOMERICA, INC.

(Exact name of registrant as specified in its charter)

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Delaware

95-2645573

(State or other jurisdiction of
incorporation or organization)

(I.R.S. Employer
Identification No.)

17571 Von Karman Avenue, Irvine, CA

92614

(Address of principal executive offices)

(Zip Code)

Registrant's telephone number including area code: (949) 645-2111

(Former name, former address and former fiscal year, if changed since last report.)

(TITLE OF EACH CLASS)

(NAME OF EACH EXCHANGE ON WHICH REGISTERED)

Common, par value \$.08

NASDAQ Capital Market

Securities registered pursuant to Section 12(g) of the Act:

(TITLE OF EACH CLASS)

COMMON STOCK, PAR VALUE \$0.08

Indicate by check whether the registrant (1) 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 Website, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (paragraph 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, 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.

Large Accelerated Filer

Accelerated Filer

Non-Accelerated Filer

Smaller Reporting Company

Indicate by check mark whether the Registrant is a shell company (as defined in Rule 12b-2 of the Act).

Yes No

Indicate the number of shares outstanding of each of the registrant's common stock, as of the latest practicable date 9,033,768 shares of common stock, par value \$0.08, as of October 15, 2018.

BIOMERICA, INC.
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PART I - FINANCIAL INFORMATION
SUMMARIZED FINANCIAL INFORMATION
ITEM 1. FINANCIAL STATEMENTS

BIOMERICA, INC. AND SUBSIDIARIES

CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS

AND COMPREHENSIVE LOSS(UNAUDITED)

Three Months Ended

		August 31,		
	2018		2017	
Net sales	\$ 1,272,870		\$ 1,444,483	
Cost of sales	(935,647)		(929,912)	
Gross profit	337,223		514,571	
Operating Expenses:				
Selling, general and administrative	400,228		452,014	
Research and development	391,781		288,585	
Total operating expenses	792,009		740,599	
Loss from operations	(454,786)		(226,028)	
Other Income (expense):				
Dividend and interest income	3,093		18,969	
Interest expense	(47)		--	
Total other income	3,046		18,969	
Loss before income tax	(451,740)		(207,059)	
Provision for income taxes	--		--	
Net loss	\$ (451,740)		\$ (207,059)	
Basic net loss per common share	\$ (.05)		\$ (0.02)	
Diluted net loss per common share	\$ (.05)		\$ (0.02)	
Weighted average number of common and common equivalent shares:				
Basic	8,930,251		8,511,260	
Diluted	8,930,251		8,511,260	
Net loss	\$ (451,740)		\$ (207,059)	
Other comprehensive gain (loss), net of tax:				
Foreign currency translation	(1,326)		(824)	
Comprehensive loss	\$ (453,066)		\$ (207,883)	

The accompanying notes are an integral part of these statements.

BIOMERICA, INC. AND SUBSIDIARIES

CONDENSED CONSOLIDATED BALANCE SHEETS

August 31, 2018 (unaudited)	May 31, 2018 (audited)
--	-------------------------------------

Current Assets:

Cash and cash equivalents

\$

583,693

\$

1,204,903

Accounts receivable, less allowance for doubtful accounts of
\$63,392 and \$57,695 as of August 31, 2018 and May 31, 2018,
respectively

1,013,687

799,940

Inventories

	2,208,457
	2,178,777
Prepaid expenses and other	
	311,291
	300,409
Total Current Assets	
	4,117,128
	4,484,029
Property and Equipment, net of accumulated depreciation and amortization of \$1,688,423 and \$1,661,128	
	361,935
	351,149

Deferred Tax Assets

10,000

10,000

Investments

165,324

165,324

Intangible Assets, net

104,782

98,923

Other Assets

116,534

113,157

Total Assets

\$

4,875,703

\$

5,222,582

The accompanying notes are an integral part of these statements.

BIOMERICA, INC. AND SUBSIDIARIES

CONDENSED CONSOLIDATED BALANCE SHEETS - Continued

August 31, 2018 (unaudited)	May 31, 2018 (audited)
--------------------------------------	---------------------------------

Current Liabilities:

Accounts payable and accrued expenses

\$

582,976

\$

686,956

Accrued compensation

214,556

209,852

Total Current Liabilities

797,532

896,808

Commitments and Contingencies (Note 6)

Shareholders' Equity:

Preferred stock, no par value authorized 5,000,000 shares,
none issued and none outstanding at August 31, 2018
and May 31, 2018

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Common stock, \$0.08 par value authorized 25,000,000 shares,
issued and outstanding 8,953,832 at August 31, 2018
and 8,888,011 at May 31, 2018

716,305

	711,040
Additional paid-in-capital	
	21,034,686
	20,843,550
Subscriptions receivable	
	(9,062)
Accumulated other comprehensive loss	
	(27,462)
	(26,136)
Accumulated deficit	
	(17,645,358)
	(17,193,618)

Total Shareholders' Equity

4,078,171

4,325,774

Total Liabilities and Shareholders' Equity

\$

4,875,703

\$

5,222,582

The accompanying notes are an integral part of these statements.

