
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, DC 20549**

FORM 10-Q

Quarterly report pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

For the quarterly period ended March 31, 2010

Transition report pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

For the transition period from _____ to _____

Commission file number 001-33957

HARVARD BIOSCIENCE, INC.

(Exact Name of Registrant as Specified in Its Charter)

Delaware
(State or Other Jurisdiction of
Incorporation or Organization)

84 October Hill Road, Holliston, MA
(Address of Principal Executive Offices)

04-3306140
(IRS Employer
Identification No.)

01746
(Zip Code)

(508) 893-8999
(Registrant's telephone number, including area code)

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

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

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

Large accelerated filer Accelerated filer
Non-accelerated filer (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

APPLICABLE ONLY TO CORPORATE ISSUERS:

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

As of April 30, 2010, there were 29,595,686 shares of Common Stock, par value \$0.01 per share, outstanding.

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Form 10-Q
For the Quarter Ended March 31, 2010

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PART I. FINANCIAL INFORMATION

Item 1. Financial Statements.

HARVARD BIOSCIENCE, INC. AND SUBSIDIARIES
CONSOLIDATED BALANCE SHEETS
(unaudited, in thousands, except share and per share amounts)

	<u>March 31,</u> <u>2010</u>	<u>December 31,</u> <u>2009</u>
<u>Assets</u>		
Current assets:		
Cash and cash equivalents	\$ 17,043	\$ 16,588
Accounts receivable, net of allowance for doubtful accounts of \$385 and \$403, respectively	14,513	14,383
Inventories	14,020	14,406
Deferred income tax assets - current	556	573
Other receivables and other assets	2,821	2,249
Total current assets	<u>48,953</u>	<u>48,199</u>
Property, plant and equipment, net	3,349	3,545
Deferred income tax assets - non-current	313	318
Amortizable intangible assets, net	20,281	21,104
Goodwill	31,362	32,108
Other indefinite lived intangible assets	1,280	1,301
Other assets	447	656
Total assets	<u>\$ 105,985</u>	<u>\$ 107,231</u>
<u>Liabilities and Stockholders' Equity</u>		
Current liabilities:		
Notes payable	\$ 6	\$ 13
Accounts payable	4,939	4,856
Deferred revenue	573	434
Accrued income taxes payable	786	369
Accrued expenses	3,520	3,680
Other liabilities - current	2,323	2,906
Total current liabilities	<u>12,147</u>	<u>12,258</u>
Long-term debt, less current installments	11,807	13,308
Deferred income tax liabilities - non-current	2,055	2,037
Other liabilities - non-current	4,060	4,371
Total liabilities	<u>30,069</u>	<u>31,974</u>
Commitments and contingencies		
Stockholders' equity:		
Preferred stock, par value \$0.01 per share, 5,000,000 shares authorized	—	—
Common stock, par value \$0.01 per share, 80,000,000 shares authorized; 35,948,108 shares issued and 29,584,436 shares outstanding	360	360
Additional paid-in-capital	185,414	184,856
Accumulated deficit	(100,236)	(102,457)
Accumulated other comprehensive income	(3,954)	(1,834)
Treasury stock at cost, 6,363,672 common shares	(5,668)	(5,668)
Total stockholders' equity	<u>75,916</u>	<u>75,257</u>
Total liabilities and stockholders' equity	<u>\$ 105,985</u>	<u>\$ 107,231</u>

See accompanying notes to unaudited consolidated financial statements.

HARVARD BIOSCIENCE, INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF OPERATIONS
(unaudited, in thousands, except per share amounts)

	Three Months Ended	
	March 31,	
	2010	2009
Revenues	\$26,300	\$19,072
Cost of product revenues	13,518	9,662
Gross profit	<u>12,782</u>	<u>9,410</u>
Sales and marketing expenses	3,807	2,372
General and administrative expenses	4,261	3,317
Research and development expenses	1,207	999
Restructuring charges	—	27
Amortization of intangible assets	531	344
Total operating expenses	<u>9,806</u>	<u>7,059</u>
Operating income	<u>2,976</u>	<u>2,351</u>
Other income (expense):		
Foreign exchange	(26)	76
Interest expense	(155)	(45)
Interest income	42	7
Other, net	(15)	53
Other income (expense), net	<u>(154)</u>	<u>91</u>
Income before income taxes	2,822	2,442
Income taxes	601	603
Net income	<u>\$ 2,221</u>	<u>\$ 1,839</u>
Income per share:		
Basic earnings per common share	<u>\$ 0.07</u>	<u>\$ 0.06</u>
Diluted earnings per common share	<u>\$ 0.07</u>	<u>\$ 0.06</u>
Weighted average common shares:		
Basic	<u>29,584</u>	<u>30,012</u>
Diluted	<u>29,941</u>	<u>30,120</u>

See accompanying notes to unaudited consolidated financial statements.

HARVARD BIOSCIENCE, INC. AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF CASH FLOWS
(unaudited, in thousands)

	Three Months Ended	
	March 31,	
	2010	2009
Cash flows from operating activities:		
Net income	\$ 2,221	\$ 1,839
Adjustments to reconcile net income to net cash provided by operating activities:		
Stock compensation expense	558	312
Depreciation	300	282
Restructuring charges	—	28
Amortization of catalog costs	67	71
Provision for allowance for doubtful accounts	(2)	(14)
Amortization of intangible assets	531	344
Amortization of deferred financing costs	22	6
Deferred income taxes	34	(1)
Changes in operating assets and liabilities:		
(Increase) decrease in accounts receivable	(481)	2,699
Increase in inventories	(36)	(302)
Increase in other receivables and other assets	(527)	(304)
Increase (decrease) in trade accounts payable	240	(899)
Increase in accrued income taxes payable	758	194
(Decrease) increase in accrued expenses	(651)	100
Increase (decrease) in deferred revenue	148	(5)
Decrease in other liabilities	(74)	(59)
Net cash provided by operating activities	<u>3,108</u>	<u>4,291</u>
Cash flows used in investing activities:		
Additions to property, plant and equipment	(204)	(299)
Additions to catalog costs	(324)	(11)
Net cash used in investing activities	<u>(528)</u>	<u>(310)</u>
Cash flows used in financing activities:		
Repayments of debt	(1,508)	(376)
Purchases of treasury stock	—	(1,206)
Net cash used in financing activities	<u>(1,508)</u>	<u>(1,582)</u>
Effect of exchange rate changes on cash	<u>(617)</u>	<u>(214)</u>
Increase in cash and cash equivalents	455	2,185
Cash and cash equivalents at the beginning of period	16,588	13,698
Cash and cash equivalents at the end of period	<u>\$17,043</u>	<u>\$15,883</u>
Supplemental disclosures of cash flow information:		
Cash paid for interest	\$ 144	\$ 35
Net cash paid for income taxes	\$ 592	\$ 406

See accompanying notes to unaudited consolidated financial statements.

HARVARD BIOSCIENCE, INC. AND SUBSIDIARIES
Notes to Unaudited Consolidated Financial Statements

1. Basis of Presentation and Summary of Significant Accounting Policies

Basis of Presentation

The unaudited consolidated financial statements of Harvard Bioscience, Inc. and its wholly-owned subsidiaries (collectively, “Harvard Bioscience,” the “Company,” “our” or “we”) as of March 31, 2010 and for the three months ended March 31, 2010 and 2009 have been prepared by the Company pursuant to the rules and regulations of the Securities and Exchange Commission (“SEC”). Certain information and footnote disclosures normally included in financial statements prepared in accordance with U.S. generally accepted accounting principles (“U.S. GAAP”) have been condensed or omitted pursuant to such rules and regulations. The December 31, 2009 consolidated balance sheet was derived from audited financial statements, but does not include all disclosures required by U.S. GAAP. However, the Company believes that the disclosures are adequate to make the information presented not misleading. These unaudited consolidated financial statements should be read in conjunction with the consolidated financial statements and the notes thereto included in the Company’s Annual Report on Form 10-K for the fiscal year ended December 31, 2009, which was filed with the SEC on March 11, 2010.

In the opinion of management, all adjustments, which include normal recurring adjustments necessary to present a fair statement of financial position as of March 31, 2010, results of operations for the three months ended March 31, 2010 and 2009 and cash flows for the three months ended March 31, 2010 and 2009, as applicable, have been made. The results of operations for the three months ended March 31, 2010 are not necessarily indicative of the operating results for the full fiscal year or any future periods.

Summary of Significant Accounting Policies

The accounting policies underlying the accompanying unaudited consolidated financial statements are those set forth in Note 2 to the consolidated financial statements included in the Company’s Annual Report on Form 10-K for the year ended December 31, 2009, which was filed with the SEC on March 11, 2010.

2. Recently Issued Accounting Pronouncements

In October 2009, the FASB issued Accounting Standard Update (“ASU”) No. 2009-13—“*Revenue Recognition (Topic 605): Multiple-Deliverable Revenue Arrangements*.” This ASU establishes the accounting and reporting guidance for arrangements including multiple revenue-generating activities. This ASU provides amendments to the criteria for separating deliverables, and measuring and allocating arrangement consideration to one or more units of accounting. The amendments in this ASU also establish a selling price hierarchy for determining the selling price of a deliverable. Significantly enhanced disclosures are also required to provide information about a vendor’s multiple-deliverable revenue arrangements, including information about the nature and terms, significant deliverables, and its performance within arrangements. The amendments also require providing information about the significant judgments made and changes to those judgments and about how the application of the relative selling-price method affects the timing or amount of revenue recognition. The amendments in this ASU are effective prospectively for revenue arrangements entered into or materially modified in the fiscal years beginning on or after June 15, 2010. Early application is permitted. We believe adoption of this new guidance will not have a material impact on our consolidated results of operations or financial position.

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3. Goodwill and Other Intangible Assets

Intangible assets consist of the following:

	March 31, 2010		December 31, 2009		Weighted Average Life (a)
	(in thousands)				
	Gross	Accumulated Amortization	Gross	Accumulated Amortization	
Amortizable intangible assets:					
Existing technology	\$10,796	\$ (7,407)	\$11,234	\$ (7,525)	5.1 years
Tradenname	4,123	(758)	4,123	(689)	13.6 years
Distribution agreement/customer relationships	17,726	(4,203)	17,884	(3,927)	12.9 years
Patents	9	(5)	9	(5)	6.1 years
Total amortizable intangible assets	<u>\$32,654</u>	<u>\$ (12,373)</u>	<u>\$33,250</u>	<u>\$ (12,146)</u>	
Unamortizable intangible assets:					
Goodwill	\$31,362		\$32,108		
Other indefinite lived intangible assets	1,280		1,301		
Total goodwill and other indefinite lived intangible assets	<u>\$32,642</u>		<u>\$33,409</u>		
Total intangible assets	<u>\$65,296</u>		<u>\$66,659</u>		

(a) Weighted average life is as of March 31, 2010.

The change in the carrying amount of goodwill for the three months ended March 31, 2010 is as follows:

	(in thousands)
Balance at December 31, 2009	\$ 32,108
Effect of change in foreign currencies	(746)
Balance at March 31, 2010	<u>\$ 31,362</u>

Intangible asset amortization expense from continuing operations was \$0.5 million and \$0.3 million for the three months ended March 31, 2010 and 2009, respectively. Amortization expense of existing amortizable intangible assets is currently estimated to be \$2.3 million for the years ending December 31, 2010 and 2011, \$2.0 million for the year ending December 31, 2012, \$1.9 million for the year ending 2013 and \$1.7 million for the year ending December 31, 2014.

4. Inventories

Inventories consist of the following:

	March 31, 2010	December 31, 2009
	(in thousands)	
Finished goods	\$ 6,812	\$ 7,116
Work in process	484	559
Raw materials	6,724	6,731
Total	<u>\$14,020</u>	<u>\$ 14,406</u>

5. Restructuring and Other Exit Costs

2009 Restructuring Plan

During the quarter ended March 31, 2009, the management of Harvard Bioscience initiated a plan to relocate the Scie-Plas operation to Hoefer's San Francisco, California facility and exit its general fabrication business as part of its ongoing business improvement initiative. During the quarter ended June 30, 2009, Biochrom's management initiated a plan to improve Biochrom's manufacturing margins. The combined costs of these activities recorded in the year ended December 31, 2009 were \$0.7 million.

During the quarter ended March 31, 2010, no charges were recorded relating to the 2009 restructuring plan. During the quarter ended March 31, 2009, we recorded charges relating to this plan of approximately \$55,000. These charges were comprised of \$9,000 in severance payments, \$28,000 in inventory impairment charges related to the discontinuance of certain product lines (included in cost of product revenues) and \$18,000 in various other costs.

Activity and liability balances related to these restructuring charges in connection with the 2009 Restructuring Plan were as follows:

	<u>Severance and Related</u>	<u>Inventory</u>	<u>Facility Closure Costs</u> (in thousands)	<u>Other</u>	<u>Total</u>
Restructuring charges	\$ 326	\$ 163	\$ 14	\$188	\$ 691
Cash payments	(323)	(4)	(14)	(88)	(429)
Non-cash charges	—	(159)	—	(88)	(247)
Currency translation	(3)	—	—	(2)	(5)
Restructuring balance at December 31, 2009	<u>\$ —</u>	<u>\$ —</u>	<u>\$ —</u>	<u>\$ 10</u>	<u>\$ 10</u>
Cash payments	—	—	—	(2)	(2)
Non-cash charges	—	—	—	—	—
Currency translation	—	—	—	—	—
Restructuring balance at March 31, 2010	<u><u>\$ —</u></u>	<u><u>\$ —</u></u>	<u><u>\$ —</u></u>	<u><u>\$ 8</u></u>	<u><u>\$ 8</u></u>

We anticipate the remaining payments related to the 2009 Restructuring Plan will occur during the second quarter of 2010.