BIOMERICA, INC. AND SUBSIDIARIES
CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS

(UNAUDITED)

	Three Months Ended	
	2018	August 31, 2017
Cash flows from operating activities:		
Net loss	\$ (451,740)	\$ (207,059)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation and amortization	43,960	48,833
Change in provision for losses on accounts receivable	5,697	7,514
Inventory reserve	715	(8)
Stock option expense	8,084	4,383
Increase in deferred rent liability	3,333	5,563
Changes in assets and liabilities:		
Accounts receivable	(219,444)	5,202
Inventories	(30,395)	(62,379)
Prepaid expenses and other assets	(14,259)	(19,053)
Accounts payable and accrued expenses	(107,313)	82,480
Accrued compensation	4,704	9,810
Net cash used in operating activities	(756,658)	(124,714)
Cash flows from investing activities:		
Increase in intangibles	(22,524)	--
Purchases of property and equipment	(38,081)	(19,974)
Net cash used in investing activities	(60,605)	(19,974)
Cash flows from financing activities:		
Proceeds from sale of common stock, net	179,504	--
Exercise of stock options	17,875	1,038
Net cash provided by financing activities	197,379	1,038
Effect of exchange rate changes in cash	(1,326)	(824)
Net decrease in cash and cash equivalents	(621,210)	(144,474)
Cash and cash equivalents at beginning of period	1,204,903	1,225,462
Cash and cash equivalents at end of period	\$ 583,693	\$ 1,080,988

Supplemental Disclosure of Cash-Flow Information:

Cash paid during the period for:

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Interest

\$ 47 \$

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The accompanying notes are an integral part of these statements.

BIOMERICA, INC. AND SUBSIDIARIES

NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (UNAUDITED)

Note 1: Basis of Presentation

Biomerica, Inc. and Subsidiaries (collectively "the Company") are primarily engaged in the development, manufacture and marketing of medical diagnostic kits.

The Company develops, manufactures, and markets medical diagnostic products designed for the early detection and monitoring of chronic diseases and medical conditions. The Company's medical diagnostic products are sold worldwide in two markets: 1) clinical laboratories and 2) point of care (physicians' offices and over-the-counter drugstores). The diagnostic test kits are used to analyze blood, urine or fecal samples from patients in the diagnosis of various diseases and other medical complications, or to measure the level of specific hormones, antibodies, antigens or other substances, which may exist in the human body in extremely small concentrations.

The information set forth in these condensed consolidated statements is unaudited and reflects all adjustments which, in the opinion of management, are necessary to present a fair statement of the consolidated results of operations of Biomerica, Inc. and subsidiaries (collectively the Company), for the periods indicated. It does not include all information and footnotes necessary for a fair presentation of financial position, results of operations, and cash flows in conformity with accounting principles generally accepted in the United States of America. All adjustments that were made are of a normal recurring nature.

The unaudited Condensed Consolidated Financial Statements and notes are presented as permitted by the requirements for Form 10-Q and do not contain certain information included in our annual financial statements and notes. The condensed consolidated balance sheet data as of May 31, 2018 was derived from audited financial statements. The accompanying interim condensed consolidated financial statements should be read in conjunction with the financial statements and related notes included in our Annual Report on Form 10-K filed with the Securities and Exchange Commission (SEC) on August 29, 2018 for the fiscal year ended May 31, 2018. The results of operations for our interim periods are not necessarily indicative of results to be achieved for our full fiscal year.

Note 2: Significant Accounting Policies

Principles of Consolidation

The condensed consolidated financial statements include the accounts of Biomerica, Inc. as well as its German subsidiary (BioEurope GmbH) and Mexican subsidiary (Biomerica de Mexico). All significant intercompany accounts and transactions have been eliminated in consolidation.

Accounting Estimates

The preparation of financial statements in conformity with accounting principles generally accepted in the United States of America (GAAP) requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements, and the reported amounts of revenues and expenses during the reported period. Actual results could materially differ from those estimates.

Concentration of Credit Risk

The Company maintains cash balances at certain financial institutions in excess of amounts insured by federal agencies. The Company does not believe it is exposed to significant credit risks.

The Company provides credit in the normal course of business to customers throughout the United States and foreign markets. At August 31, 2018 and May 31, 2018, the Company had one customer which accounted for 53.8% and 53.3%, respectively, of gross accounts receivable. The Company had one customer which accounted for approximately 45.5% and 41.8%, of consolidated sales for the three months ended August 31, 2018 and August 31, 2017, respectively.

For the quarters ended August 31, 2018 and 2017, one company accounted for approximately 21.3% and 20.6% of the purchases or raw materials, respectively.

Cash and Cash Equivalents

Cash and cash equivalents consist of demand deposits and money market accounts with original maturities of less than three months.

Accounts Receivable

The Company extends unsecured credit to its customers on a regular basis. International accounts are required to prepay until they establish a history with the Company and at that time, they are extended credit at levels based on a number of criteria. Credit levels are approved by designated upper level management. Domestic customers are extended initial \$500 credit limits until they establish a history with the Company or submit credit information. All increases in credit limits are also approved by designated upper level management. Management evaluates receivables on a quarterly basis and adjusts the allowance for doubtful accounts accordingly. Balances over ninety days old are usually reserved for.

Occasionally certain long-standing customers, who routinely place large orders, may have unusually large receivables balances relative to the total gross receivables. Management monitors the payments for these large balances closely and very often requires payment of existing invoices before shipping new sales orders.