6. Acquisition

On September 2, 2009, the Company through its newly formed wholly-owned subsidiary, DAC Acquisition Holding, Inc., acquired substantially all of the assets of Denville Scientific, Inc. ("Denville"), a Delaware corporation with its principal offices in New Jersey.

Under the terms of the Asset Purchase Agreement, the Company made payments of approximately \$20.8 million in cash during 2009, and will make one additional contingent payment. We anticipate the final payment of approximately \$1.9 million will be paid in the second quarter of 2010. The Company expects to fund the final installment of the purchase price from its existing cash balances, its credit facility or both.

Denville is a supplier of molecular biology products, with a focus on liquid handling consumables utilized in research laboratories. We believe that the acquisition of Denville Scientific will bring to Harvard Bioscience a well-established business with an excellent organic growth history, an extensive field sales organization throughout the United States and a significant consumables business.

The balance of the contingent consideration of approximately \$1.9 million was recorded in other current liabilities in our consolidated balance sheets as of December 31, 2009 and March 31, 2010.

The amounts of Denville's revenue and net income included in the consolidated statement of operations for the three months ended March 31, 2010 are \$5.7 million and \$0.4 million, respectively.

The following consolidated pro forma information is based on the assumption that the acquisition occurred on January 1, 2009. Accordingly, the historical results have been adjusted to reflect amortization expense and interest costs that would have been recognized on such a pro forma basis. The unaudited pro forma information is presented for comparative purposes only and is not necessarily indicative of the financial position or results of operations which would have been reported had we completed the acquisition during these periods or which might be reported in the future.

Pro Forma	<u>Three Months Ended March 31, 2009</u> (in thousands)
Revenues	\$ 24,267
Net income	\$ 1,985

7. Warranties

Warranties are estimated and accrued for at the time sales are recorded. A rollforward of product warranties is as follows:

	<u>Beginning Balance</u>	<u>Payments</u> (in thousands)	<u>Additions</u>	<u>Ending Balance</u>
Year ended December 31, 2009	\$ 186	(56)	32	\$ 162
Three months ended March 31, 2010	\$ 162	(1)	8	\$ 169

8. Comprehensive Income

As of March 31, 2010, accumulated other comprehensive income consisted of cumulative foreign currency translation adjustments of \$(1.2) million and, in accordance with FASB ASC 715-20, "Compensation – Retirement Benefits, Defined Benefit Plans," \$(2.7) million to reflect the under-funded status of the Company's pension plans net of tax.

The components of total comprehensive income were as follows:

	<u>Three Months Ended March 31,</u>	
	<u>2010</u>	<u>2009</u>
	(in thousands)	
Net income	\$ 2,221	\$ 1,839
Other comprehensive loss	(2,120)	(982)
Comprehensive income	<u>\$ 101</u>	<u>\$ 857</u>

Other comprehensive income for the three months ended March 31, 2010 and 2009 consisted of foreign currency translation adjustments.

9. Employee Benefit Plans

Certain of the Company's United Kingdom subsidiaries, Harvard Apparatus Limited and Biochrom Limited, maintain contributory, defined benefit or defined contribution pension plans for substantially all of their employees. The components of the Company's defined benefit pension expense were as follows:

	<u>Three Months Ended March 31,</u>	
	<u>2010</u>	<u>2009</u>
	(in thousands)	
Components of net periodic benefit cost:		
Service cost	\$ 45	\$ 33
Interest cost	195	181
Expected return on plan assets	(148)	(132)
Net amortization loss	38	23
Net periodic benefit cost	<u>\$ 130</u>	<u>\$ 105</u>

For the three months ended March 31, 2010 and 2009, the Company made no contribution to its defined benefit plans. The Company expects to contribute approximately \$0.5 million to its defined benefit plans during 2010.

10. Capital Stock**Stock Repurchase Program**

On December 6, 2007, the Board of Directors authorized the repurchase by the Company of up to \$10 million of its common stock in the open market or through privately negotiated transactions over the next 24 months. On November 3, 2009, the Board of Directors extended this program for an additional year. Under the program, shares may be repurchased from time to time and in such amounts as market conditions warrant, subject to regulatory considerations and any applicable contractual restrictions.

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During the three months ended March 31, 2010, the Company repurchased no shares pursuant to this program. During the three months ended March 31, 2009, the company repurchased in the open market 434,074 shares of common stock at an aggregate cost of \$1.2 million, including commissions under the stock repurchase program.

At March 31, 2010, we had \$5.0 million remaining under the stock repurchase program authorization.

Repurchased shares have been recorded as treasury stock and will be held until the Company's Board of Directors designates that these shares be retired or used for other purposes.

Stock Option Plans

We account for share-based payment awards in accordance with the provisions of FASB ASC 718, which requires us to recognize compensation expense for all share-based payment awards made to employees and directors including employee stock options and employee stock purchases related to the Employee Stock Purchase Plan ("ESPP"). Stock-based compensation expense recognized under FASB ASC 718 for the three months ended March 31, 2010 and 2009 was \$0.6 million and \$0.3 million, respectively, which consisted of stock-based compensation expense related to employee stock options and the ESPP.

Valuation and Expense Information Under Share-Based Payment Accounting

Stock-based compensation expense related to employee stock options and the ESPP under FASB ASC 718 for the three months ended March 31, 2010 and 2009, respectively, was allocated as follows:

	Three Months Ended March 31,	
	2010	2009
	(in thousands)	
Cost of sales	\$ 16	\$ 11
Sales and marketing	11	(4)
General and administrative	528	304
Research and development	3	1
Total stock-based compensation	<u>\$ 558</u>	<u>\$ 312</u>

We did not capitalize any stock-based compensation. No significant tax benefit on the stock-based compensation was recorded in the three months ended March 31, 2010 and 2009 since the Company has established a valuation allowance against net deferred tax assets.

Stock-based compensation expense recognized in the consolidated statement of operations for the three months ended March 31, 2010 and 2009 is based on awards ultimately expected to vest and has been reduced for annualized estimated forfeitures of 4.83% and 5.96%, respectively. Share-based-payment accounting requires forfeitures to be estimated at the time of grant and revised, if necessary, in subsequent periods if actual forfeitures differ from those estimates. Forfeitures were estimated based on historical experience.

Weighted Average Common Shares Outstanding

Basic income per share is based upon net income divided by the number of weighted average common shares outstanding during the period. The calculation of diluted net income per share assumes conversion of stock options into common stock using the treasury method. The weighted average number of shares used to compute basic and diluted earnings per share consists of the following:

	Three Months Ended March 31,	
	2010	2009
Basic	29,584,436	30,011,732
Effect of assumed conversion of employee and director stock options	356,781	108,378
Diluted	<u>29,941,217</u>	<u>30,120,110</u>

Excluded from the shares used in calculating the diluted earnings per common share in the above table are options to purchase approximately 5,781,850 and 5,615,572 shares of common stock for the three months ended March 31, 2010 and 2009, respectively, as the impact of these shares would be anti-dilutive.