Inventories

The Company values inventory at the lower of cost (determined using a combination of specific lot identification and the first-in, first-out methods) or net realizable value. Management periodically reviews inventory for excess quantities and obsolescence. Management evaluates quantities on hand, physical condition, and technical functionality as these characteristics may be impacted by anticipated customer demand for current products and new product introductions. The reserve is adjusted based on such evaluation, with a corresponding provision included in cost of sales. Abnormal amounts of idle facility expenses, freight, handling costs and wasted material are recognized as current period charges and the allocation of fixed production overhead is based on the normal capacity of the Company's production facilities.

Inventories approximate the following at:

August 31,	May 31,
2018	2018

Raw materials	
\$	
	989,000
\$	
	1,000,000
Work in progress	
	923,000
	854,000
Finished products	
	296,000
	325,000
Total	
\$	
	2,208,000
\$	
	2,179,000

Reserves for inventory obsolescence are recorded as necessary to reduce obsolete inventory to estimated realizable value or to specifically reserve for obsolete inventory that the Company intends to dispose of. As of August 31, 2018 and May 31, 2018, inventory reserves were approximately \$52,000.

Property and Equipment, net

Property and equipment are stated at cost. Expenditures for additions and major improvements are capitalized. Repairs and maintenance costs are charged to operations as incurred. When property and equipment are retired or otherwise disposed of, the related cost and accumulated depreciation or amortization is removed from the accounts, and gains or losses from retirements and dispositions are credited or charged to income.

Depreciation and amortization are provided over the estimated useful lives of the related assets, ranging from 5 to 10 years, using the straight-line method. Leasehold improvements are amortized over the lesser of the estimated useful life of the asset or the term of the lease. Depreciation and amortization expense on property and equipment and leasehold improvements amounted to \$27,295 and \$31,222 for the three months ended August 31, 2018 and 2017, respectively.

Intangible Assets, net

Intangible assets include trademarks, product rights, licenses, technology rights and patents, and are accounted for based on Accounting Standards Codification (ASC) 350, Intangibles Goodwill and Other (ASC 350). In that regard, intangible assets that have indefinite useful lives are not amortized but are tested at least annually for impairment or more frequently if events or changes in circumstances indicate that the asset might be impaired. Intangible assets are being amortized using the straight-line method over the useful life; not to exceed 18 years for marketing and distribution rights, 10 years for purchased technology use rights, licenses, and 17 years for patents. Amortization amounted to \$16,665 and \$17,611 for the three months ended August 31, 2018 and 2017, respectively.

Share-Based Compensation

The Company follows the guidance of the accounting provisions of ASC 718, Share-based Compensation (ASC 718), which requires the use of the fair-value based method to determine compensation for all arrangements under which employees and others receive shares of stock or equity instruments (options). The fair value of each option award is estimated on the date of grant using the Black-Scholes option pricing model that uses assumptions for expected volatility, expected dividends, expected forfeiture rate, expected term, and the risk-free interest rate.

Expected volatilities are based on weighted averages of the historical volatility of the Company s stock and other factors estimated over the expected term of the options. The expected forfeiture rate is based on historical forfeitures experienced. The expected term of options granted is derived using the simplified method which computes expected term as the average of the sum of the vesting term plus the contract term as historically the Company had limited activity surrounding its options. The risk-free rate is based on the U.S. Treasury yield curve in effect at the time of grant for the period of the expected term.

The following summary presents the options and warrants granted, exercised, expired, cancelled and outstanding as of August 31, 2018:

	Option	Exercise
	Shares	Price
		Weighted
		Average
Outstanding May 31, 2018	1,138,625	\$ 1.65
Granted	--	--
Forfeited	(3,500)	0.99
Exercised	(21,500)	0.83

Outstanding August 31, 2018	1,113,625	\$	1.67
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In the quarter ended August 31, 2018, options to purchase 21,500 shares of common stock were exercised at a prices ranging from \$0.83 to \$0.85. Net proceeds to the Company were \$17,875.

Revenue Recognition

Revenues from product sales are recognized at the time the product is shipped, customarily FOB shipping point, at which point title passes. An allowance is established when necessary for estimated returns as revenue is recognized. As of August 31, 2018 and May 31, 2018, the allowance for returns was \$0. In conjunction with sales to certain customers, the Company provides free products upon attaining certain levels of purchases by the customer. The Company accounts for these free products in accordance with ASC 605-50, Revenue Recognition - Customer Payments and Incentives and recognizes the cost of the product as part of cost of sales.

Investments

From time-to-time, the Company makes investments in privately-held companies. The Company determines whether the fair values of any investments in privately-held entities have declined below their carrying value whenever adverse events or changes in circumstances indicate that recorded values may not be recoverable. If the Company considers any such decline to be other than temporary (based on various factors, including historical financial results, and the overall health of the investee's industry), a write-down to estimated fair value is recorded. The Company currently has not written down the investment and no events have occurred which could indicate the carrying value to be less than the fair value. Investments represent the Company's investment in a Polish distributor which is primarily engaged in distributing medical devices. The Company owns approximately 6% of the investee, and accordingly, applies the cost method to account for the investment. Under the cost method, investments are recorded at cost, with gains and losses recognized as of the sale date, and income recorded when received.