11. Revolving Credit Facility

During 2003, we entered into a \$20.0 million credit facility with Brown Brothers Harriman & Co. On December 1, 2006, we amended the terms of the credit facility extending the maturity date from January 1, 2007 to December 1, 2009.

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On August 7, 2009, we entered into an amended and restated \$20.0 million revolving credit loan agreement with Bank of America, as agent, and Bank of America and Brown Brothers Harriman & Co as lenders. The amended and restated revolving credit facility will mature on August 7, 2012. Borrowings under the credit facility bear interest at the London Interbank Offered Rate (“LIBOR”) plus 4.0%. The amended and restated facility includes covenants relating to income, debt coverage and cash flow, as well as minimum working capital requirements. The credit facility also contains limitations on our ability to incur additional indebtedness and requires lender approval for acquisitions funded with cash, promissory notes and/or other consideration in excess of \$6.0 million and for acquisitions funded solely with equity in excess of \$10.0 million.

As of March 31, 2010 and December 31, 2009, we had \$11.8 million and \$13.3 million, respectively, outstanding under our credit facility. The borrowings under the credit facility were related to our recent acquisition of Denville Scientific. As of March 31, 2010, we were in compliance with all financial covenants contained in the credit facility; we were not subject to any borrowing restrictions under the financial covenants and had available borrowing capacity under our revolving credit facility of \$8.2 million.

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

Forward Looking Statements

This Quarterly Report on Form 10-Q contains statements that are not statements of historical fact and are forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. The forward-looking statements are principally, but not exclusively, contained in “Item 2: Management’s Discussion and Analysis of Financial Condition and Results of Operations.” These statements involve known and unknown risks, uncertainties and other factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. Forward-looking statements include, but are not limited to, statements about management’s confidence or expectations, and our plans, objectives, expectations and intentions that are not historical facts. In some cases, you can identify forward-looking statements by terms such as “may,” “will,” “should,” “could,” “would,” “expects,” “plans,” “anticipates,” “believes,” “goals,” “sees,” “estimates,” “projects,” “predicts,” “intends,” “think,” “potential,” “objectives,” “optimistic,” “strategy,” and similar expressions intended to identify forward-looking statements. These statements reflect our current views with respect to future events and are based on assumptions and subject to risks and uncertainties. Given these uncertainties, you should not place undue reliance on these forward-looking statements. Factors that may cause the Company’s actual results to differ materially from those in the forward-looking statements include the Company’s failure to identify potential acquisition candidates, successfully integrate acquired businesses or technologies, successfully negotiate favorable pricing and other terms with acquisition candidates to enable potential acquisitions to close, complete consolidations of business functions, expand our distribution channels, expand our product offerings, introduce new products or commercialize new technologies, including in the field of regenerative medicine, unanticipated costs relating to acquisitions, unanticipated costs arising in connection with the Company’s consolidation of business functions and any restructuring initiatives, decreased demand for the Company’s products due to changes in our customers’ needs, our ability to obtain regulatory approvals, including FDA approval, for our products including any products in the field of regenerative medicine, the current size or anticipated size of the regenerative medicine market, the existence and size of opportunities in the regenerative medicine market, our financial position, general economic outlook, or other circumstances, overall economic trends, the seasonal nature of purchasing in Europe, economic, political and other risks associated with international revenues and operations, the impact of the current economic and financial crisis, additional costs of complying with recent changes in regulatory rules applicable to public companies, our ability to manage our growth, our ability to retain key personnel, competition from our competitors, technological changes resulting in our products becoming obsolete, future changes to the operations or the activities of our subsidiaries due to manufacturing consolidations, our ability to meet the financial covenants contained in our credit facility, our ability to protect our intellectual property and operate without infringing on others’ intellectual property, potential costs of any lawsuits to protect or enforce our intellectual property, economic and political conditions generally and those affecting pharmaceutical and biotechnology industries, research funding levels from endowments at our university customers, impact of any impairment of our goodwill or intangible assets, our acquisition of Genomic Solutions failing to qualify as a tax-free reorganization for federal tax purposes, the amount of earn-out consideration that the Company receives in connection with the disposition of the Company’s Capital Equipment Business segment and factors that may impact the receipt of this consideration, such as the revenues of the businesses disposed of, plus factors described under the heading “Item 1A. Risk Factors” in the Company’s Annual Report on Form 10-K for the fiscal year ended December 31, 2009, filed with the SEC on March 11, 2010. Our results may also be affected by factors of which we are not currently aware. Also, these forward-looking statements represent our estimates and assumptions only as of the date of this report. We may not update these forward-looking statements, even though our situation may change in the future, unless we have obligations under the federal securities laws to update and disclose material developments related to previously disclosed information.

General

Our strategy focuses on creating value through combining tuck-under acquisitions with organic growth and operational improvements. One of the major drivers of our success in the first quarter of 2010 was our acquisition of Denville Scientific (“Denville”), which occurred in September 2009. Denville continued to perform well during the first quarter of 2010 and has been

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accretive to our earnings per share since the acquisition. In addition to Denville's positive impact on the first quarter, our core business also performed well. Order growth was widespread but we were particularly pleased to see increases in the sales of our new syringe pumps launched last year.

We expect the main drivers of organic growth going forward to be our continued expansion of our distribution channels and the introduction of new products. Throughout last year and continuing into the first quarter 2010, we maintained our investments in adding field sales people in our Harvard Apparatus, Biochrom and Denville businesses and in introducing new products. In 2009, we launched two new syringe pump products at our Harvard Apparatus business, the KDS Legato 200 and the Harvard Apparatus PHD Ultra., which contributed to the growth in the first quarter 2010, and we expect to launch the third in this series in the second quarter of 2010. We believe these new pumps significantly improve the performance and ease of use of our syringe pump product line.

We expect to launch a major new spectrophotometer at our Biochrom business. Spectrophotometry is the core of our Biochrom product line, and we believe this new product platform will provide both improvements to the technical specifications, such as accuracy and reproducibility, as well as ease of use by adding a color touch screen user interface. This new product has been designed as a platform that we believe will enable us to significantly expand the range of our products at very little extra cost. In addition to our traditional strength in single beam spectrophotometers, we will now be adding dual beam instruments (which are inherently more accurate than single beam instruments) and variable bandwidth instruments, which provide significant extra flexibility to the user. With these new products we believe we will be able to access a larger segment of the entire spectrophotometer market. We think that these new products will help to drive organic growth during 2010 and beyond.

We continue to pursue our tuck-under acquisition strategy.