Shipping and Handling Fees and Costs

Shipping and handling fees billed to customers are classified as net sales and shipping and handling costs are classified as cost of sales. The Company included shipping and handling fees billed to customers in net sales. The Company included shipping and handling costs associated with inbound freight and unreimbursed shipping to customers in cost of sales.

Research and Development

Research and development costs are expensed as incurred.

Income Taxes

The Company has provided a valuation allowance on deferred tax assets of approximately \$1,640,000 and \$1,549,000 as of August 31, 2018 and May 31, 2018, respectively.

Foreign Currency Translation

The subsidiaries located in Germany and Mexico are accounted for primarily using local functional currency. Accordingly, assets and liabilities of these subsidiaries are translated using exchange rates in effect at the end of the period, and revenues and costs are translated using average exchange rates for the period. The subsidiaries in Germany and Mexico each have one bank account which according to exchange rates in effect at the end of each period need to be adjusted for that fluctuation. The resulting adjustments are presented as a separate component of accumulated other comprehensive loss.

Deferred Rent

Incentive payments received from landlords are recorded as deferred lease incentives and are amortized over the underlying lease term on a straight-line basis as a reduction of rent expense. When the terms of an operating lease provide for periods of free rent, rent concessions, and/or rent escalations, the Company establishes a deferred rent liability for the difference between the scheduled rent payment and the straight-line rent expense recognized. This

deferred rent liability is amortized over the underlying lease term on a straight-line basis as a reduction of rent expense.

Net Loss Per Share

Basic loss per share is computed as net loss divided by the weighted average number of common shares outstanding for the period. Diluted loss per share reflects the potential dilution that could occur from common shares issuable through stock options using the treasury stock method. The total amount of anti-dilutive options not included in the earnings per share calculation for the three months ended August 31, 2018 and 2017 was 1,113,625 and 945,000 respectively.

The following table illustrates the required disclosure of the reconciliation of the numerators and denominators of the basic and diluted earnings per share computations:

	2018	Three Months Ended
	2017	August 31,
Numerator		
Loss from continuing operations		
\$		(451,740)
\$		(207,059)
Denominator for basic net loss per common share		8,930,251

	8,511,260
Effect of dilutive securities:	
Options	--
	--
Denominator for diluted net loss per common share	
	8,930,251
	8,511,260
Basic net loss per common share	
\$	(0.05)
	(0.02)
Diluted net loss per common share	
\$	(0.05)

\$

(0.02)

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Recent Accounting Pronouncements

In August 2014, the Financial Accounting Standards Board (FASB) issued Accounting Standards Update (ASU) 2014-15, Presentation of Financial Statements – Going Concern (Subtopic 205-40), which addresses Disclosure of Uncertainties about an Entity’s Ability to Continue as a Going Concern . In connection with preparing financial statements for each annual and interim reporting period, an entity’s management should evaluate whether there are conditions or events, considered in the aggregate, that raise substantial doubt about the entity’s ability to continue as a going concern within one year after the date that the financial statements are issued (or within one year after the date that the financial statements are available to be issued when applicable). Management’s evaluation should be based on relevant conditions and events that are known and reasonably knowable at the date that the financial statements are issued (or at the date that the financial statements are available to be issued when applicable). The amendments in this update are effective for the annual period ending after December 15, 2016, and for annual periods and interim periods thereafter. Early adoption is permitted. Management adopted the provisions of this statement and is taking them into account in the preparation of the financial statements for the period ended August 31, 2018.

In May 2014, the FASB issued ASU 2014-09, Revenue from Contracts with Customers (ASU 2014-09). ASU 2014-09 is a comprehensive new revenue recognition model requiring a company to recognize revenue to depict the transfer of goods or services to a customer at an amount reflecting the consideration it expects to receive in exchange for those goods or services. In adopting, ASU 2014-09, companies may use either a full retrospective or a modified retrospective approach. ASU 2014-09 is effective for the first interim period within annual reporting periods beginning December 15, 2016, and early adoption is not permitted. During August 2015, the FASB voted to defer the effective date of the above mentioned revenue recognition guidance by one year to December 15, 2017 for interim and annual reporting periods beginning after that date and permitted early adoption of the standard, but not before the original effective date of December 15, 2016. Management adopted the provisions of this statement and is taking them into account in the preparation of the financial statements for the period ended August 31, 2018.

On January 5, 2016, the FASB issued ASU 2016-01, Financial Instruments-Overall (Subtopic 825-10): Recognition and Measurement of Financial Assets and Financial Liabilities (ASU-2016-01). The release affects public and private companies that hold financial assets or owe financial liabilities. ASU-2016-01 will take effect for public companies for fiscal years beginning after December 15, 2017. Management adopted the provisions of this statement and is taking them into account in the preparation of the financial statements for the period ended August 31, 2018.

On February 25, 2016, the FASB issued ASU 2016-02, Leases (Topic 842) (ASU-2016-02). ASU-2016-02 defines whether a contract is a lease. If it is a lease, the Company is required to recognize the lease assets and liabilities. ASU-2016-02 is effective for public companies for the annual periods beginning after December 15, 2018. Management is evaluating the provisions of this statement and has not determined what impact the adoption of ASU-2016-02 will have on the Company’s financial position or results of operations.

On August 26, 2016, the FASB issued ASU 2016-15, Statement of Cash Flows (Topic 230): Classification of Certain Cash Receipts and Cash Payments. This Update addresses eight specific cash flow issues with the objective of reducing the existing diversity in practice. ASU-2016-15 will take effect for public companies for the fiscal years beginning after December 15, 2017, and interim periods within those fiscal years. Management adopted the provisions of this statement and is taking them into account in the preparation of the financial statements for the period ended August 31, 2018.

On November 27, 2016, the FASB issued ASU 2016-18, Statement of Cash Flows (Topic 230): Restricted Cash. This update addresses the fact that diversity exists in the classification and presentation of changes in restricted cash on the statement of cash flows under Topic 230, Statement of Cash Flows. ASU-2016-18 will take effect for public companies for the fiscal years beginning after December 15, 2017, and interim periods within those fiscal years. Management adopted the provisions of this statement and is taking them into account in the preparation of the financial statements for the period ended August 31, 2018.

In January 2017 the FASB issued ASU 2017-04, Intangibles – Goodwill and Other (Topic 350), Simplifying the test for Goodwill Impairment. This update addresses how an entity should perform its annual, or interim, goodwill impairment test by comparing the fair value of a reporting unit with its carrying amount. ASU 2017-04 will take effect for public companies for the fiscal years beginning after December 15, 2019. Management is evaluating the provisions of this statement and has not determined what impact the adoption of ASU-2017-04 will have on the Company's financial position or results of operations.

On February 15, 2018, the FASB issued ASU 2018-02, Reclassification of Certain Tax Effects From Accumulated Comprehensive Income (ASU 2018-02). ASU 2018-02 will give companies the option to reclassify stranded tax effects caused by the newly-enacted U.S. TAX Cuts and Jobs Act (TCJA) from accumulated other comprehensive income (ASCI) to retained earnings. ASU 2018-02 will take effect for all companies for the fiscal years beginning after December 15, 2018, and interim periods within those fiscal years. Management is evaluating the provisions of this statement and has not determined what impact the adoption of ASU 2018-02 will have on the Company's financial position or results of operations.

On June 20, 2018, the FASB issued ASU 2018-07, Compensation – Stock Compensation (Topic 718): Improvements to Nonemployee Share-Based Payment Accounting. ASU 2018-07 is intended to reduce cost and complexity and to improve financial reporting for share-based payments to nonemployees (for example, service providers, external legal counsel, suppliers, etc.). ASU 2018-07 will be effective for public companies for December 31, 2019 financial statements and for nonpublic entities for December 31, 2020 financial statements. Early adoption is permitted, but no earlier than entity’s adoption date for ASC Topic 606, Revenue from Contracts with Customers. Management is evaluating the provisions of this statement and has not determined what impact the adoption of ASU-2018-7 will have on the Company’s financial position or results of operations.

Other recent ASU's issued by the FASB and guidance issued by the Securities and Exchange Commission did not, or are not believed by management to, have a material effect on the Company’s present or future consolidated financial statements.

Note 3: Accounts Payable and Accrued Expenses

The Company’s accounts payable and accrued expense balances consist of the following at:

	August 31,		May 31,
	2018		2018
Accounts payable	\$ 548,283	\$	655,599
Deferred rent	34,690		31,357
Total	\$ 582,973	\$	686,956

Note 4: Shareholders’ Equity

As described in the Company’s Form 10Q, filed with the Securities and Exchange Commission on April 15, 2018, the Form 10K report, filed with the Securities and Exchange Commission on August 29, 2018, and the Form S-3 Registration Statement and Prospectus filed on June 30, 2017 and December 4, 2017, respectively, the Company entered into an At Market Issuance Sales Agreement, whereby, the Company may raise additional working capital and funds for continued development of current research projects. These funds will be needed to fund current research and development projects and bring them to the next state of completion. Management expects to raise additional funds throughout the year from the At Market Issuance Agreement to fund operations as necessary. During the quarter that ended August 31, 2018, the Company received \$179,504 in net proceeds from the sale of its common stock through this Agreement.

Note 5: Geographic Information

Financial information about foreign and domestic operations and export sales is approximately as follows:

	Three Months Ended	
	2018	August 31, 2017
Revenues from sales to unaffiliated customers:		
United States	\$ 143,000	\$ 187,000
Asia	648,000	620,000
Europe	392,000	532,000
South America	71,000	75,000
Middle East	19,000	26,000
Other	--	4,000
	\$ 1,273,000	\$ 1,444,000

No other geographic concentrations exist where net sales exceed 10% of total net sales.

As of August 31, 2018 and May 31, 2018, approximately \$676,000 and \$657,000 of Biomerica's gross inventory and approximately \$43,000 and \$41,000, of Biomerica's property and equipment, net of accumulated depreciation and amortization, was located in Mexicali, Mexico, respectively.

Note 6: Commitments and Contingencies

On June 18, 2009, the Company entered into an agreement to lease a building in Irvine, California. The lease commenced September 1, 2009 and ended August 31, 2016. The initial base rent was set at \$18,490 per month with scheduled annual increases through the end of the lease term. The rent was \$22,080. In November 2015, the Company signed the First Amendment to Lease to extend the lease until August 31, 2021. The initial base rent for the lease amendment which started September 1, 2016 is \$21,000 per month. September 1, 2018 the rent increased to \$22,279 per month.