We believe that through execution of our strategy of organic growth, tuck-under acquisitions and operational improvements that we will be able to strengthen the company and position ourselves well for when the economy recovers. While we expect the initiatives discussed above to positively impact our business, the success of these initiatives is subject to a number of factors, including fluctuations in foreign exchange rates, the current economic and financial condition and their impact on our customers and our ability to obtain credit on terms favorable to us, the competitiveness of our new products, the strength of our intellectual property underlying these products, the success of our marketing efforts and those of our distributors and the other factors described under the heading "Item 1A. Risk Factors" in our Annual Report on Form 10-K filed with the SEC on March 11, 2010.

Our goal is to develop and sell products that improve life science research and as such, we monitor our operating metrics and when appropriate, effect organizational changes to leverage infrastructure and distribution channels. These changes may be effected as a result of various events, including acquisitions, the worldwide economy, general market conditions and personnel changes.

Financing

On August 7, 2009, we entered into an amended and restated \$20.0 million revolving credit loan agreement with Bank of America, as agent, and Bank of America and Brown Brothers Harriman & Co as lenders. The amended and restated revolving credit facility will mature on August 7, 2012. Borrowings under the credit facility bear interest at LIBOR plus 4.0%. The new facility includes covenants relating to income, debt coverage and cash flow, as well as minimum working capital requirements. The credit facility also contains limitations on our ability to incur additional indebtedness and requires lender approval for acquisitions funded with cash, promissory notes and/or other consideration in excess of \$6.0 million and for acquisitions funded solely with equity in excess of \$10.0 million.

At March 31, 2010, we had \$11.8 million outstanding under our credit facility with Bank of America and Brown Brothers Harriman & Co.

Historically, we have funded acquisitions with debt, capital raised by issuing equity and cash flow from operations. In order to continue the acquisition portion of our growth strategy beyond what our current cash balances and cash flow from operations can support, we will need to raise more capital, either by incurring additional debt, issuing equity or a combination.

Components of Operating Income

Revenues. We generate revenues by selling apparatus, instruments, devices and consumables through our catalog, our distributors, our direct sales force and our website. For products primarily priced under \$10,000, we typically distribute a new, comprehensive catalog every one to three years, initially in a series of bulk mailings, first to our existing customers, followed by mailings to targeted markets of potential customers. Over the life of the catalog, distribution will also be made periodically to potential and existing customers through direct mail and trade shows and in response to e-mail and telephone inquiries. From time to time, we also distribute catalog supplements that promote selected areas of our catalog or new products to targeted subsets of our customer base. Future editions of our comprehensive catalog and our catalog supplements will be timed at least in part with the incidence of new product introductions. Our end user customers are research scientists at pharmaceutical and biotechnology companies, universities and government laboratories. Revenue from catalog sales in any period is influenced by the amount of time elapsed since

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the last mailing of the catalog, the number of catalogs mailed and the number of new items included in the catalog. We launched our latest comprehensive catalog in March 2010, with approximately 850 pages, 11,000 products and approximately 65,000 copies printed. Revenues from direct sales to end users, derived through our catalog and the electronic version of our catalog on our website, represented approximately 25% and 30%, respectively, of our revenues for the three months ended March 31, 2010 and for the year ended December 31, 2009.

Products sold under brand names of distributors, including GE Healthcare, are typically priced in the range of \$5,000-\$15,000. They are mainly scientific instruments like spectrophotometers and plate readers that analyze light to detect and quantify a very wide range of molecular and cellular processes or apparatus like gel electrophoresis units. We also use distributors for both our catalog products and our higher priced products, for sales in locations where we do not have subsidiaries or where we have distributors in place for acquired businesses. For the three months ended March 31, 2010 and for the year ended December 31, 2009, approximately 46% and 48%, respectively, of our revenues were derived from sales to distributors.

For the three months ended March 31, 2010, approximately 68% of our revenues were derived from products we manufacture; approximately 10% were derived from complementary products we distribute in order to provide the researcher with a single source for all equipment needed to conduct a particular experiment and 22% were derived from distributed products sold under our brand names. For the year ended December 31, 2009, approximately 76% of our revenues were derived from products we manufacture; approximately 15% were derived from complementary products we distribute in order to provide the researcher with a single source for all equipment needed to conduct a particular experiment and 9% were derived from distributed products sold under our brand names.

For the three months ended March 31, 2010 and for the year ended December 31, 2009, approximately 43% and 52%, respectively, of our revenues were derived from sales made by our non-U.S. operations. A large portion of our international sales during these periods consisted of sales to GE Healthcare, the distributor for our spectrophotometers and plate readers. GE Healthcare distributes these products to customers around the world, including to many customers in the United States, from its distribution center in Upsalla, Sweden. As a result, we believe our international sales would have been a lower percentage of our revenues if we had shipped our products directly to our end-users. Changes in the relative proportion of our revenue sources between catalog sales, direct sales and distribution sales are primarily the result of a different sales proportion of acquired companies.

Cost of product revenues. Cost of product revenues includes material, labor and manufacturing overhead costs, obsolescence charges, packaging costs, warranty costs, shipping costs and royalties. Our cost of product revenues may vary over time based on the mix of products sold. We sell products that we manufacture and products that we purchase from third parties. The products that we purchase from third parties have a higher cost of product revenues as a percent of revenue because the profit is effectively shared with the original manufacturer. We anticipate that our manufactured products will continue to have a lower cost of product revenues as a percentage of revenues as compared with the cost of non-manufactured products for the foreseeable future. Additionally, our cost of product revenues as a percent of product revenues will vary based on mix of direct to end user sales and distributor sales, mix by product line and mix by geography.

Sales and marketing expenses. Sales and marketing expense consists primarily of salaries and related expenses for personnel in sales, marketing and customer support functions. We also incur costs for travel, trade shows, demonstration equipment, public relations and marketing materials, consisting primarily of the printing and distribution of our catalogs, supplements and the maintenance of our websites. We may from time to time expand our marketing efforts by employing additional technical marketing specialists in an effort to increase sales of selected categories of products in our catalog. We may also from time to time expand our direct sales organizations in an effort to concentrate on key accounts or promote certain product lines.

General and administrative expenses. General and administrative expense consists primarily of salaries and other related costs for personnel in executive, finance, accounting, information technology and human relations functions. Other costs include professional fees for legal and accounting services, facility costs, investor relations, insurance and provision for doubtful accounts.

Research and development expenses. Research and development expense consists primarily of salaries and related expenses for personnel and spending to develop and enhance our products and to support collaboration agreements. Other research and development expense includes fees for consultants and outside service providers, and material costs for prototype and test units. We expense research and development costs as incurred. We believe that investment in product development is a competitive necessity and plan to continue to make these investments in order to realize the potential of new technologies that we develop, license or acquire for existing markets. Additionally, we are working to develop new products aimed at long term opportunities in the emerging field of regenerative medicine.