In November 2016, the Company's Mexican subsidiary, Biomerica de Mexico, entered into a 10-year lease for approximately 8,100 square feet of manufacturing space with initial base rent of \$2,926 per month. The Company has a one 10-year option to renew at the end of the initial lease period. Biomerica, Inc. is not a guarantor of such lease. Biomerica de Mexico also leases a smaller unit on a month-to-month basis for use in one manufacturing process. In addition, the Company leases a small office in Lindau, Germany, as headquarters for BioEurope GmbH, its Germany subsidiary.

On December 1, 2017, Biomerica, Inc. (the Company) entered into an At Market Issuance Sales Agreement (the At Market Issuance Sales Agreement) with an agent (Agent), pursuant to which the Company may offer and sell from time to time up to an aggregate of \$7,000,000 of shares of the Company's common stock, par value \$0.08 per share (the Placement Shares), through the Agent.

The Placement Shares have been registered under the Securities Act of 1933, as amended (the Securities Act), pursuant to the Registration Statement on Form S-3 (File No. 333-219130) (the Registration Statement), which was originally filed with the Securities and Exchange Commission (SEC) on June 30, 2017 and declared effective by the SEC on July 20, 2017, the base prospectus contained within the Registration Statement, and a prospectus supplement that was filed with the SEC on December 1, 2017.

Sales of the Placement Shares, if any, pursuant to the At Market Issuance Sales Agreement, may be made in sales deemed to be at the market offerings as defined in Rule 415 promulgated under the Securities Act. The Agent will act as sales agent and will use commercially reasonable efforts to sell on the Company's behalf all of the Placement Shares requested to be sold by the Company, consistent with its normal trading and sales practices, on mutually agreed terms between the Agent and the Company.

The Company has no obligation to sell any of the Placement Shares under the At Market Issuance Sales Agreement, and may at any time suspend offers under the At Market Issuance Sales Agreement or terminate the At Market Issuance Sales Agreement. The Company intends to use the net proceeds from this offering for general corporate purposes, including, without limitation, sales and marketing activities, clinical studies and product development, making acquisitions of assets, businesses, companies or securities, capital expenditures, and for working capital needs.

During the quarter ended August 31, 2018 the Company sold 44,321 shares of common stock under the S-3 registration statement and received net proceeds of \$170,443.

Note 7: Subsequent Events

In September 2018, the Company granted 139,000 options to purchase common stock to various employees at the exercise price of \$3.62 per share under the 2017 Stock Option and Restricted Stock Option plan. The shares are exercisable one quarter one year from date of grant and one quarter per year thereafter and expire five years from date of grant.

Subsequent to August 31, 2018, the Company sold 52,936 shares of its common stock under its At Market Issuance Sales Agreement and S-3 Registration Statement for total net proceeds of approximately \$180,000.

Subsequent to August 31, 2018, 27,000 options to purchase shares of the Company's common stock were exercised at prices ranging from \$0.71 to \$1.04. Gross proceeds to the Company were \$20,500.

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

EXCEPT FOR HISTORICAL INFORMATION CONTAINED HEREIN, THE STATEMENTS IN THIS FORM 10-K MAY BE FORWARD-LOOKING STATEMENTS WITHIN THE MEANING OF SECTION 21E OF THE SECURITIES EXCHANGE ACT OF 1934 AND SECTION 27A OF THE SECURITIES ACT OF 1933. FORWARD-LOOKING STATEMENTS INVOLVE KNOWN AND UNKNOWN RISKS AND UNCERTAINTIES WHICH MAY CAUSE BIOMERICA'S RESULTS IN FUTURE PERIODS TO DIFFER MATERIALLY FROM FORECASTED RESULTS. THESE RISKS AND UNCERTAINTIES INCLUDE, AMONG OTHER THINGS, THE CONTINUED DEMAND FOR THE COMPANY'S PRODUCTS, AVAILABILITY OF RAW MATERIALS, THE STATE OF THE ECONOMY, RESULTS OF RESEARCH AND DEVELOPMENT ACTIVITIES AND THE CONTINUED ABILITY OF THE COMPANY TO MAINTAIN THE LICENSES AND APPROVALS REQUIRED. THESE AND OTHER RISKS ARE DESCRIBED IN THE COMPANY'S ANNUAL REPORT ON FORM 10-K AND IN THE COMPANY'S OTHER FILINGS WITH THE SECURITIES AND EXCHANGE COMMISSION.

EXCEPT AS MAY BE REQUIRED BY APPLICABLE LAW, WE MAY NOT UPDATE OR REVISE OUR FORWARD-LOOKING STATEMENTS AND THE LACK OF SUCH UPDATE DOES NOT IMPLY THAT ACTUAL EVENTS ARE AS ORIGINALLY EXPRESSED BY SUCH FORWARD-LOOKING STATEMENTS. YOU SHOULD READ THE DISCLOSURES IN THIS REPORT AND OTHER REPORTS WHICH WE FILE WITH THE SECURITIES AND EXCHANGE COMMISSION.