Stock compensation expenses. Stock-based compensation expense recognized under FASB ASC 718, "Compensation – Stock Compensation," was \$0.6 million and \$0.3 million for the three months ended March 31, 2010 and 2009, respectively. This stock-based compensation expense was related to employee stock options and the employee stock purchase plan and was recorded as a component of cost of product revenues, sales and marketing expenses, general and administrative expenses and research and development expenses.

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Income Taxes

As described in Note 13 to our consolidated financial statements in our Annual Report on Form 10-K for the year ended December 31, 2009 filed with the SEC on March 11, 2010, we have recorded a full valuation allowance against most United States deferred tax assets, net of deferred tax liabilities except deferred tax liabilities related to indefinite lived intangible assets. As we gain more insight into the amount and predictability of taxable income from Denville, our investment requirements for regenerative medicine and certain other factors impacting our U.S. tax position, we may reverse some or all of the valuation allowance in future periods.

Selected Results of Operations

Three months ended March 31, 2010 compared to three months ended March 31, 2009:

	Three Months Ended March 31,		Dollar Change	% Change
	2010	2009		
	(dollars in thousands, unaudited)			
Revenues	\$26,300	\$19,072	\$7,228	37.9%
Cost of product revenues	13,518	9,662	3,856	39.9%
Gross margin percentage	48.6%	49.3%	N/A	-1.5%
Sales and marketing expenses	3,807	2,372	1,435	60.5%
General and administrative expenses	4,261	3,317	944	28.5%
Research and development expenses	1,207	999	208	20.8%

Revenues.

Revenues increased \$7.2 million, or 37.9%, to \$26.3 million for the three months ended March 31, 2010 compared to \$19.1 million for the same period in 2009. Our Denville Scientific subsidiary, which we acquired on September 2, 2009, contributed approximately \$5.7 million to first quarter 2010 revenues. The effect of a weakened U.S. dollar increased the Company's first quarter revenues by \$0.7 million, or 3.7%, compared with the same period in 2009. Adjusting for the effect of foreign currency fluctuation and excluding Denville, revenues were up \$0.8 million, or 4.3%, year-to-year and reflected organic growth across our Harvard Apparatus, Biochrom and electrophoresis businesses.

Cost of product revenues.

Cost of product revenues increased \$3.9 million, or 39.9%, to \$13.5 million for the three months ended March 31, 2010 compared with \$9.7 million for the three months ended March 31, 2009. The increase in cost of product revenues included \$3.6 million attributable to our Denville Scientific subsidiary and \$0.4 million from the currency effect of a weaker U.S. dollar, which were partially offset by the effects of cost reductions related to our operational improvement initiatives. Gross profit as a percentage of revenues decreased to 48.6% for the three months ended March 31, 2010 compared with 49.3% for the same period in 2009. The decrease in gross profit as a percentage of revenues was primarily due to the impact of Denville Scientific, which has lower gross margins than our overall average margin. First quarter 2010 gross margin as a percentage of revenues, excluding Denville, was 51.7%, which reflected the effects of operational improvement initiatives completed during 2009, ongoing cost improvement efforts and a more favorable sales mix in the first quarter of 2010 compared with the first quarter of 2009.

Sales and marketing expense.

Sales and marketing expenses increased \$1.4 million, or 60.5%, to \$3.8 million for the three months ended March 31, 2010 compared with \$2.4 million for the three months ended March 31, 2009. This increase was primarily due to expenses from our recently acquired Denville Scientific subsidiary of \$1.1 million, increased marketing efforts in all of our businesses and a \$0.1 million adverse impact of currency exchange rates.

General and administrative expense.

General and administrative expenses increased \$0.9 million, or 28.5%, to \$4.3 million for the three months ended March 31, 2010 compared with \$3.3 million for the three months ended March 31, 2009. The year-to-year quarterly increase was primarily due to \$0.3 million of expenses related to our Denville subsidiary acquisition, a \$0.2 million increase in stock compensation expense, \$0.1 million from the effect of foreign exchange and a \$0.3 million increase in other general and administrative areas combined.

Research and development expense.

Research and development expenses increased 20.8%, or \$0.2 million, to \$1.2 million for the three months ended March 31, 2010 compared with \$1.0 million for the same period in 2009. The increase in research and development expenses was primarily due to \$0.1 million of spending related to regenerative medicine and increased development efforts in the Biochrom group, which were partially offset by decreased expenses in the Harvard Apparatus group.

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Amortization of intangible assets.

Amortization of intangibles was \$0.5 million and \$0.3 million for the three months ended March 31, 2010 and 2009, respectively.

Other income, net.

Other income and expense, net, was \$0.2 million expense and \$0.1 million income for the three months ended March 31, 2010 and 2009, respectively. Net interest expense was \$0.1 million for the three months ended March 31, 2010 compared to net interest expense of \$38,000 for the three months ended March 31, 2009. The increase in net interest expense was primarily due to higher average debt balances in the first quarter of 2010 compared to the first quarter of 2009. Other income, net, also included foreign exchange losses of \$26,000 for the three months ended March 31, 2010 and foreign exchange gains of \$0.1 million for the three months ended March 31, 2009. These exchange gains and losses were primarily the result of currency fluctuations on intercompany transactions between our subsidiaries.

Income taxes.

Income tax expense was approximately \$0.6 million for the three months ended March 31, 2010 and 2009. The effective income tax rate was 21.3% for the three months ended March 31, 2010, compared with 24.7% for the same period of 2009. The difference between our effective tax rate and the US statutory tax rate is principally attributable to foreign tax rate differential and changes in our valuation allowance.

Restructuring

During the quarter ended March 31, 2009, the management of Harvard Bioscience initiated a plan to relocate the Scie-Plas operation to Hoefer's San Francisco location and exit the Scie-Plas general fabrication business as part of our ongoing business improvement initiative.

During the year ended December 31, 2009, we recorded restructuring charges in our Scie-Plas, Biochrom and Hoefer businesses related to the 2009 restructuring plan of approximately \$0.7 million. These charges were comprised of \$0.3 million in severance payments, \$0.2 million in inventory impairment charges related to the discontinuance of certain product lines (included in cost of product revenues) and \$0.2 million in various other costs.

No charges were recorded during the three months ended March 31, 2010 related to the 2009 restructuring plan.

Liquidity and Capital Resources

Historically, we have financed our business through cash provided by operating activities, the issuance of common stock and preferred stock, and bank borrowings. Our liquidity requirements have arisen primarily from investing activities, including funding of acquisitions, working capital and capital expenditures.

The Company ended the first quarter of 2010 with cash and cash equivalents of \$17.0 million compared to \$16.6 million at December 31, 2009. As of March 31, 2010 and December 31, 2009, the Company had \$11.8 million and \$13.3 million, respectively, outstanding under its credit facility. The borrowings under the credit facility are related to our acquisition of Denville Scientific. Total cash and equivalents, net of debt, was \$5.2 million and \$3.3 million at March 31, 2010 and December 31, 2009, respectively.