OVERVIEW

Biomerica, Inc. and Subsidiaries ("Biomerica", the "Company", "we" or "our") develops, manufactures, and markets medical diagnostic products designed for the early detection and monitoring of chronic diseases and medical conditions. Our medical diagnostic products are sold worldwide in two markets: 1) clinical laboratories and 2) point of care (physicians' offices and over-the-counter drugstores). Our diagnostic test kits are used to analyze blood or urine from patients in the diagnosis of various diseases and other medical complications, or to measure the level of specific hormones, antibodies, antigens or other substances, which may exist in the human body in extremely small concentrations.

RESULTS OF OPERATIONS

Consolidated net sales for Biomerica were \$1,272,870 for the three months ended August 31, 2018 as compared to \$1,444,483 for the same period in the previous year. This represents a decrease of \$171,613, or 11.9%. The decrease was primarily due to decreased sales of approximately \$140,000 in Europe which was due to lower sales to certain distributors and timing of certain orders. This was partially offset by an increase in Asia of approximately \$28,000.

Sales in the U.S. were down by approximately \$44,000 from the prior year due to higher sales in the prior year of contract manufacturing.

For the three months ended August 31, 2018 as compared to August 31, 2017, cost of sales increased as a percentage of sales from 64.4% of sales, or \$929,912, to 73.5% of sales, or \$935,647. Cost of sales as a percentage of sales increased primarily due to the lower sales volume in relationship to fixed expenses as well as higher material costs.

For the three months ended August 31, 2018 compared to 2017, selling, general and administrative costs were \$400,228 as compared to \$452,014, a decrease of \$51,786, or 11.5%. This was due to lower consulting and outside services in fiscal 2019.

For the three months ended August 31, 2018 and 2017, research and development expenses were \$391,781 as compared to \$288,585, an increase of \$103,196, or 35.1%. The increase was due to higher costs (materials, legal, regulatory and wages) incurred in the development, regulatory approval and applications for patents of new products in the gastroenterology area including costs involved with the clinical studies which will continue in future quarters. During August 2018, the Company received its first patent related to this project. The patent was issued in Korea and covers the InFoods IBS product.

For the three months ended August 31, 2018 as compared to August 31, 2017, dividend and interest income decreased from \$18,969 to \$3,093.

LIQUIDITY AND CAPITAL RESOURCES

As of August 31, 2018 and May 31, 2018, the Company had cash and cash equivalents in the amount of \$583,693 and \$1,204,903 and working capital of \$3,319,596 and \$3,587,221, respectively.

During the three months ended August 31, 2018, the Company's operations used cash of \$756,658 compared to cash used in operations of \$124,714 in the same period of the prior fiscal year. Cash used by operations in fiscal 2018 was primarily a result of an increase in accounts receivable of \$219,444 and a decrease in accounts payable and accrued expenses of \$107,313 and a net loss of \$451,070 as compared to fiscal 2017 which had increased accounts payable and accrued expenses of \$82,480 which was offset against an increase in inventories of \$62,379 and a net loss of \$207,059. Cash used in investing activities in fiscal 2018 was \$38,081 for purchases of fixed assets and \$22,524 for increased intangibles compared to fiscal 2017 of \$19,974, which was the result of property and equipment purchases. Cash provided by financing activities in fiscal 2018 was \$197,379 which was a result of \$17,875 for stock option exercises and \$179,504 from the net proceeds from the sale of common stock compared to \$1,038 in fiscal 2017, which was a result of the exercise of a stock option.

The Company has been working on new products for the gastroenterology market. Patent applications for the new products have been filed and the Company has been working on obtaining additional patents and U.S. regulatory approvals. The Company has been spending significant funds on the research, development and related costs and expects this will continue in order to obtain the desired patents and approvals.

As described in the Company's Form 10Q, filed with the Securities and Exchange Commission on April 15, 2018, the Form 10K report, filed with the Securities and Exchange Commission on August 29, 2018, and the Form S-3 Registration Statement and Prospectus filed on June 30, 2017 and December 4, 2017, respectively, the Company entered into an At Market Issuance Sales Agreement, whereby, the Company intends to raise additional working capital and funds for continued development of current research projects. These funds will be needed to fund current research and development projects and bring them to the next state of completion. Management expects to raise additional funds throughout the year from the At Market Issuance Agreement to fund operations as necessary. During the quarter that ended August 31, 2018, the Company received \$179,504 in net proceeds from the sale of its common stock through this Agreement.

Off Balance Sheet Arrangements None.

CRITICAL ACCOUNTING POLICIES

The preparation of condensed consolidated financial statements in conformity with accounting principles generally accepted in the United States of America requires us to make a number of estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements. Such estimates and assumptions affect the reported amounts of revenues and expenses during the reporting period. We base our estimates on historical experience and on various other assumptions that we believe to be reasonable under the circumstances. Actual results may differ materially from these estimates under different assumptions or conditions. We continue to monitor significant estimates made during the preparation of our financial statements. On an ongoing basis, we evaluate estimates and assumptions based upon historical experience and various other factors and circumstances. We believe our estimates and assumptions are reasonable in the circumstances; however, actual results may differ from these estimates under different future conditions.

We believe that the estimates and assumptions that are most important to the portrayal of our financial condition and results of operations, in that they require subjective or complex judgments, form the basis for the accounting policies deemed to be most critical to us. These relate to revenue recognition, bad debts, inventory overhead application, and inventory reserve. We believe estimates and assumptions related to these critical accounting policies are appropriate under the circumstances; however, should future events or occurrences result in unanticipated consequences, there could be a material impact on our future financial conditions or results of operations. We suggest that our significant accounting policies be read in conjunction with this Management's Discussion and Analysis of Financial Condition and Results of Operations. Please refer to Note 2 for information on Significant Accounting Policies.