Overview of Cash Flows
(in thousands, unaudited)

	Three Months Ended March 31,	
	2010	2009
Cash flows from operations:		
Net income	\$ 2,221	\$ 1,839
Changes in assets and liabilities	(623)	1,424
Other adjustments to operating cash flows	1,510	1,028
Net cash provided by operating activities	3,108	4,291
Investing activities:		
Net cash used in investing activities	(528)	(310)
Financing activities:		
Repayments of debt, net	(1,508)	(376)
Other financing activities	0	(1,206)
Net cash used in financing activities	(1,508)	(1,582)
Effect of exchange rate changes on cash	(617)	(214)
Increase in cash and cash equivalents	<u>\$ 455</u>	<u>\$ 2,185</u>

Our operating activities generated cash of \$3.1 million for the three months ended March 31, 2010 compared to \$4.3 million for the three months ended March 31, 2009. The decrease in cash flows from operations was primarily due to changes in working capital year to year.

Our investing activities used cash of \$0.5 million during the three months ended March 31, 2010 compared to \$0.3 million during the three months ended March 31, 2009. Investing activities during the first quarters of both 2009 and 2010 included purchases of property, plant and equipment and expenditures for our catalogs. During the first quarter of 2010, catalog costs reflect the publication and distribution of an 850-page Harvard Apparatus catalog. We expect to make approximately \$1.2 million of capital expenditures during 2010.

Our financing activities have historically consisted of borrowings and repayments under a revolving credit facility, long-term debt, the issuance of preferred stock and common stock, including the common stock issued in our initial public offering, and repurchases of our common stock under our stock repurchase program. During the three months ended March 31, 2010, financing activities used cash of \$1.5 million, which consisted of repayments of borrowings under our credit facility. These borrowings under our credit facility related to our acquisition of Denville Scientific in September 2009. During the first quarter of 2009, we made repayments of \$0.4 million of notes payable to banks and we repurchased in the open market approximately 0.4 million shares of our common stock at a cost of \$1.2 million, including commissions.

On August 7, 2009, we entered into an amended and restated \$20.0 million revolving credit loan agreement with Bank of America, as agent, and Bank of America and Brown Brothers Harriman & Co as lenders. The amended and restated revolving credit facility will mature on August 7, 2012. Borrowings under the credit facility bear interest at LIBOR plus 4.0%. The new facility includes covenants relating to income, debt coverage and cash flow, as well as minimum working capital requirements. The credit facility also contains limitations on our ability to incur additional indebtedness and requires lender approval for acquisitions funded with cash, promissory notes and/or other consideration in excess of \$6.0 million and for acquisitions funded solely with equity in excess of \$10.0 million.

Our forecast of the period of time through which our financial resources will be adequate to support our operations is a forward-looking statement that involves risks and uncertainties, and actual results could vary as a result of a number of factors. Based on our current operations and current operating plans, we expect that our available cash, cash generated from current operations and debt capacity will be sufficient to finance current operations and capital expenditures for 12 months and beyond. However, we may use substantial amounts of capital to accelerate product development or expand our sales and marketing activities. We may need to raise additional capital in order to make significant acquisitions. Additional capital raising activities will dilute the ownership interests of existing stockholders to the extent we raise capital by issuing equity securities and we cannot assure you that we will be successful in raising additional capital on favorable terms or at all.

Impact of Foreign Currencies

We sell our products in many countries and a substantial portion of our sales, costs and expenses are denominated in foreign currencies, especially the British pound sterling and the Euro.

Changes in foreign currency exchange rates resulted in increases in revenues of \$0.7 million and expenses of \$0.6 million (net \$0.1 million) during the three months ended March 31, 2010.

The loss associated with the translation of foreign equity into U.S. dollars was approximately \$2.1 million during the three months ended March 31, 2010 compared to a loss associated with the translation of foreign equity into U.S. dollars of approximately \$1.0 million during the three months ended March 31, 2009. In addition, currency fluctuations resulted in approximately \$0.03 million in foreign currency losses during the three months ended March 31, 2010 and \$0.1 million in foreign currency gains during the three months ended March 31, 2009.

As of March 31, 2010 and December 31, 2009, we had \$11.8 million and \$13.3 million, respectively, outstanding under our credit facility. The borrowings under our credit facility were related to our September 2009 acquisition of Denville Scientific.

Critical Accounting Policies

The critical accounting policies underlying the accompanying unaudited consolidated financial statements are those set forth in Part II, Item 7 included in the Company's Annual Report on Form 10-K for the year ended December 31, 2009 filed with the SEC on March 11, 2010.

Recent Accounting Pronouncements

In October 2009, the FASB issued ASU No. 2009-13 – "Revenue Recognition (Topic 605): Multiple-Deliverable Revenue Arrangements." This ASU establishes the accounting and reporting guidance for arrangements including multiple revenue-generating activities. This ASU provides amendments to the criteria for separating deliverables, measuring and allocating arrangement consideration to one or more units of accounting. The amendments in this ASU also establish a selling price hierarchy for determining the selling price of a deliverable. Significantly enhanced disclosures are also required to provide information about a vendor's multiple-deliverable revenue arrangements, including information about the nature and terms, significant deliverables, and its performance within arrangements. The amendments also require providing information about the significant judgments made and changes to those judgments and about how the application of the relative selling-price method affects the timing or amount of revenue recognition. The amendments in this ASU are effective prospectively for revenue arrangements entered into or materially modified in the fiscal years beginning on or after June 15, 2010. Early application is permitted. We believe adoption of this new guidance will not have a material impact on our consolidated results of operations or financial position.

Item 3. Quantitative and Qualitative Disclosures About Market Risk.

We manufacture and test the majority of products in research centers in the United States, the United Kingdom, Germany and Spain. We sell our products globally through our direct catalog sales, direct sales force and indirect distributor channels. As a result, our financial results are affected by factors such as changes in foreign currency exchange rates and weak economic conditions in foreign markets.

We collect amounts representing a substantial portion of our revenues and pay amounts representing a substantial portion of our operating expenses in foreign currencies. As a result, changes in currency exchange rates from time to time may affect our operating results. In order to mitigate the impact of changes in foreign currency exchange rates, we use derivative financial instruments (or "foreign currency contracts") to hedge the foreign currency effects on the value of certain loans between subsidiaries and do not designate these derivative instruments as accounting hedges.

We are exposed to market risk from changes in interest rates primarily through our financing activities. As of March 31, 2010, we had \$11.8 million outstanding under our revolving credit facility, which bears interest at LIBOR plus 4.0%. At March 31, 2010, the interest rate on this debt was 4.25%. Assuming no other changes which would affect the margin of the interest rate under our revolving credit facility, the effect of interest rate fluctuations on outstanding borrowings under our revolving credit facility as of March 31, 2010 over the next twelve months is quantified and summarized as follows:

<u>If compared to the rate as of March 31, 2010</u>	<u>Interest Expense Increase (in thousands)</u>
Interest rates increase by 1.0%	\$ 118
Interest rates increase by 2.0%	\$ 236

Item 4. Controls and Procedures.

Disclosure controls and procedures (as defined in Rules 13a-15(e) and 15d-15(e) under the Exchange Act) are designed to ensure that information required to be disclosed in reports filed or submitted under the Exchange Act is recorded, processed, summarized, and reported within the time periods specified in SEC rules and forms and that such information is accumulated and communicated to management, including the Chief Executive Officer and the Chief Financial Officer, to allow timely decisions regarding required disclosures.

As required by Rules 13a-15(e) and 15d-15(e) under the Securities Exchange Act of 1934,, our management, under the supervision and with the participation of our Chief Executive Officer and Chief Financial Officer, conducted an evaluation of the effectiveness of the design and operation of our disclosure controls and procedures as of March 31, 2010. Our disclosure controls and procedures are designed to provide reasonable assurance that information required to be disclosed by us in the reports that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms, and our management necessarily was required to apply its judgment in evaluating and implementing our disclosure controls and procedures. Based upon the evaluation described above, our Chief Executive Officer and Chief Financial Officer have concluded that they believe that our disclosure controls and procedures were effective, as of the end of the period covered by this report, in providing reasonable assurance that information required to be disclosed by us in the reports that we file or submit under the Exchange Act is accumulated and communicated to our management, including our Chief Executive Officer and Chief Financial Officer, to allow timely decisions regarding required disclosures, and is recorded, processed, summarized and reported within the time periods specified in the SEC's rules and forms.

We continue to review our internal controls over financial reporting, and may from time to time make changes aimed at enhancing their effectiveness and to ensure that our systems evolve with our business. These efforts have led to various changes in our internal controls over financial reporting. There were no changes in our internal controls over financial reporting that occurred during the first quarter ended March 31, 2010 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

PART II. OTHER INFORMATION

Item 1A. Risk Factors

To our knowledge and except to the extent additional factual information disclosed in this Quarterly Report on Form 10-Q relates to such risk factors, there have been no material changes in the risk factors described in “Item 1A. Risk Factors” in our Annual Report on Form 10-K for the year ended December 31, 2009, which was filed with the SEC on March 11, 2010.

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

On December 6, 2007, the Board of Directors authorized the repurchase by the Company of up to \$10 million of its common stock in the open market or through privately negotiated transactions over 24 months. Under the program, shares could be repurchased from time to time and in such amounts as market conditions warranted, subject to regulatory considerations and any applicable contractual restrictions. On November 3, 2009, the Board of Directors extended this program for an additional year.

During the life of the program, we have repurchased in the open market 1,702,888 shares of common stock at an aggregate cost of \$5.0 million, including commissions under the stock repurchase program. There were no purchases by the Company of the Company’s common stock during the three months ended March 31, 2010.

Item 6. Exhibits

Exhibit Index

31.1+	Certification of Chief Financial Officer of Harvard Bioscience, Inc., pursuant to Rules 13a-14(a) and 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
31.2+	Certification of Chief Executive Officer of Harvard Bioscience, Inc., pursuant to Rules 13a-14(a) and 15d-14(a), as adopted pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
32.1*	Certification of Chief Financial Officer of Harvard Bioscience, Inc., pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
32.2*	Certification of Chief Executive Officer of Harvard Bioscience, Inc., pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.

+ Filed herewith.

* This certification shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, or otherwise subject to the liability of that section, nor shall it be deemed to be incorporated by reference into any filing under the Securities Act of 1933 or the Securities Exchange Act of 1934.

SIGNATURES

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

Date: May 7, 2010

HARVARD BIOSCIENCE, INC.

By: _____ /s/ CHANE GRAZIANO

Chane Graziano
Chief Executive Officer

By: _____ /s/ THOMAS MCNAUGHTON

Thomas McNaughton
Chief Financial Officer

Certification

I, Thomas McNaughton, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Harvard Bioscience, 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 consolidated 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: May 7, 2010

/s/ Thomas McNaughton

Thomas McNaughton
Chief Financial Officer

Certification

I, Chane Graziano, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Harvard Bioscience, 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 consolidated 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: May 7, 2010

/s/ Chane Graziano

Chane Graziano
Chief Executive Officer

CERTIFICATION OF PERIODIC FINANCIAL REPORT
PURSUANT TO 18 U.S.C. SECTION 1350

The undersigned officer of Harvard Bioscience, Inc. (the "Company") hereby certifies to his knowledge that the Company's Quarterly Report on Form 10-Q for the quarterly period ended March 31, 2010 (the "Report") to which this certification is being furnished as an exhibit, as filed with the Securities and Exchange Commission on the date hereof, fully complies with the requirements of Section 13(a) or 15(d), as applicable, of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), and that the information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company. This certification is provided solely pursuant to 18 U.S.C. Section 1350 and Item 601(b)(32) of Regulation S-K ("Item 601(b)(32)") promulgated under the Securities Act of 1933, as amended (the "Securities Act"), and the Exchange Act. In accordance with clause (ii) of Item 601(b)(32), this certification (A) shall not be deemed "filed" for purposes of Section 18 of the Exchange Act, or otherwise subject to the liability of that section, and (B) shall not be deemed to be incorporated by reference into any filing under the Securities Act or the Exchange Act, except to the extent that the Company specifically incorporates it by reference.

Date: May 7, 2010

/s/ Thomas McNaughton

Name: Thomas McNaughton

Title: Chief Financial Officer

CERTIFICATION OF PERIODIC FINANCIAL REPORT
PURSUANT TO 18 U.S.C. SECTION 1350

The undersigned officer of Harvard Bioscience, Inc. (the "Company") hereby certifies to his knowledge that the Company's Quarterly Report on Form 10-Q for the quarterly period ended March 31, 2010 (the "Report") to which this certification is being furnished as an exhibit, as filed with the Securities and Exchange Commission on the date hereof, fully complies with the requirements of Section 13(a) or 15(d), as applicable, of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), and that the information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company. This certification is provided solely pursuant to 18 U.S.C. Section 1350 and Item 601(b)(32) of Regulation S-K ("Item 601(b)(32)") promulgated under the Securities Act of 1933, as amended (the "Securities Act"), and the Exchange Act. In accordance with clause (ii) of Item 601(b)(32), this certification (A) shall not be deemed "filed" for purposes of Section 18 of the Exchange Act, or otherwise subject to the liability of that section, and (B) shall not be deemed to be incorporated by reference into any filing under the Securities Act or the Exchange Act, except to the extent that the Company specifically incorporates it by reference.

Date: May 7, 2010

/s/ Chane Graziano

Name: Chane Graziano
Title: Chief Executive Officer