Item 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK.

We are a smaller reporting company as defined by Rule 12b-2 of the Securities Exchange Act of 1934 and are not required to provide the information under this item.

Item 4. CONTROLS AND PROCEDURES

Our management evaluated the effectiveness of our disclosure controls and procedures as defined in Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934, as amended, or the Exchange Act, as of the end of the period covered by this report. Our management recognizes that any controls and procedures, no matter how well designed and operated, can provide only reasonable assurance of achieving their objectives and management is required to apply its judgment in evaluating the cost-benefit relationship of possible controls and procedures. The disclosure controls and procedures have been designed to provide reasonable assurance of achieving their objectives and the Chief Executive Officer and Chief Financial Officer have concluded that our disclosure controls and procedures are effective at the "reasonable assurance" level. Based upon that evaluation, the Chief Executive Officer and Chief Financial Officer concluded that the disclosure controls and procedures were effective to ensure that information required to be disclosed in the reports that we file and submit under the Exchange Act is (1) recorded, processed, summarized and reported within the time periods specified in the Commission's rules and forms; and (2) accumulated and communicated to the Company's management, including its Chief Executive Officer and Chief Financial Officer, as appropriate, to allow timely decisions regarding required disclosure.

There have been no changes in our internal control over financial reporting identified in connection with the evaluation that occurred during our last fiscal quarter that has materially affected, or that is reasonably likely to materially affect, our internal control over financial reporting.

PART II. OTHER INFORMATION

Item 1. LEGAL PROCEEDINGS. None.

Item 1A. RISKS AND UNCERTAINTIES.

You should read the following factors in conjunction with the factors discussed elsewhere in this and our other filings with the Securities and Exchange Commission and in materials incorporated by reference in these filings. The following is intended to highlight certain factors that may affect the financial condition and results of operations of Biomerica, Inc. and are not meant to be an exhaustive discussion of risks that apply to companies such as Biomerica, Inc. Like other businesses, Biomerica, Inc. is susceptible to macroeconomic downturns in the United States or abroad, as were experienced in recent history that may affect the general economic climate and performance of Biomerica, Inc. or its customers.

Aside from general macroeconomic downturns, the additional material factors that could affect future financial results include, but are not limited to: Terrorist attacks and the impact of such events; diminished access to raw materials that directly enter into our manufacturing process; shipping labor disruption or other major degradation of the ability to ship our products to end users; inability to successfully control our margins which are affected by many factors including competition and product mix; protracted shutdown of the U.S. border due to an escalation of terrorist or counter terrorist activity; any changes in our business relationships with international distributors or the economic climate they operate in; any event that has a material adverse impact on our foreign manufacturing operations may adversely affect our operations as a whole; failure to manage the future expansion of our business could have a material adverse effect on our revenues and profitability; possible costs or difficulty in complying with government regulations and the delays in receiving required regulatory approvals or the enactment of new adverse regulations or regulatory requirements; numerous competitors, some of which have substantially greater financial and other resources than we do; potential claims and litigation brought by patients or medical professionals alleging harm caused by the use of or exposure to our products; quarterly variations in operating results caused by a number of factors, including business and industry conditions; concentrations of sales with certain distributors could adversely affect the results of the Company if the Company were to lose the sales of that distributor and other factors beyond our control; high balances carried on accounts receivables from concentrated customers; and the costs of recalls, should such occasion arise. All these factors make it difficult to predict operating results for any particular period.

Item 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS. None.

Item 3. DEFAULTS UPON SENIOR SECURITIES. None.

Item 4. MINE SAFETY DISCLOSURES. None.

Item 5. OTHER INFORMATION. None.

Item 6. EXHIBITS.

The following exhibits are filed or furnished as part of this quarterly report on Form 10-Q:

Exhibit No.	Description
31.1*	Certification Pursuant to Section 302 of the Sarbanes-Oxley Act Zackary S. Irani
31.2*	Certification Pursuant to Section 302 of the Sarbanes-Oxley Act Janet Moore
32.1*	Certification Pursuant to Section 906 of the Sarbanes-Oxley Act Zackary S. Irani
32.2*	Certification Pursuant to Section 906 of the Sarbanes-Oxley Act Janet Moore

101 Interactive data files pursuant to Rule 405 Regulation S-T, as follows:

101.INS-XBRL Instance Document

101.SCH-XBRL Taxonomy Extension Schema Document

101.CAL-XBRL Taxonomy Extension Calculation Linkbase Document

101.DEF XBRL Taxonomy Extension Definition Linkbase Document

101.LAB-XBRL Taxonomy Extension Label Linkbase Document

101.PRE-XBRL Taxonomy Extension Presentation Linkbase Document

*Filed herewith.

SIGNATURES

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

BIOMERICA, INC.

By:

/S/ Zackary S. Irani

Zackary S. Irani

Chief Executive Officer

(Principal Executive Officer)

Date: October 15, 2018

By:

/S/ Janet Moore

Janet Moore

Chief Financial Officer

(Principal Financial Officer)

Date: October 15, 2018